

# APPROVED DRUG PRODUCTS

WITH

THERAPEUTIC  
EQUIVALENCE  
EVALUATIONS

38<sup>th</sup> EDITION

THE PRODUCTS IN THIS LIST HAVE BEEN APPROVED UNDER  
SECTION 505 OF THE FEDERAL FOOD, DRUG, AND COSMETIC ACT.

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES  
FOOD AND DRUG ADMINISTRATION  
OFFICE OF MEDICAL PRODUCTS AND TOBACCO  
CENTER FOR DRUG EVALUATION AND RESEARCH  
OFFICE OF GENERIC DRUGS  
OFFICE OF GENERIC DRUG POLICY

2018

# **APPROVED DRUG PRODUCTS with THERAPEUTIC EQUIVALENCE EVALUATIONS**

**The products in this list have been approved under section 505 of the Federal Food, Drug, and Cosmetic Act. This volume is current through December 31, 2017.**

## **38<sup>th</sup> EDITION**



**U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES  
FOOD AND DRUG ADMINISTRATION  
OFFICE OF MEDICAL PRODUCTS AND TOBACCO  
CENTER FOR DRUG EVALUATION AND RESEARCH  
OFFICE OF GENERIC DRUGS  
OFFICE OF GENERIC DRUG POLICY**

**2018**

**FOOD AND DRUG ADMINISTRATION  
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS  
With  
Therapeutic Equivalence Evaluations**

**CONTENTS**

	<i>PAGE</i>
PREFACE TO THIRTY EIGHTH EDITION.....	iv
1.0 INTRODUCTION .....	vi
1.1 Content and Exclusion .....	vi
1.2 Therapeutic Equivalence-Related Terms .....	vi
1.3 Further Guidance on Bioequivalence .....	ix
1.4 Reference Listed Drug and Reference Standard.....	ix
1.5 General Policies and Legal Status .....	x
1.6 Practitioner/User Responsibilities .....	xi
1.7 Therapeutic Equivalence Evaluations Codes .....	xiii
1.8 Description of Certain Special Situations .....	xx
1.9 Therapeutic Equivalence Code Change for a Drug Entity .....	xxiii
1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product.....	xxiv
1.11 Discontinued Section .....	xxiv
1.12 Changes to the Orange Book.....	xxiv
1.13 Availability of the Edition .....	xxv
2.0 HOW TO USE THE DRUG PRODUCTS LISTS .....	2-1
2.1 Key Sections for Using the Drug Product Lists .....	2-1
2.2 Drug Product Illustration .....	2-3
2.3 Therapeutic Equivalence Evaluations Illustration .....	2-4
 <b>DRUG PRODUCT LISTS</b>	
Prescription Drug Product List .....	3-1
OTC Drug Product List .....	4-1
Drug Products with Approval under Section 505 of the FD&C Act Administered by the Center for Biologics Evaluation and Research List .....	5-1
Discontinued Drug Product List .....	6-1
Orphan Products Designations and Approvals List .....	7-1
Drug Products Which Must Demonstrate <i>in vivo</i> Bioavailability Only if Product Fails to Achieve Adequate Dissolution .....	8-1
 <b>APPENDICES</b>	
A. Product Name Index .....	A-1
B. Product Name Index Listed by Applicant .....	B-1
C. Uniform Terms .....	C-1
 <b>PATENT AND EXCLUSIVITY INFORMATION ADDENDUM .....</b> AD1	
A. Patent and Exclusivity Lists .....	ADA1
B. Patent and Exclusivity Terms .....	ADB1

**FOOD AND DRUG ADMINISTRATION  
CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPROVED DRUG PRODUCTS  
With  
Therapeutic Equivalence Evaluations**

**PREFACE TO THIRTY EIGHTH EDITION**

The publication, *Approved Drug Products With Therapeutic Equivalence Evaluations* (the List, commonly known as the Orange Book), identifies drug products approved on the basis of safety and effectiveness by the Food and Drug Administration (FDA) under the Federal Food, Drug, and Cosmetic Act (the FD&C Act). The main criterion for the inclusion of any product is that the product is the subject of an application with an approval that has not been withdrawn for safety or efficacy reasons. Inclusion of products in the Orange Book is independent of any current regulatory action through administrative or judicial means against a drug product. In addition, the Orange Book contains therapeutic equivalence evaluations for approved multisource prescription drug products. These evaluations have been prepared to serve as public information and advice to state health agencies, prescribers, and pharmacists to promote public education in the area of drug product selection and to foster containment of health care costs. Therapeutic equivalence evaluations in this publication are not official FDA actions affecting the legal status of products under the FD&C Act.

**Background of the Publication.** To contain drug costs, virtually every state has adopted laws and/or regulations that encourage the substitution of drug products. These state laws generally require either that substitution be limited to drugs on a specific list (the positive formulary approach) or that it be permitted for all drugs except those prohibited by a particular list (the negative formulary approach). Because of the number of requests in the late 1970s for FDA assistance in preparing both positive and negative formularies, it became apparent that FDA could not serve the needs of each state on an individual basis. The Agency also recognized that providing a single list based on common criteria would be preferable to evaluating drug products on the basis of differing definitions and criteria in various state laws. As a result, on May 31, 1978, the Commissioner of the Food and Drug Administration sent a letter to officials of each state announcing FDA's intent to provide a list of all prescription drug products that are approved by FDA for safety and effectiveness, along with therapeutic equivalence determinations for multisource prescription products.

The Orange Book was distributed as a proposal in January 1979. It included only currently marketed prescription drug products approved by FDA through new drug applications (NDAs) and abbreviated new drug applications (ANDAs) under the provisions of Section 505 of the FD&C Act.

The therapeutic equivalence evaluations in the Orange Book reflect FDA's application of specific criteria to the multisource prescription drug products listed in the Orange Book and approved under Section 505 of the FD&C Act. These evaluations are presented in the form of code letters that indicate the basis for the evaluation made. An explanation of the codes appears in the *Introduction*.

A complete discussion of the background and basis of FDA's therapeutic equivalence evaluation policy was published in the *Federal Register* on January 12, 1979 (44 FR 2932). The final rule, which includes FDA's responses to the public comments on the proposal, was published in the *Federal Register* on October 31, 1980 (45 FR 72582). The first publication of the Orange Book in October 1980, concurrent with finalization of the rule, incorporated appropriate corrections and additions. Each subsequent edition has included new approvals and made appropriate changes in data.

On September 24, 1984, the President signed into law the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments). The Hatch-Waxman Amendments require that FDA, among other things, make publicly available a list of approved drug products with monthly supplements. The Orange Book and its monthly Cumulative Supplements satisfy this requirement. The *Addendum* to this publication identifies drugs that qualify under the FD&C Act for periods of exclusivity and provides patent information concerning the listed drugs. The *Addendum* also provides additional information that may be helpful to those submitting an NDA or ANDA to the Agency.

The Agency intends to use this publication to further its objective of obtaining input and comment on the publication itself and related Agency procedures. Therefore, if you have comments on how the publication can be improved, please send them to the Director, Division of Legal and Regulatory Support, Office of Generic Drug Policy, Office of Generic Drugs, Center for Drug Evaluation and Research, 7620 Standish Place, Rockville, MD 20855-2773. Comments received are publicly available to the extent allowable under the Freedom of Information Act and FDA regulations.

# 1.0 INTRODUCTION

## 1.1 Content and Exclusion

The Orange Book is composed of four parts: (1) approved prescription drug products with therapeutic equivalence evaluations; (2) approved over-the-counter (OTC) drug products for those drugs that may not be marketed without NDAs or ANDAs because they are not covered under existing OTC monographs; (3) drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research; and (4) a cumulative list of approved products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing.<sup>1</sup> This publication also includes indices of prescription and OTC drug products by trade name (proprietary name) or established name (if no trade name exists) and by applicant name (holder of the approved application), which have been abbreviated for this publication. Established names for active ingredients generally conform to official compendial names or *United States Adopted Names* (USAN) as described in (21 CFR 299.4(e)). A list of uniform terms is provided in Appendix C.

The *Addendum* contains patent and exclusivity information for the Prescription, OTC, Discontinued Drug Product Lists, and for the Drug Products with Approval under Section 505 of the FD&C Act Administered by the Center for Biologics Evaluation and Research. The publication may include additional information that the Agency deems appropriate to disseminate.

Prior to the 6th Edition, the publication had excluded OTC drug products and drug products with approval under Section 505 of the FD&C Act administered by the Center for Biologics Evaluation and Research. The Hatch-Waxman Amendments required the Agency to begin publishing an up-to-date list of all marketed drug products, OTC as well as prescription, that have been approved for safety and efficacy and for which new drug applications are required.

Under the FD&C Act, some drug products are given tentative approvals. The Agency will not include drug products with tentative approvals in the Orange Book because a drug product that is granted tentative approval is not an approved drug product. Tentative approval lists by month are available on FDA's website [Drugs@FDA](#). When the tentative approval becomes a final approval through a subsequent action letter to the applicant, the Agency will list the drug product and the date of approval in the appropriate approved drug product list.

Distributors or repackagers of products listed in the Orange Book are not identified.

## 1.2 Therapeutic Equivalence-Related Terms

---

<sup>1</sup> Newly approved products are added to parts 1, 2, or 3, of the Orange Book, depending on the dispensing requirements (prescription or OTC) or approval authority, unless the Orange Book staff is otherwise notified before publication.

**Pharmaceutical Equivalents.** Drug products are considered pharmaceutical equivalents if they contain the same active ingredients, are of the same dosage form and route of administration, and are formulated to contain the same amount of active ingredient and to meet the same or compendial or other applicable standards (i.e., strength, quality, purity, and identity).<sup>2</sup> They may differ in characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, within certain limits, labeling.

**Pharmaceutical Alternatives.** Drug products are considered pharmaceutical alternatives if they contain the same therapeutic moiety, but are different salts, esters, or complexes of that moiety, or are different dosage forms or strengths (e.g., tetracycline hydrochloride, 250mg capsules vs. tetracycline phosphate complex, 250mg capsules; quinidine sulfate, 200mg tablets vs. quinidine sulfate, 200mg capsules).<sup>3</sup> Different dosage forms and strengths within a product line by a single manufacturer are thus pharmaceutical alternatives, as are extended-release products when compared with immediate-release or standard-release formulations of the same active ingredient.

**Therapeutic Equivalents.** Drug products are considered to be therapeutic equivalents only if they are pharmaceutical equivalents for which bioequivalence has been demonstrated, and they can be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.<sup>4</sup>

FDA classifies as therapeutically equivalent those products that meet the following general criteria: (1) they are approved as safe and effective; (2) they are pharmaceutical equivalents in that they (a) contain identical amounts of the same active drug ingredient in the same dosage form and route of administration, and (b) meet compendial or other applicable standards of strength, quality, purity, and identity; (3) they are bioequivalent in that (a) they do not present a known or potential bioequivalence problem, and they meet an acceptable *in vitro* standard, or (b) if they do present such a known or potential problem, they are shown to meet an appropriate bioequivalence standard; (4) they are adequately labeled; and (5) they are manufactured in compliance with Current Good Manufacturing Practice regulations. *The concept of therapeutic equivalence, as used to develop the Orange Book, applies only to drug products containing the same active ingredient(s) and does not encompass a comparison of different therapeutic agents used for the same condition (e.g., meperidine hydrochloride vs. morphine sulfate for the treatment of pain).* Any drug product in the Orange Book repackaged and/or distributed by other than the applicant is considered to be therapeutically equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., BN). Distributors or repackagers of an applicant's drug product are not identified in the Orange Book.

FDA considers drug products to be therapeutically equivalent if they meet the criteria outlined above, even though they may differ in certain other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time and certain aspects of labeling (e.g., the presence of specific pharmacokinetic information), and storage conditions. When such differences are important in the care of a particular patient, it may be appropriate for the prescribing physician to require that a specific product be dispensed as a medical necessity. With this limitation, however, FDA believes that

---

<sup>2</sup> See generally 21 CFR 314.3(b).

<sup>3</sup> See generally 21 CFR 314.3(b).

<sup>4</sup> See generally 21 CFR 314.3(b).

products classified as therapeutically equivalent can be substituted with the full expectation that the substituted product will produce the same clinical effect and safety profile as the prescribed product.

**Strength.** Strength refers to the amount of drug substance contained in, delivered, or deliverable from a drug product, which includes: (1)(a) the total quantity of drug substance in mass or units of activity in a dosage unit or container closure (e.g., weight/unit dose, weight/volume or weight/weight in a container closure, or units/volume or units/weight in a container closure); and/or, as applicable, (b) the concentration of the drug substance in mass or units of activity per unit volume or mass (e.g., weight/weight, weight/volume, or units/volume); or (2) such other criteria the Agency establishes for determining the amount of drug substance contained in, delivered, or deliverable from a drug product if the weights and measures described in clause (1)(a) do not apply (e.g., certain drug-device combination products for which the amount of drug substance is emitted per use or unit time).<sup>5</sup> Note that if the criteria the Agency establishes for determining and expressing the amount of drug substance in a product evolves over time, the Agency generally does not intend to revise the expressions of strength for drug products already included in the Orange Book, but rather intends to apply the criteria prospectively to drug products added to the Orange Book.

Although the strength of drug products in the Orange Book is generally expressed in terms of the amount of drug substance (active ingredient) in the drug product, it is sometimes expressed in terms of the amount of the active moiety. For example, certain drug products included in the Orange Book include a designation of "EQ" next to their expression of strength. This "EQ" designation generally is used in connection with salt drug products to indicate that the strength of such drug product is being expressed in terms of the equivalent strength of the active moiety (e.g., "EQ 200MG BASE"), rather than in terms of the strength of the active ingredient.

**Bioavailability.** Bioavailability is the rate and extent to which the active ingredient or active moiety is absorbed from a drug product and becomes available at the site of drug action. For drug products that are not intended to be absorbed into the bloodstream, bioavailability may be assessed by scientifically valid measurements intended to reflect the rate and extent to which the active ingredient or active moiety becomes available at the site of drug action.

**Bioequivalence.** Bioequivalence is the absence of a significant difference in the rate and extent to which the active ingredient or active moiety in pharmaceutical equivalents or pharmaceutical alternatives becomes available at the site of drug action when administered at the same molar dose under similar conditions in an appropriately designed study. Section 505 (j)(8)(B) of the FD&C Act describes one set of conditions under which a test and reference listed drug (see Section 1.4) shall be considered bioequivalent:

the rate and extent of absorption of the [test] drug do not show a significant difference from the rate and extent of absorption of the [reference] drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses; or

the extent of absorption of the [test] drug does not show a significant difference from the extent of absorption of the [reference] drug when administered at the same molar dose of the therapeutic ingredient under

---

<sup>5</sup> See generally 21 CFR 314.3(b).

similar experimental conditions in either a single dose or multiple doses and the difference from the [reference] drug in the rate of absorption of the drug is intentional, is reflected in its proposed labeling, is not essential to the attainment of effective body drug concentrations on chronic use, and is considered medically insignificant for the drug.

Where these above methods are not applicable (e.g., for drug products that are not intended to be absorbed into the bloodstream), other scientifically valid *in vivo* or *in vitro* test methods to demonstrate bioequivalence may be appropriate.

For example, bioequivalence may sometimes be demonstrated using an *in vitro* bioequivalence standard, especially when such an *in vitro* test has been correlated with human *in vivo* bioavailability data. In other situations, bioequivalence may sometimes be demonstrated through comparative clinical trials or pharmacodynamic studies.

### **1.3 Further Guidance on Bioequivalence**

FDA's regulations and guidance documents provide additional information regarding bioequivalence and bioavailability, including methodologies and statistical criteria used to establish the bioequivalence of drug products.<sup>6</sup>

### **1.4 Reference Listed Drug and Reference Standard**

A reference listed drug (21 CFR 314.3(b)) means the listed drug identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA. Generally, a reference listed drug is a drug product approved in a new drug application under Section 505(c) of the FD&C Act based on full reports of investigations of safety and effectiveness. For an ANDA based on an approved suitability petition (a petitioned ANDA), the reference listed drug generally is the listed drug referenced in the approved suitability petition.<sup>7</sup>

A reference standard is the drug product selected by FDA that an applicant seeking approval of an ANDA must use in conducting an *in vivo* bioequivalence study required for approval. FDA generally selects a single reference standard that ANDA applicants must use in *in vivo* bioequivalence testing. Ordinarily, FDA will select the reference listed drug as the reference standard. However, in some instances (e.g., where the reference listed drug has been withdrawn from sale and FDA has determined it was not withdrawn for reasons of safety or effectiveness, and FDA selects an ANDA as the reference standard), the reference listed drug and the reference standard may be different.

---

<sup>6</sup> We note that prior editions of the Preface to the Orange Book included a section entitled "Statistical Criteria for Bioequivalence." Please see FDA's regulations and guidance documents for additional information regarding bioequivalence and bioavailability. See FDA Drugs guidance Web page at

<http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/default.htm>; FDA Drugs guidance (Product-Specific Recommendations for Generic Drug Development) Web page at

<http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/ucm075207.htm>; see generally 21 CFR part 320.

<sup>7</sup> 21 CFR 314.94(a)(3)(i).

FDA identifies reference listed drugs in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists. Listed drugs identified as reference listed drugs represent drug products upon which an applicant can rely in seeking approval of an ANDA. FDA intends to update periodically the reference listed drugs identified in the Prescription Drug Product, OTC Drug Product, and Discontinued Drug Product Lists, as appropriate.

FDA also identifies reference standards in the Prescription Drug Product and OTC Drug Product Lists. Listed drugs identified as reference standards represent the FDA's best judgment at this time as to the appropriate comparator for purposes of conducting any *in vivo* bioequivalence studies required for approval.

In some instances when FDA has not designated a listed drug as a reference listed drug, such listed drug may be shielded from generic competition. If FDA has not designated a reference listed drug for a drug product the applicant intends to duplicate, the potential applicant may ask FDA to designate a reference listed drug for that drug product. Potential applicants should consult agency guidance related to referencing approved drug products in ANDA submissions for information on submitting such a request. Section 1.7, *Therapeutic Equivalence Evaluations Codes (products meeting necessary bioequivalence requirements)* explains the character coding system (e.g., **AB**, **AB1**, **AB2**, **AB3**...) for multisource drug products listed under the same heading with two or more reference listed drugs.

A potential applicant should consult Agency guidance related to referencing approved drug products in ANDA submissions for information on submitting a request for selection of a reference standard. FDA may, on its own initiative, select a new reference standard when doing so will help to ensure that applications for generic drugs may be submitted and evaluated, e.g., in the event that the listed drug currently selected as the reference standard has been withdrawn from sale for other than safety and efficacy reasons.

If an applicant has a question related to the appropriate reference standard, it is recommended that an applicant planning to conduct an *in vivo* bioequivalence study submit a controlled correspondence to the Office of Generic Drugs.

## 1.5 General Policies and Legal Status

The Orange Book contains public information and advice. It does not mandate the drug products that are purchased, prescribed, dispensed, or substituted for one another, nor does it, conversely, mandate the products that should be avoided. To the extent that the Orange Book sets forth FDA's evaluations of the therapeutic equivalence of drug products that have been approved, it contains FDA's advice to the public, to practitioners, and to the states regarding drug product selection. These evaluations do not constitute determinations that any product is in violation of the FD&C Act or that any product is preferable to any other. Therapeutic equivalence evaluations are a scientific judgment based upon evidence, while generic substitution may involve social and economic policy administered by the states, intended to reduce the cost of drugs to consumers. To the extent that the Orange Book identifies drug products approved under Section 505 of the FD&C Act, it sets forth information that the Agency is required to publish and that the public is entitled to under the Freedom of Information Act. Exclusion of a drug product from the Orange Book does not necessarily mean that the drug product is in violation of Section 505 of the FD&C Act, that such a product is not safe or effective, or that such a product is not therapeutically equivalent to other drug products. Rather, the exclusion may

be based on the fact that FDA has not evaluated the safety, effectiveness, and quality of the drug product.

## 1.6 Practitioner/User Responsibilities

***Professional care and judgment should be exercised in using the Orange Book.*** Evaluations of therapeutic equivalence for prescription drugs are based on scientific and medical evaluations by FDA. Products evaluated as therapeutically equivalent can be expected, in the judgment of FDA, to have equivalent clinical effect and no difference in their potential for adverse effects when used under the conditions of their labeling. However, these products may differ in other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, in some instances, labeling. If products with such differences are substituted for each other, there is a potential for patient confusion due to differences in color or shape of tablets, inability to provide a given dose using a partial tablet if the proper scoring configuration is not available, or decreased patient acceptance of certain products because of flavor. For example, there may also be allergic reactions in rare cases due to a coloring or a preservative ingredient, as well as differences in cost to the patient.

FDA evaluation of therapeutic equivalence in no way relieves practitioners of their professional responsibilities in prescribing and dispensing such products with due care and with appropriate information to individual patients. In those circumstances where the characteristics of a specific product, other than its active ingredient, are important in the therapy of a particular patient, the physician's prescribing of that product may be appropriate. Pharmacists must also be familiar with the different characteristics of therapeutically equivalent products, e.g., expiration dates/times and labeling directions for storage of the different products (particularly for reconstituted products), so they can properly advise patients when one product is substituted for another.

***Multisource and single-source drug products.*** In the Orange Book, FDA has evaluated for therapeutic equivalence only multisource prescription drug products approved under Section 505 of the FD&C Act, which in most instances means those pharmaceutical equivalents available from more than one manufacturer. For such products, a therapeutic equivalence code is included and product information is highlighted in bold face and underlined. Those products with approved applications that are single-source (i.e., there is only one approved product available for that active ingredient, dosage form, route of administration, and strength) are also included in the Orange Book, but no therapeutic equivalence code is included with such products. Any drug product in the Orange Book repackaged and/or distributed by other than the applicant (e.g., an authorized generic) is considered to be therapeutically equivalent to the applicant's drug product even if the applicant's drug product is single source or coded as non-equivalent (e.g., BN). The details of these codes and the policies underlying them are discussed in Section 1.7, *Therapeutic Equivalence Evaluations Codes*. Distributors or repackagers of an applicant's drug product are not identified in the Orange Book.

***Products in the Orange Book are identified by the names of the holders of approved applications (applicants) who may not necessarily be the manufacturer of the product.*** There are numerous entities other than the applicant that may be involved in the development, manufacturing, and/or marketing of a product. The applicant may have had its product manufactured by a contract manufacturer and may simply be distributing the product for which it has obtained approval. In many instances, however, the manufacturer of the product is also the applicant. The name of the manufacturer is

permitted by regulation to appear on the label, even when the manufacturer is not the applicant or marketer.

Although the products in the Orange Book are identified by the names of the applicants, circumstances, such as changing corporate ownership, have sometimes made identification of the applicant difficult. The Agency believes, based on continuing document review and communication with firms, that the applicant designations in the Orange Book are, in most cases, correct.

To relate firm name information on a product label to that in the Orange Book, the following should be noted: the applicant's name always appears in the Orange Book. This applies whether the applicant (firm name on the Form FDA 356h in the application) is the manufacturer or marketer (firm name in largest letters on the label) or not. However, the applicant's name may not always appear on the label of the product.

If the applicant is the marketer, its name appears in the Orange Book and on the label; if the applicant is not the marketer, and the Agency is aware of a corporate relationship (e.g., parent and subsidiary) between the applicant and the marketer, the name of the applicant appears in the Orange Book and both firm names may appear on the label. Firms with known corporate relationships are displayed in Appendix B. If there is no known corporate relationship between the applicant and the marketer, the applicant's name appears in the Orange Book; however, unless the applicant is the manufacturer, packager, or distributor, the applicant's name may not appear on the label. In this case, the practitioner, from labeling alone, will not be able to relate the marketed product to an applicant cited in the Orange Book, and hence to a specific approved drug product. In such cases, to assure that the product in question is the subject of an approved application, the firm named on the label should be contacted.

To relate trade name (proprietary name) information on a product label to that in the Orange Book, the following should be noted: if the applicant is the marketer, the applicant's name appears in the Orange Book and on the label; if the Agency is aware of a corporate relationship between the applicant and the marketer, the trade name (proprietary name) of the drug product (established name of the active ingredient, if no trade name exists) appears in the Orange Book. If a corporate relationship exists between an applicant and a marketer and both firms are distributing the drug product, the FDA reserves the right to select the trade name of either the marketer or the applicant to appear in the Orange Book. If there is no known corporate relationship between the applicant and the marketer, the established drug name (i.e., non-proprietary name) appears in the Orange Book.

***Every product in the Orange Book is subject at all times to regulatory action.*** From time to time, approved products may be found in violation of one or more provisions of the FD&C Act. In such circumstances, the Agency may commence appropriate enforcement action to correct the violation, if necessary, by securing removal of the product from the market by voluntary recall, seizure, or other enforcement actions. Such regulatory actions are, however, independent of the inclusion of a product in the Orange Book. The main criterion for inclusion of a product is that it has an application that has been approved and that has not been withdrawn for safety or efficacy reasons. FDA believes that retention of a violative product in the Orange Book will not have any significant adverse health consequences, because other legal mechanisms are available to the Agency to prevent the product's actual marketing. FDA may however, change a product's therapeutic equivalence rating if the circumstances giving rise to the violation change or otherwise call into question the Agency's assessment of whether a product meets the criteria for therapeutic equivalence.

## **1.7 Therapeutic Equivalence Evaluations Codes**

Generally, drug products that the Agency considers multisource have been assigned a therapeutic equivalence code. The coding system for therapeutic equivalence evaluations is designed to allow users to determine quickly whether the Agency has evaluated a particular approved product (e.g., a particular strength of an approved drug) as therapeutically equivalent to other pharmaceutically equivalent products (first letter) and to provide additional information on the basis of FDA's evaluations (second letter). With some exceptions (e.g., therapeutic equivalence evaluations for certain 505(b)(2) applications), the therapeutic equivalence evaluation date is the same as the approval date.

The two basic categories into which multisource drugs have been placed are indicated by the first letter of the relevant therapeutic equivalence code as follows:

**A Drug products that FDA considers to be therapeutically equivalent to other pharmaceutically equivalent products, i.e., drug products for which:**

- (1) there are no known or suspected bioequivalence problems. These are designated **AA**, **AN**, **AO**, **AP**, or **AT**, depending on the dosage form; or
- (2) actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. These are designated **AB**.

**B Drug products that FDA at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products, i.e.,**

drug products for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence. Often the problem is with specific dosage forms rather than with the active ingredients. These are designated **BC**, **BD**, **BE**, **BN**, **BP**, **BR**, **BS**, **BT**, **BX**, or **B\***.

Individual drug products have been evaluated as therapeutically equivalent to the reference product in accordance with the definitions and policies outlined below:

### **"A" CODES**

**Drug products that are considered to be therapeutically equivalent to other pharmaceutically equivalent products.**

"A" products are those for which there are no known or suspected bioequivalence problems or for which actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. Drug products designated with an "A" code fall under one of two main policies:

- (1) for those active ingredients or dosage forms for which no *in vivo* bioequivalence issue is known or suspected, the information necessary to show bioequivalence between pharmaceutically equivalent products is either presumed and considered self-evident (based on other information in the application for some dosage forms (e.g., solutions)) or satisfied by a showing that an acceptable *in vitro* dissolution standard is met. A

therapeutically equivalent rating is assigned such products so long as they are manufactured in accordance with Current Good Manufacturing Practice regulations and meet the other requirements of their approved applications (these are designated **AA**, **AN**, **AO**, **AP**, or **AT**, depending on the dosage form, as described below); or

- (2) for those Drug Efficacy Study Implementation (DESI) drug products containing active ingredients or dosage forms that have been identified by FDA as having actual or potential bioequivalence problems, and for post-1962 drug products in a dosage form presenting a potential bioequivalence problem, an evaluation of therapeutic equivalence is assigned to pharmaceutical equivalents only if the approved application contains adequate scientific evidence establishing through *in vivo* and/or *in vitro* studies the bioequivalence of the product to a selected reference product (these products are designated as **AB**).

There are some general principles that may affect the substitution of pharmaceutically equivalent products in specific cases. Prescribers and dispensers of drugs should be alert to these principles so as to deal appropriately with situations that require professional judgment and discretion.

There may be labeling differences among pharmaceutically equivalent products that require attention on the part of the health professional (e.g., pharmaceutically equivalent powders to be reconstituted for administration as oral or injectable liquids may vary with respect to their expiration time or storage conditions after reconstitution). FDA's determination that such products are therapeutically equivalent is applicable only when each product is reconstituted, stored, and used under the conditions specified in its labeling.

The Agency may use notes in this publication to point out special situations, such as potential differences between two drug products that have been evaluated as bioequivalent and otherwise therapeutically equivalent, when they should be brought to the attention of health professionals. These notes are contained in Section 1.8, *Description of Certain Special Situations*.

For example, in certain instances, there may be variations among pharmaceutically equivalent products in their use or in conditions of administration. Such differences may be due to patent or exclusivity rights associated with such use. When such variations may, in the Agency's opinion, affect prescribing or substitution decisions by health professionals, a note may be added to Section 1.8.

Also, occasionally a situation may arise in which changes in a listed drug product after its approval (for example, a change in dosing interval) may have an impact on the substitutability of already approved generic versions of that product that were rated by the Agency as therapeutically equivalent to the listed product. When such changes in the listed drug product are considered by the Agency to have a significant impact on therapeutic equivalence, the Agency will change the therapeutic equivalence ratings for other versions of the drug product unless the manufacturers of those other versions of the product provide additional information to assure equivalence under the changed conditions. Pending receipt of the additional data, the Agency may add a note to Section 1.8, or, in rare cases, may even change the therapeutic equivalence rating.

In some cases (e.g., Isolyte® S w/ Dextrose 5% in Plastic Container and Plasma-Lyte® 148 and Dextrose 5% in Plastic Container), closely related products are listed as containing the same active ingredients, but in

somewhat different amounts. In determining which of these products are pharmaceutically equivalent, generally the Agency has considered products to be pharmaceutically equivalent with labeled strengths of an ingredient that do not vary by more than 1%.

Different salts, esters or other noncovalent derivatives (such as a complex, chelate, or clathrate) of the same active moiety are regarded as different active ingredients. For the purpose of this publication, products containing such different active ingredients are considered pharmaceutical alternatives and, thus, not therapeutically equivalent. Anhydrous and hydrated entities, as well as different polymorphs, are considered to be the same active ingredient and are expected to meet the same standards for identity to be considered pharmaceutical equivalents and therapeutic equivalents.

The codes in this book are not intended to preclude health care professionals from converting pharmaceutically different concentrations into pharmaceutical equivalents using accepted professional practice.

Where package size variations have therapeutic implications, products so packaged have not been considered pharmaceutically equivalent. For example, some oral contraceptives are supplied in 21-tablet and 28-tablet packets; the 28-tablet packets contain 7 placebo or iron tablets. These two packaging configurations are not regarded as pharmaceutically equivalent; thus, they are not designated as therapeutically equivalent.

Preservatives and other inactive ingredients may differ among some therapeutically equivalent drug products. These differences do not affect FDA's evaluation of therapeutic equivalence except in cases where these components may influence bioequivalence or routes of administration.

The specific sub-codes for those drugs evaluated as therapeutically equivalent and the policies underlying these sub-codes follow:

## **AA Products in conventional dosage forms not presenting bioequivalence problems**

Multisource drug products coded as **AA** contain active ingredients and are in dosage forms that are not regarded as presenting either actual or potential bioequivalence problems or drug quality or standards issues. However, all oral dosage forms must, nonetheless, meet an appropriate *in vitro* bioequivalence standard that is acceptable to the Agency in order to be approved.

## **AB, AB1, AB2, AB3... Products meeting necessary bioequivalence requirements**

Multisource drug products listed under the same heading (i.e., identical active ingredients(s), dosage form, and route(s) of administration) and having the same strength (see Section 1.2, *Therapeutic Equivalence-Related Terms, Strength*) generally will be coded **AB** if data and information are submitted demonstrating bioequivalence.

In certain instances, a number is added to the end of the **AB** code to make a three character code (i.e., **AB1**, **AB2**, **AB3**, etc.). Three-character codes generally are assigned only in situations when more than one reference listed drug of the same strength has been designated under the same heading. Two or more reference listed drugs are generally selected only when there are at least two potential reference listed drug products that are not identified as bioequivalent to each other. If a study is submitted that demonstrates bioequivalence to a specific listed drug product, the generic product will be given the same three-character code

as the reference listed drug it was compared against. For example, Adalat® CC and Procardia XL®, extended-release tablets, are listed under the active ingredient nifedipine. These drug products, listed under the same heading, are not bioequivalent to each other. Adalat® CC and Procardia XL® have been assigned ratings of **AB1** and **AB2**, respectively. Generic drug products deemed by FDA to be bioequivalent to Adalat® CC and Procardia XL® have been approved. As a result, the generic drug products bioequivalent to Adalat® CC have been assigned a rating of **AB1** and those bioequivalent to Procardia XL® have been assigned a rating of **AB2**. (The assignment of an **AB1** or **AB2** rating to a specific product does not imply product preference.) Even though drug products of distributors and/or repackagers are not included in the Orange Book, they are considered therapeutically equivalent to the applicant's drug product if the applicant's drug product is rated either with an **AB** or three-character code or is single source in the Orange Book. Drugs coded as **AB** under a heading are considered therapeutically equivalent only to other drugs coded as **AB** under that heading. Drugs coded with a three-character code under a heading are considered therapeutically equivalent only to other drugs coded with the same three-character code under that heading.

#### **AN Solutions and powders for aerosolization**

Uncertainty regarding the therapeutic equivalence of aerosolized products arises primarily because of differences in the drug delivery system. Solutions and powders intended for aerosolization that are marketed for use in general-use delivery systems are considered to be pharmaceutically and therapeutically equivalent and are coded **AN**. Those products that are compatible only with a specific delivery system or those products that are packaged in and with a specific delivery system are coded **BN**, unless they have met an appropriate bioequivalence standard and are otherwise determined to be therapeutically equivalent. Solutions or suspensions in a specific delivery system will be coded **AN** if the bioequivalence standard is based upon *in vitro* methodology, if bioequivalence needs to be demonstrated by *in vivo* methodology then the drug products will be coded **AB**.

#### **AO Injectable oil solutions**

The absorption of drugs in injectable (parenteral) oil solutions may vary substantially with the type of oil employed as a vehicle and the concentration of the active ingredient. Injectable oil solutions are therefore considered to be pharmaceutically and therapeutically equivalent only when the active ingredient, its concentration, and the type of oil used as a vehicle are all identical.

#### **AP Injectable aqueous solutions and, in certain instances, intravenous non-aqueous solutions**

It should be noted that even though injectable (parenteral) products under a specific listing may be evaluated as therapeutically equivalent, there may be important differences among the products in the general category, Injectable; Injection. For example, historically some injectable products that are rated therapeutically equivalent are labeled for different routes of administration. In addition, some products evaluated as therapeutically equivalent may have different preservatives or no preservatives at all. Injectable products available as dry powders for reconstitution, concentrated sterile solutions for dilution, or sterile solutions ready for injection are pharmaceutical alternative drug

products. They are not rated as therapeutically equivalent (AP) to each other even if these pharmaceutical alternative drug products are designed to produce the same concentration prior to injection and are similarly labeled. Consistent with accepted professional practice, it is the responsibility of the prescriber, dispenser, or individual administering the product to be familiar with a product's labeling to assure that it is given only by the route(s) of administration stated in the labeling.

Certain commonly used large volume intravenous products in glass containers are not included in the Orange Book (e.g., dextrose injection 5%, dextrose injection 10%, sodium chloride injection 0.9%) since these products are on the market without FDA approval and the FDA has not published conditions for marketing such parenteral products under approved NDAs. When packaged in plastic containers, however, FDA regulations require approved applications prior to marketing. Approval then depends on, among other things, the extent of the available safety data involving the specific plastic component of the product. All large volume parenteral products are manufactured under similar standards, regardless of whether they are packaged in glass or plastic. Thus, FDA has no reason to believe that the packaging container of large volume parenteral drug products that are pharmaceutically equivalent would have any effect on their therapeutic equivalence.

Consistent with the definition of strength included in Section 1.2, *Therapeutic Equivalence-Related Terms*, the strength of parenteral drug products generally is identified by both the total drug content and the concentration of drug substance in a container approved by FDA.<sup>8</sup> In the past, the strength of liquid parenteral drug products in the Orange Book has not been fully displayed. Rather, the strength of liquid parenteral drug products in the Orange Book has been displayed in terms of concentration, expressed as xmg/mL. Generally, the amount of dry powder or lyophilized powder in a container is identified as the strength, expressed as xmg/vial.

After the enactment of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA), which amended the FD&C Act, it became evident that the format of the Orange Book with respect to parenteral solutions should be changed to reflect that each strength of a drug is considered to be a separate listed drug. The Orange Book now displays the strength of all new approvals of parenteral solutions. Previously, we would have displayed only the concentration of an approved parenteral solution, e.g. 50mg/mL. If this application had a 20 mL and 60 mL container approved, we would now display two product strengths, listing both total drug content and concentration of drug substance in the relevant approved container, e.g. 1gm/20mL (50mg/mL) and 3gm/60mL (50mg/mL).

## **AT Topical products**

There are a variety of topical dosage forms available for dermatologic, ophthalmic, otic, rectal, and vaginal administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays and suppositories. Even though different topical dosage forms may contain the same active ingredient and potency, these dosage forms are not considered pharmaceutically equivalent. Therefore, they are not considered therapeutically equivalent. All solutions and DESI drug products

---

<sup>8</sup> The strengths of certain parenteral drug products, including contrast agents, may be expressed as a percentage.

containing the same active ingredient in the same topical dosage form for which a waiver of *in vivo* bioequivalence has been granted and for which chemistry and manufacturing processes are adequate to demonstrate bioequivalence, are considered therapeutically equivalent and coded **AT**. Pharmaceutically equivalent topical products that raise questions of bioequivalence, including all post-1962 non-solution topical drug products, are coded **AB** when supported by adequate bioequivalence data, and **BT** in the absence of such data.

### **"B" CODES**

**Drug products that FDA, at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products.**

"**B**" products, for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence, often have a problem with specific dosage forms rather than with the active ingredients. Drug products designated with a "**B**" code fall under one of three main policies:

- (1) the drug products contain active ingredients or are manufactured in dosage forms that have been identified by the Agency as having documented bioequivalence problems or a significant potential for such problems and for which no adequate studies demonstrating bioequivalence have been submitted to FDA; or
- (2) the quality standards are inadequate or FDA has an insufficient basis to determine therapeutic equivalence; or
- (3) the drug products are under regulatory review.

The specific coding definitions and policies for the "**B**" sub-codes are as follows:

#### **B\* Drug products requiring further FDA investigation and review to determine therapeutic equivalence**

The code **B\*** is assigned to products previously assigned an **A** or **B** code when FDA receives new information that raises a significant question regarding therapeutic equivalence that can be resolved only through further Agency investigation and/or review of data and information submitted by the applicant. The **B\*** code signifies that the Agency will take no position regarding the therapeutic equivalence of the product until the Agency completes its investigation and review.

#### **BC Extended-release dosage forms (capsules, injectables and tablets)**

Extended-release tablets are formulated in such a manner as to make the contained drug substance available over an extended period of time following ingestion.

Although bioavailability studies have been conducted on these dosage forms, they may be subject to bioavailability differences, primarily because applicants developing extended-release products for the same active ingredient rarely employ the same formulation approach. FDA, therefore, does not consider different extended-release dosage forms containing the same active ingredient in equal strength to be therapeutically equivalent unless equivalence between individual products

in both rate and extent has been specifically demonstrated through appropriate bioequivalence studies. Extended-release products for which such bioequivalence data have not been submitted are coded **BC**, while those for which such data are available have been coded **AB**.

#### **BD Active ingredients and dosage forms with documented bioequivalence problems**

The **BD** code denotes products containing active ingredients with known bioequivalence problems and for which adequate studies have not been submitted to FDA demonstrating bioequivalence. Where studies showing bioequivalence have been submitted, the product has been coded **AB**.

#### **BE Delayed-release oral dosage forms**

Where the drug may be destroyed or inactivated by the gastric juice or where it may irritate the gastric mucosa, the use of "enteric" coatings is indicated. Such coatings are intended to delay the release of the medication until the tablet has passed through the stomach. Drug products in delayed-release dosage forms containing the same active ingredients are subject to significant differences in absorption. Unless otherwise specifically noted, the Agency considers different delayed-release products containing the same active ingredients as presenting a potential bioequivalence problem and codes these products **BE** in the absence of *in vivo* studies showing bioequivalence. If adequate *in vivo* studies have demonstrated the bioequivalence of specific delayed-release products, such products are coded **AB**.

#### **BN Products in aerosol-nebulizer drug delivery systems**

This code applies to drug solutions or powders that are marketed only as a component of, or as compatible with, a specific drug delivery system. There may, for example, be significant differences in the dose of drug and particle size delivered by different products of this type. Therefore, the Agency does not consider different metered aerosol dosage forms containing the same active ingredient(s) in equal strengths to be therapeutically equivalent unless the drug products meet an appropriate bioequivalence standard; such products are coded **AB**.

#### **BP Active ingredients and dosage forms with potential bioequivalence problems**

FDA's bioequivalence regulations (21 CFR 320.33) contain criteria and procedures for determining whether a specific active ingredient in a specific dosage form has a potential for causing a bioequivalence problem. It is FDA's policy to consider an ingredient meeting these criteria as having a potential bioequivalence problem even in the absence of positive data demonstrating inequivalence. Pharmaceutically equivalent products containing these ingredients in oral dosage forms are coded **BP** until adequate bioequivalence data are submitted, after which such products are coded **AB**. Injectable suspensions containing an active ingredient suspended in an aqueous or oleaginous vehicle have also been coded **BP**. Injectable suspensions are subject to bioequivalence problems because differences in particle size, polymorphic structure of the suspended active ingredient, or the suspension formulation can significantly affect the rate of release and absorption. FDA does not consider pharmaceutical equivalents of these products bioequivalent without adequate evidence of bioequivalence; such products would be coded **AB**.

## **BR Suppositories or enemas that deliver drugs for systemic absorption**

The absorption of active ingredients from suppositories or enemas that are intended to have a systemic effect (as distinct from suppositories administered for local effect) can vary significantly from product to product. Therefore, FDA considers pharmaceutically equivalent systemic suppositories or enemas bioequivalent only if *in vivo* evidence of bioequivalence is available. In those cases where *in vivo* evidence is available, the products are coded **AB**. If such evidence is not available, the products are coded **BR**.

## **BS Products having drug standard deficiencies**

If the drug standards for an active ingredient in a particular dosage form are found by FDA to be deficient so as to prevent an FDA evaluation of either pharmaceutical or therapeutic equivalence, all drug products containing that active ingredient in that dosage form are coded **BS**. For example, if the standards permit a wide variation in pharmacologically active components of the active ingredient such that pharmaceutical equivalence is in question, all products containing that active ingredient in that dosage form are coded **BS**.

## **BT Topical products with bioequivalence issues**

This code applies mainly to post-1962 dermatologic, ophthalmic, otic, rectal, and vaginal products for topical administration, including creams, ointments, gels, lotions, pastes, and sprays, as well as suppositories not intended for systemic drug absorption. Topical products evaluated as having acceptable clinical performance, but that are not bioequivalent to other pharmaceutically equivalent products or that lack sufficient evidence of bioequivalence, will be coded **BT**.

## **BX Drug products for which the data are insufficient to determine therapeutic equivalence**

The code **BX** is assigned to specific drug products for which the data that have been reviewed by the Agency are insufficient to determine therapeutic equivalence under the policies stated in this document. In these situations, the drug products are presumed to be therapeutically inequivalent until the Agency has determined that there is adequate information to make a full evaluation of therapeutic equivalence.

## **1.8 Description of Certain Special Situations**

Certain drugs listed in the Orange Book present special situations that merit further discussion. The following are descriptions of certain examples of those special situations:

**Amino Acid and Protein Hydrolysate Injections.** These products differ in the amount and kinds of amino acids they contain and, therefore, are not considered pharmaceutical equivalents. For this reason, these products are not considered therapeutically equivalent. At the same time, the Agency believes that it is appropriate to point out that where nitrogen balance is the sole therapeutic objective and individual amino acid content is not a consideration, pharmaceutical alternatives with the same total amount of nitrogen content may be considered therapeutically equivalent.

**Gaviscon®.** Gaviscon® is an OTC product that has been marketed since September 1970. The active ingredients in this product, aluminum hydroxide and magnesium trisilicate, were reviewed by the Agency's OTC Antacid Panel and were considered to be safe and effective ingredients (Category I) by that Panel. However, the tablet failed to pass the antacid test that is required of all antacid products. The Agency, therefore, placed the tablet in Category III for lack of effectiveness. A full NDA with clinical studies was submitted by Marion Laboratories, Inc., and approved by FDA on December 9, 1983. Gaviscon®'s activity in treating reflux acidity is made possible by the physical-chemical properties of the inactive ingredients, sodium bicarbonate and alginic acid. Therefore, *all ANDAs that cite Gaviscon® tablets as the reference listed drug must contain the inactive ingredients sodium bicarbonate and alginic acid.* A full NDA will be required to support the effectiveness of the drug product if different inactive ingredients are to be substituted for sodium bicarbonate or alginic acid or if different proportions of these ingredients are to be used.

**Levothyroxine Sodium.** Because there are multiple reference listed drugs of levothyroxine sodium tablets and some reference listed drugs' sponsors have conducted studies to establish their drugs' therapeutic equivalence to other reference listed drugs, FDA has determined that its usual practice of assigning two or three character TE codes may be potentially confusing and inadequate for these drug products. Accordingly, FDA provides the following explanation and chart of therapeutic equivalence evaluations for levothyroxine sodium tablet drug products.

Levothyroxine Sodium (Mylan ANDA 076187), Levoxyl (King Pharms NDA 021301), Synthroid (AbbVie NDA 021402), and Levo-T (CEDIPROF NDA 021342) tablets have been determined to be therapeutically equivalent to corresponding strengths of Unithroid (Jerome Stevens NDA 021210) tablets.

Levo-T (CEDIPROF NDA 021342), Levothyroxine Sodium (Mylan ANDA 076187), and Unithroid (Jerome Stevens NDA 021210) tablets have been determined to be therapeutically equivalent to corresponding strengths of Synthroid (AbbVie NDA 021402) tablets.

Levo-T (CEDIPROF NDA 021342), Unithroid (Jerome Stevens NDA 021210), and Levothyroxine Sodium (Mylan ANDA 076187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levoxyl (King Pharms NDA 021301) tablets.

Levothyroxine Sodium (Mylan ANDA 076187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Thyro-Tabs (Lloyd NDA 021116) tablets.<sup>9</sup>

The chart outlines TE codes for all 0.025 mg products in the active section of the Orange Book. Other product strengths may be similar. Therapeutic equivalence has been established between products that have the same AB+number TE code. More than one TE code may apply to some products. One

---

<sup>9</sup> Lloyd's Thyro-Tabs tablets (NDA 021116) (previously known as Levothroid) is currently listed in the Discontinued Drug Product List section of the Orange Book and Mylan's levothyroxine product (ANDA 076187) has been selected as the reference standard for ANDA applicants to use to establish bioequivalence to Thyro-Tabs. If an ANDA that uses Mylan's levothyroxine product as its reference standard is approved, the ANDA will receive an AB4 rating. The ANDA applicant also may obtain an AB rating for its product to the other reference listed drugs (i.e., Unithroid, Synthroid, and Levoxyl) by submitting supplements that demonstrate that the generic product is bioequivalent to these other reference listed drugs and satisfies all other therapeutic equivalence criteria with respect to these reference listed drugs. See Letter from Janet Woodcock, M.D., Director, Center for Drug Evaluation and Research, FDA to Teri Nataline, Principal Consultant, Lachman Consultant Services, Inc., Docket No. FDA-2015-P-0403 (May 27, 2016).

common TE code indicates therapeutic equivalence between products.

Trade Name	Applicant	Strength	TE Code	Appl No	Product No
UNITHROID	STEVENS J	0.025MG	AB1	021210	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB1	076187	001
LEVOXYL	KING PHARMS	0.025MG	AB1	021301	001
SYNTHROID	ABBVIE	0.025MG	AB1	021402	001
LEVO-T	CEDIPROF INC	0.025MG	AB1	021342	001
SYNTHROID	ABBVIE	0.025MG	AB2	021402	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB2	076187	001
LEVO-T	CEDIPROF INC	0.025MG	AB2	021342	001
UNITHROID	STEVENS J	0.025MG	AB2	021210	001
LEVOXYL	KING PHARMS	0.025MG	AB3	021301	001
LEVO-T	CEDIPROF INC	0.025MG	AB3	021342	001
UNITHROID	STEVENS J	0.025MG	AB3	021210	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB3	076187	001
THYRO-TABS	LLOYD	0.025MG	N/A <sup>10</sup>	021116	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB4	076187	001

**Patent Certification(s) and Reference Standard for ANDAs Duplicating a Drug Product Approved in a Petitioned ANDA.** To submit an ANDA for a generic drug that is not the same as its reference listed drug because it has one different active ingredient (in a fixed combination drug product), or has a different route of administration, dosage form, or strength than that of the reference listed drug, an applicant first must obtain permission from FDA through what is known as a suitability petition pursuant to section 505(j)(2)(C) of the FD&C Act. A petitioned ANDA relies on the reference listed drug described in the suitability petition. An ANDA for a drug that is the same as a drug product approved in a petitioned ANDA should utilize the drug product approved in the petitioned ANDA as a reference standard. However, the reference listed drug for any such ANDA is generally the listed drug referenced in the approved suitability petition. The ANDA must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug that served as the basis for the approved suitability petition.<sup>11</sup> (This concept also generally applies to an ANDA applicant that utilizes a reference standard that is not a reference listed drug, as such an application must include appropriate patent certification(s) and an exclusivity statement with respect to the reference listed drug.)

**Waived exclusivity.** If an NDA submitted under Section 505(b) of the FD&C

<sup>10</sup> Thryo-Tabs is in the Discontinued Drug Product List and therefore no longer is assigned a TE code.

<sup>11</sup> If after approval of a suitability petition and before approval of an ANDA submitted pursuant to the approved petition, a drug product is approved in an NDA for the change described in the petition, the suitability petition and the listed drug identified in the petition can no longer be the basis of submission for such ANDA. Under these circumstances, an applicant seeking approval for a drug product with the change approved in the suitability petition must submit a new ANDA that identifies the drug product approved under such NDA as the RLD and comply with applicable regulatory requirements. See 21 CFR 314.93(f)(2).

Act qualifies for exclusivity under the FD&C Act, the exclusivity is generally listed in the Patent and Exclusivity Section of the Orange Book. If a drug product has received this exclusivity, the FDA will not accept for review and/or will not approve a 505(b)(2) application or an ANDA under Section 505(j) of the FD&C Act, as applicable, in accordance with the relevant exclusivity. If the listed drug is also protected by one or more patents, the approval date for the ANDA or 505(b)(2) application will be determined based on an analysis of the applicant's patent certification(s) or statement(s) for each relevant patent and the effect of relevant exclusivity listed in the Orange Book. However, the holder of the NDA may waive its exclusivity as to any or all ANDAs and 505(b)(2) applications that might otherwise be blocked by such exclusivity. If an NDA sponsor waives its right to the exclusivity protection, qualified ANDAs or 505(b)(2) applications may be accepted for review and/or approved, as applicable, pursuant to the NDA holder's exclusivity being waived. An NDA for which the holder has waived its exclusivity as to all ANDAs and 505(b)(2) applications will be coded with a "W" in the Patent and Exclusivity Section of the Orange Book. The applicant whose product might otherwise be blocked by this exclusivity should indicate in the exclusivity statement in its application that the holder of the listed drug has waived its exclusivity.

## **1.9 Therapeutic Equivalence Code Change for a Category of Multisource Drug Products**

The Agency will use the following procedures when, in response to a petition or on its own initiative, it is considering a change in the therapeutic equivalence code for approved multisource drug products. Such changes will generally occur when the Agency becomes aware of new scientific information affecting the therapeutic equivalence of an entire category of multisource drug products in the Orange Book (e.g., information concerning the active ingredient or the dosage form), rather than information concerning a single drug product within the category. These procedures will be used when a change in therapeutic equivalence code is under consideration for all drug products found in the Prescription Drug Product List under a specific drug entity and dosage form. The change may be from the code signifying that the drug does not present a bioequivalence problem (e.g., **AA**) to a code signifying an actual or potential bioequivalence problem (e.g., **BP**), or vice versa. This procedure does not apply to a change of a particular product code (e.g., a change from **BP** to **AB** or from **AB** to **BX**).

Before making a change in a therapeutic equivalence code for an entire category of multisource drug products as described above, the Agency will announce in the *Introduction to the Cumulative Supplement* that it is considering the change and will invite comments. Comments, along with scientific data, may be sent to the Director, Office of Bioequivalence, Office of Generic Drugs, Center for Drug Evaluation and Research, HFD-650, 7620 Standish Place, Rockville, MD 20855.

The comment period will generally be 60 days in length, and the closing date for comments will be listed in the description of the proposed change for each drug entity.

The most useful type of scientific data submitted to support comments is an *in vivo* bioavailability/bioequivalence study conducted on batches of the subject drug products. Comments including scientific data from an *in vivo* bioavailability/bioequivalence study should present a full description of the analytical procedures and equipment used, a validation of the analytical methodology, including the standard curve, a description of the method of calculating results, and a description of the pharmacokinetic and statistical models used in analyzing the data. Anecdotal or testimonial information is the least useful to the Agency, and submission of comments based on such

information is discouraged. However, when there is supporting published or unpublished scientific literature, copies should be submitted with comments.

## **1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product**

The procedure described in Section 1.9 does not apply to a change in a single drug product code. For example, a change in a single drug product's code from **BP** to **AB** as a result of the submission of an acceptable bioequivalence study ordinarily will not be the subject of notice and comment in the Cumulative Supplement. Likewise, a change in a single drug product's code from **AB** to **BX** (e.g., as a result of new information raising a significant question as to bioequivalence) does not require notice and comment. The Agency's responsibility to provide the public with the Agency's most current information related to therapeutic equivalence may require a change in a drug product's code prior to any formal notice and opportunity for the applicant to be heard. The publication in the *Federal Register* of a proposal to withdraw approval of a drug product will ordinarily result in a change in a product's code from **AB** to **BX** if this action has not already been taken.

We recognize that certain drug products approved in 505(b)(2) applications may not have therapeutic equivalence codes, and that FDA may undertake therapeutic equivalence evaluations with respect to such drug products. A person seeking to have a therapeutic equivalence rating for a drug product approved in a 505(b)(2) application may petition the Agency through the citizen petition procedure (see 21 CFR 10.25(a) and 21 CFR 10.30).

## **1.11 Discontinued Section**

Those drug products in the discontinued section of the Orange Book (Discontinued Drug Product List) for which a determination has been made that the products were not withdrawn for safety or effectiveness reasons have been annotated with a footnote following the product strength: "Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*". The determinations listed in Orange Book are only reflective of determinations made since 1995 and published in the Federal Register. The identification of these drug products in the Discontinued Drug Product List should avoid the submission of multiple citizen petitions requesting a determination for the same drug product.

Generally, approved products are added to the Discontinued Drug Product List when the applicant notifies the Orange Book staff of the products' not-marketed status. Products may also be added to the Discontinued Drug Product List if annual reports or other submissions to the Agency indicate the product is not being marketed or as a result of other Agency administrative actions.<sup>12</sup> Changes to the Orange Book are not affected by the drug registration and listing requirements of Section 510 of the FD&C Act.

## **1.12 Changes to the Orange Book**

Every effort is made to ensure the Annual Edition is current and accurate. Applicants are requested to inform the FDA Orange Book Staff of any changes or corrections, including any change in a product's marketing status that would result in the product being moved to the Discontinued Drug

---

<sup>12</sup> See, e.g., Section 506I(d) of the FD&C Act.

Product List. FDA notes that under Section 506I(a) of the FD&C Act, applicants must notify the Agency in writing 180 days prior to withdrawing a drug product from sale, or if 180 days is not practicable, not later than the date of withdrawal from sale. Furthermore, Section 506I(b) of the FD&C Act requires that applicants notify the Agency in writing within 180 days of approval of a drug product if such drug product will not be available for sale within 180 days of approval. A request to include a newly approved product in the Discontinued Drug Product List, rather than parts 1, 2 or 3 of the Orange Book (as discussed in Section 1.1), must be submitted to the Orange Book staff by the end of the month in which the product is approved to ensure that the product is not included in the "active" portions of the next published Orange Book update.

In addition, FDA Orange Book Staff generally will act on requests to change a proprietary name for a listed drug only after approval of a supplement for the relevant change in proprietary name. To the extent that conventions for describing product identification information (i.e., active ingredients, dosage forms, routes of administration, product names, applicants, strengths) evolve over time, the Agency generally does not intend to revise such information for drug products already included in the Orange Book, but rather intends to apply the change prospectively to drug products added to the Orange Book.

You can contact the Orange Book Staff by email at [orangebook@fda.hhs.gov](mailto:orangebook@fda.hhs.gov). If you do not have access to email, you can contact the Orange Book Staff by mail at:

FDA/CDER Orange Book Staff  
Office of Generic Drug Policy  
Office of Generic Drugs  
7620 Standish Place  
Rockville, MD 20855-2773

### **1.13 Availability of the Edition**

Commencing with the 25<sup>th</sup> edition, the Annual Edition and current monthly Cumulative Supplement are available in a Portable Document Format (PDF) at the [Orange Book](#) home page by clicking on Publications. The PDF annual format duplicates previous paper versions except for the Orphan Products Designations and Approvals List. An annual subscription of the PDF format may be obtained from the U.S. Government Publishing Office, 866-512-1800.

## 2. HOW TO USE THE DRUG PRODUCT LISTS

### 2.1 Key Sections for Using the Drug Product Lists

This publication contains illustrations, along with Drug Product Lists, indices, and lists of abbreviations and terms which facilitate their use.

**Illustrations.** The annotated *Drug Product Illustration*, see Section 2.2, and the *Therapeutic Equivalence Evaluations Illustration*, see Section 2.3, are offered to provide further clarification. These depict the format found in the Prescription Drug Product List (the only list in which therapeutic equivalence evaluation codes are displayed).

**Drug Product Lists.** The Prescription and OTC Drug Product Lists, arranged alphabetically by active ingredient(s), contain product identification information (active ingredients, dosage forms, routes of administration, product names, applicants, strengths) for single and multiple ingredient drug products. Also shown are the application number and drug product number (FDA internal computer data use only) and approval dates for those drug products approved on or after January 1, 1982. The application number preceded by "N" is a New Drug Application (NDA or commonly the innovator). The application number preceded by an "A" is an Abbreviated New Drug Application (ANDA or commonly the generic).

The Discontinued Drug Product List, arranged alphabetically by active ingredient(s), contains product identification information (dosage form, product name, strength, and application number).

If a prescription drug product is available from more than one source (multisource), a therapeutic equivalence code will appear in front of the applicant's name. If a product is therapeutically equivalent to one or more products or to an appropriate reference, it will be designated with a code beginning with "A" and the entry will be underlined and printed in bold font for emphasis.

Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For purposes of this publication, this alphabetical sort takes precedence over United States Pharmacopeia official monograph order (i.e., Reserpine, Hydralazine Hydrochloride, Hydrochlorothiazide). For example, product information labeled as Reserpine, Hydrochlorothiazide and Hydralazine Hydrochloride appears under the active ingredient heading *Hydralazine Hydrochloride; Hydrochlorothiazide; Reserpine*. A cross-reference to the product information (for prescription and OTC products) appears for each additional active ingredient in the product. For combination drug products, the ingredient strengths are separated by semicolons and appear in the same relative sequence as the ingredients in the heading. Available strengths of the dosage form from an applicant appear on separate lines.

To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if necessary. Then, find the ingredient in the applicable Drug Product List. Proceed to the dosage form and route of administration and compare products within that ingredient heading only. Therapeutic equivalence or inequivalence for prescription products is determined on the basis of the therapeutic equivalence codes provided within that specific dosage form and route heading. The OTC Drug Product List, Discontinued Drug Product List, and

Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List have their data arranged similarly.

The Discontinued Drug Product List contains approved products that have never been marketed, have been discontinued from marketing and we have not determined that they were withdrawn for safety or effectiveness reasons, are for military use, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing. All products having a "@" in the December Cumulative Supplement of the previous Edition List have been added to the Discontinued Drug Product List appearing in this Edition. In addition, approved drug products that are not in the commercial distribution channel e.g., approved drug products in applications for export only are also listed in the Discontinued Drug Product List.

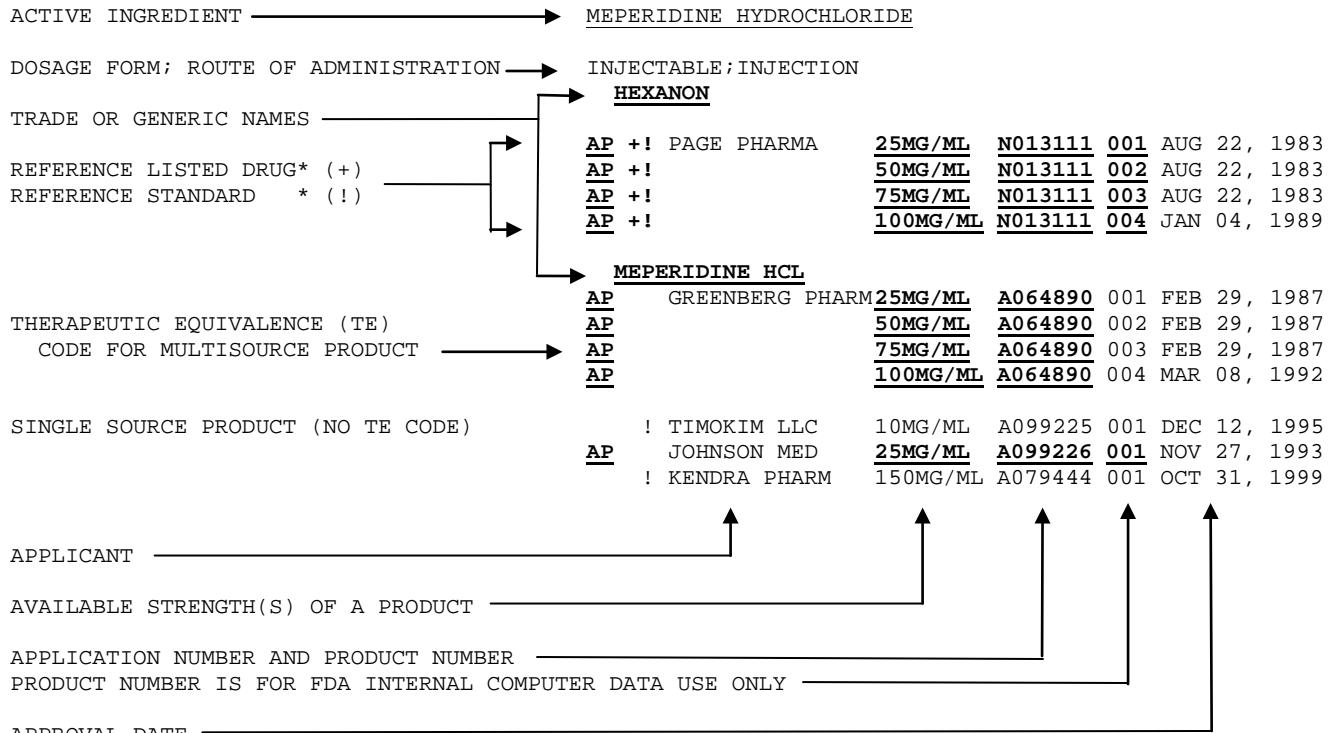
**Product Name Index** (*Prescription and OTC Drug Product Lists*). This is an index of drug products by trade name or established name of the active ingredient, if no trade name exists. The second term of each entry indicates the active ingredient name under which product information can be found in the appropriate Drug Product List. For those drug products with multiple active ingredients, only the first active ingredient (in alphabetical order) will appear. OTC products are so designated.

**Product Name Index Listed by Applicant** (*Prescription and OTC Drug Product Lists*). This is an index that cross-references applicants to drug products. The bolded and underlined entry represents the applicant name abbreviation used in this publication. Each complete applicant name that is represented by the abbreviated name is marked with an asterisk (\*). Listed under each complete applicant name is the first alphabetically arranged ingredient under which product information can be found in the appropriate Drug Product List. OTC products are so designated. To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if appropriate.

**Uniform Terms.** To improve readability, uniform terms are used to designate dosage forms, routes of administration, and abbreviations used to express strengths. These terms are listed in Appendix C. In some cases, the terms used may differ from those used in product labels and other labeling.

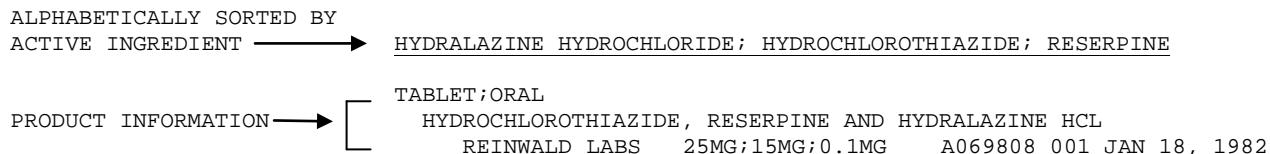
## 2.2 DRUG PRODUCT ILLUSTRATION

### SINGLE INGREDIENT



\*NOTE: REFERENCE LISTED DRUG AND REFERENCE STANDARD ARE DISCUSSED IN THE PREFACE SECTION 1.4

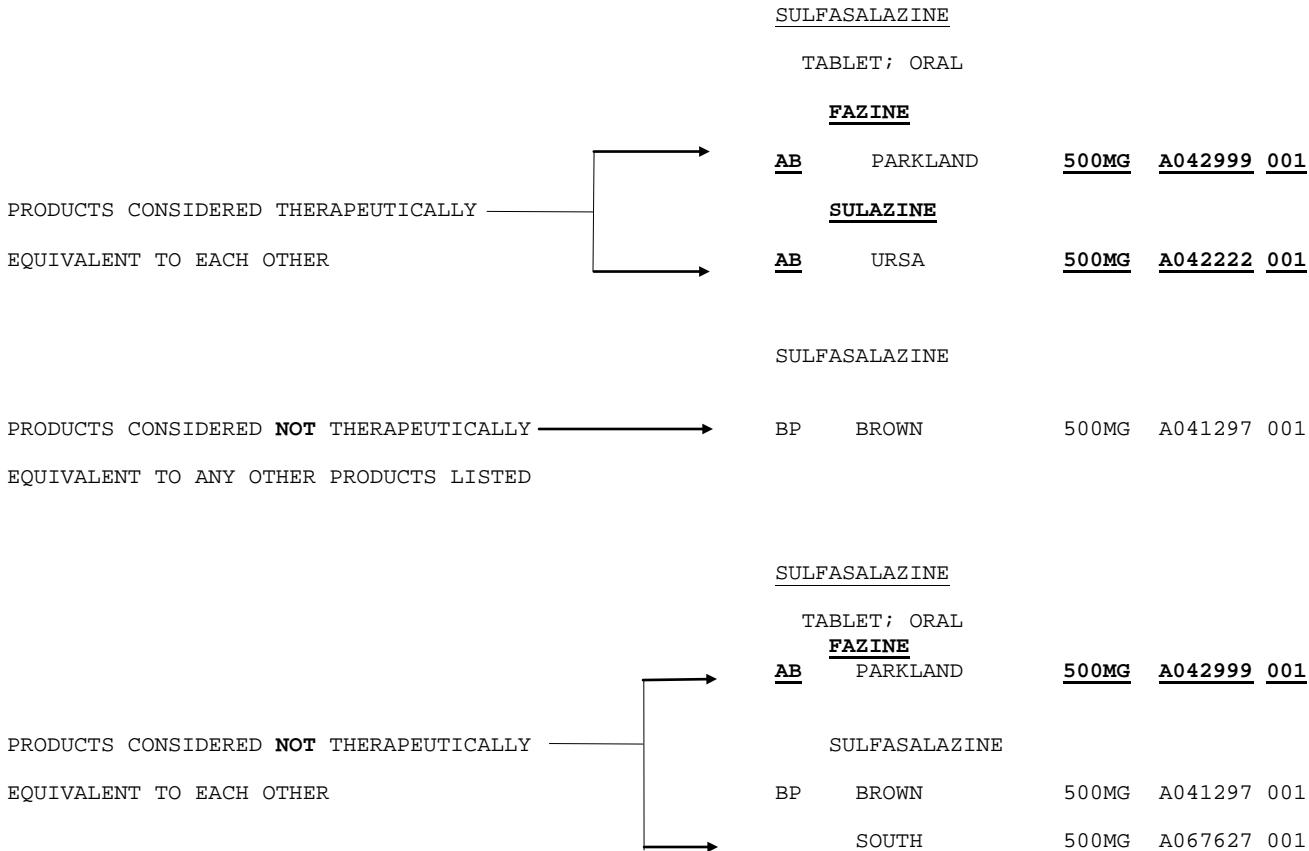
### MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION



THIS EXAMPLE IS FOR PURPOSE OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

## 2.3 THERAPEUTIC EQUIVALENCE EVALUATIONS ILLUSTRATION

DRUG PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "**A**") UNDER AN INGREDIENT AND DOSAGE FORM HEADING ARE CONSIDERED THERAPEUTICALLY EQUIVALENT ONLY TO OTHER PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "**A**") AND **NOT** TO THOSE CODED **BP** (OR ANY CODE BEGINNING WITH "**B**") AND ANY PRODUCTS NOT LISTED. DRUG PRODUCTS CODED **BP** (OR ANY CODE BEGINNING WITH A "**B**") ARE **NOT** CONSIDERED THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCT. FOR A COMPLETE EXPLANATION OF THE **TE** CODES REFER TO SECTION 1.7 OF THE INTRODUCTION.



NOTE: BOLD FONT AND UNDERLINING DENOTES MULTISOURCE PRODUCTS WHICH ARE CONSIDERED THERAPEUTICALLY EQUIVALENT.

THIS EXAMPLE IS FOR PURPOSES OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-1 (of 436)

**ABACAVIR SULFATE**

SOLUTION;ORAL

**ABACAVIR SULFATE**

**AA** HETERO LABS LTD III **EQ 20MG BASE/ML**

**A201107 001** Sep 26, 2016

**ZIAGEN**

**AA** +! VIIV HLTHCARE **EQ 20MG BASE/ML**

**N020978 001** Dec 17, 1998

TABLET;ORAL

**ABACAVIR SULFATE**

**AB** APOTEX INC **EQ 300MG BASE**

**A201570 001** Dec 17, 2012

**AB** AUROBINDO PHARMA LTD

**A077844 001** Dec 17, 2012

**AB** CIPLA LTD **EQ 300MG BASE**

**A078119 001** Nov 21, 2017

**AB** HETERO LABS LTD III **EQ 300MG BASE**

**A091560 001** Sep 13, 2013

**AB** MYLAN PHARMS INC **EQ 300MG BASE**

**A091294 001** Jun 18, 2012

**AB** STRIDES PHARMA **EQ 300MG BASE**

**A091050 001** Oct 28, 2016

**ZIAGEN**

**AB** +! VIIV HLTHCARE **EQ 300MG BASE**

**N020977 001** Dec 17, 1998

**ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE**

TABLET;ORAL

**TRIUMEQ**

+! VIIV HLTHCARE **EQ 600MG BASE;EQ 50MG BASE;300MG**

N205551 001 Aug 22, 2014

**ABACAVIR SULFATE; LAMIVUDINE**

TABLET;ORAL

**ABACAVIR SULFATE AND LAMIVUDINE**

**AB** AUROBINDO PHARMA LTD **EQ 600MG BASE;300MG**

**A206151 001** Mar 28, 2017

**AB** CIPLA LTD **EQ 600MG BASE;300MG**

**A091144 001** Mar 28, 2017

**AB** LUPIN LTD **EQ 600MG BASE;300MG**

**A204990 001** Mar 28, 2017

**AB** TEVA PHARMS USA **EQ 600MG BASE;300MG**

**A079246 001** Sep 29, 2016

**EPZICOM**

**AB** +! VIIV HLTHCARE **EQ 600MG BASE;300MG**

**N021652 001** Aug 02, 2004

**ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE**

TABLET;ORAL

**ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE**

**AB** LUPIN LTD **EQ 300MG BASE;150MG;300MG**

**A202912 001** Dec 05, 2013

**TRIZIVIR**

**AB** +! VIIV HLTHCARE **EQ 300MG BASE;150MG;300MG**

**N021205 001** Nov 14, 2000

**ABALOPARATIDE**

SOLUTION;SUBCUTANEOUS

**TYMLOS**

+! RADIUS HEALTH INC **3.12MG/1.56ML (2MG/ML)**

N208743 001 Apr 28, 2017

**ABEMACICLIB**

TABLET;ORAL

**VERZENIO**

+ ELI LILLY AND CO **50MG**

N208716 001 Sep 28, 2017

+ **100MG**

N208716 002 Sep 28, 2017

+ **150MG**

N208716 003 Sep 28, 2017

+! **200MG**

N208716 004 Sep 28, 2017

**ABIRATERONE ACETATE**

TABLET;ORAL

**ZYTIGA**

+! JANSSEN BIOTECH **250MG**

N202379 001 Apr 28, 2011

+ **500MG**

N202379 002 Apr 14, 2017

**ACALABRUTINIB**

CAPSULE;ORAL

**CALQUENCE**

+! ASTRAZENECA **100MG**

N210259 001 Oct 31, 2017

**ACAMPROSATE CALCIUM**

TABLET, DELAYED RELEASE;ORAL

**ACAMPROSATE CALCIUM**

**AB** BARR LABS DIV TEVA **333MG**

**A200143 001** Nov 18, 2013

**AB** ! GLENMARK GENERICS **333MG**

**A202229 001** Jul 16, 2013

**AB** MYLAN PHARMS INC **333MG**

**A200142 001** Mar 11, 2014

**AB** ZYDUS PHARMS USA INC **333MG**

**A205995 001** May 26, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-2 (of 436)

ACARBOSE

TABLET;ORAL

ACARBOSE

<u>AB</u>	EMCURE PHARMS LTD	<u>25MG</u>	<u>A202271 001</u>	Feb 07, 2012
<u>AB</u>		<u>50MG</u>	<u>A202271 002</u>	Feb 07, 2012
<u>AB</u>		<u>100MG</u>	<u>A202271 003</u>	Feb 07, 2012
<u>AB</u>	IMPAX LABS	<u>25MG</u>	<u>A078441 001</u>	May 14, 2009
<u>AB</u>		<u>50MG</u>	<u>A078441 002</u>	May 14, 2009
<u>AB</u>		<u>100MG</u>	<u>A078441 003</u>	May 14, 2009
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A091053 001</u>	Jan 06, 2011
<u>AB</u>		<u>50MG</u>	<u>A091053 002</u>	Jan 06, 2011
<u>AB</u>		<u>100MG</u>	<u>A091053 003</u>	Jan 06, 2011
<u>AB</u>	STRIDES PHARMA	<u>25MG</u>	<u>A090912 001</u>	Jul 27, 2011
<u>AB</u>		<u>50MG</u>	<u>A090912 002</u>	Jul 27, 2011
<u>AB</u>		<u>100MG</u>	<u>A090912 003</u>	Jul 27, 2011
<u>AB</u>	VIRTUS PHARM	<u>25MG</u>	<u>A091343 001</u>	Oct 17, 2013
<u>AB</u>		<u>50MG</u>	<u>A091343 002</u>	Oct 17, 2013
<u>AB</u>		<u>100MG</u>	<u>A091343 003</u>	Oct 17, 2013
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A077532 001</u>	May 07, 2008
<u>AB</u>		<u>50MG</u>	<u>A077532 002</u>	May 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A077532 003</u>	May 07, 2008
<u>AB</u>	WEST-WARD PHARMS INT	<u>25MG</u>	<u>A078470 001</u>	May 07, 2008
<u>AB</u>		<u>50MG</u>	<u>A078470 002</u>	May 07, 2008
<u>AB</u>		<u>100MG</u>	<u>A078470 003</u>	May 07, 2008
<u>PRECPOSE</u>				
<u>AB</u>	+! BAYER HLTHCARE	<u>25MG</u>	<u>N020482 004</u>	May 29, 1997
<u>AB</u>	+	<u>50MG</u>	<u>N020482 001</u>	Sep 06, 1995
<u>AB</u>	+	<u>100MG</u>	<u>N020482 002</u>	Sep 06, 1995

ACEBUTOLOL HYDROCHLORIDE

CAPSULE;ORAL

ACEBUTOLOL HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARM	<u>EQ 200MG BASE</u>	<u>A075047 001</u>	Dec 30, 1999
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A075047 002</u>	Dec 30, 1999
<u>AB</u>	MYLAN	<u>EQ 200MG BASE</u>	<u>A074288 001</u>	Apr 24, 1995
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A074288 002</u>	Apr 24, 1995
<u>SECTRAL</u>				
<u>AB</u>	+ PROMIUS PHARMA	<u>EQ 200MG BASE</u>	<u>N018917 001</u>	Dec 28, 1984
<u>AB</u>	+	<u>EQ 400MG BASE</u>	<u>N018917 003</u>	Dec 28, 1984

ACETAMINOPHEN

SOLUTION;IV (INFUSION)

ACETAMINOPHEN

<u>AP</u>	CUSTOPHARM INC	<u>1GM/100ML (10MG/ML)</u>	<u>A202605 001</u>	Jun 13, 2016
<u>AP</u>	SANDOZ INC	<u>1GM/100ML (10MG/ML)</u>	<u>A204052 001</u>	Mar 22, 2016
<u>OFIRMEV</u>				
<u>AP</u>	+! MALLINCKRODT IP ACETAMINOPHEN	<u>1GM/100ML (10MG/ML)</u>	<u>N022450 001</u>	Nov 02, 2010

FRESENIUS KABI USA 1GM/100ML (10MG/ML)

N204767 001 Oct 28, 2015

ACETAMINOPHEN; BUTALBITAL

CAPSULE;ORAL

BUTALBITAL AND ACETAMINOPHEN

MIKART INC 300MG;50MG

A207313 001 Dec 27, 2017

TABLET;ORAL

BUTALBITAL AND ACETAMINOPHEN

<u>AA</u>	CNTY LINE PHARMS	<u>325MG;50MG</u>	<u>A205120 001</u>	Oct 30, 2015
<u>AA</u>	LARKEN LABS INC	<u>325MG;50MG</u>	<u>A203484 002</u>	Dec 04, 2015
<u>AA</u>	MIKART INC	<u>300MG;50MG</u>	<u>A207386 001</u>	Nov 15, 2016
<u>AA</u>	! NEXGEN PHARMA	<u>300MG;50MG</u>	<u>A090956 001</u>	Aug 23, 2011
<u>AA</u>	TEDOR PHARMA INC	<u>300MG;50MG</u>	<u>A207635 001</u>	Jun 05, 2017
<u>BUTAPAP</u>				
<u>AA</u>	! MIKART ALLZITAL	<u>325MG;50MG</u>	<u>A089987 001</u>	Oct 26, 1992

LARKEN LABS INC 325MG;25MG

A203484 001 Dec 04, 2015

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AA</u>	! NEXGEN PHARMA	<u>300MG;50MG;40MG</u>	<u>A040885 001</u>	Nov 16, 2009
<u>AB</u>	NUVO PHARM INC	<u>300MG;50MG;40MG</u>	<u>A207118 001</u>	Oct 28, 2016
<u>AB</u>	TEDOR PHARMA INC	<u>300MG;50MG;40MG</u>	<u>A206615 001</u>	Aug 04, 2017
	MAYNE PHARMA INC	325MG;50MG;40MG	A089007 001	Mar 17, 1986

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-3 (of 436)

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

SOLUTION;ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE  
! MIKART 325MG/15ML;50MG/15ML;40MG/15ML

A040387 001 Jan 31, 2003

TABLET;ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<b>AA</b>	ACTAVIS LABS UT INC	<u>325MG;50MG;40MG</u>
<b>AA</b>	CNTY LINE PHARMS	<u>325MG;50MG;40MG</u>
<b>AA</b>	HIKMA PHARMS	<u>325MG;50MG;40MG</u>
<b>AA</b>	LANNETT HOLDINGS INC	<u>325MG;50MG;40MG</u>
<b>AA</b>	MIKART	<u>325MG;50MG;40MG</u>
<b>AA</b>	SPECGX LLC	<u>325MG;50MG;40MG</u>
<b>AA</b> !	VINTAGE PHARMS	<u>325MG;50MG;40MG</u>

<b>A088616</b>	<b>001</b>	Nov 09, 1984
<b>A204984</b>	<b>001</b>	Jan 10, 2017
<b>A089718</b>	<b>001</b>	Jun 12, 1995
<b>A200243</b>	<b>001</b>	Sep 13, 2012
<b>A089175</b>	<b>001</b>	Jan 21, 1987
<b>A087804</b>	<b>001</b>	Jan 24, 1985
<b>A040511</b>	<b>001</b>	Aug 27, 2003

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

<b>AB</b>	NEXGEN PHARMA INC	<u>325MG;50MG;40MG;30MG</u>
<b>AB</b>	VINTAGE PHARMS	<u>325MG;50MG;40MG;30MG</u>
<b>FIORICET W/ CODEINE</b>		
<b>AB</b>	+! ACTAVIS LABS UT INC	<u>325MG;50MG;40MG;30MG</u>
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE		
NEXGEN PHARMA INC 300MG;50MG;40MG;30MG		

<b>A076560</b>	<b>001</b>	Jun 10, 2004
<b>A075929</b>	<b>001</b>	Apr 22, 2002
<b>N020232</b>	<b>001</b>	Jul 30, 1992
A076560	002	Jul 19, 2012

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

TREZIX

WRASER PHARMS LLC 320.5MG;30MG;16MG

A204785 001 Nov 26, 2014

TABLET;ORAL

ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE  
LARKEN LABS INC 325MG;30MG;16MG

A204209 001 Sep 30, 2016

ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

<b>AA</b>	HI TECH PHARMA	<u>120MG/5ML;12MG/5ML</u>
<b>AA</b>	MIKART	<u>120MG/5ML;12MG/5ML</u>
<b>AA</b> !	PHARM ASSOC	<u>120MG/5ML;12MG/5ML</u>
<b>AA</b>	VINTAGE PHARMS	<u>120MG/5ML;12MG/5ML</u>
<b>AA</b>	WOCKHARDT BIO AG	<u>120MG/5ML;12MG/5ML</u>

<b>A040119</b>	<b>001</b>	Apr 26, 1996
<b>A089450</b>	<b>001</b>	Oct 27, 1992
<b>A087508</b>	<b>001</b>	
<b>A091238</b>	<b>001</b>	Nov 10, 2011
<b>A087006</b>	<b>001</b>	

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

<b>AA</b>	AMNEAL PHARMS NY	<u>300MG;30MG</u>
<b>AA</b>	AUROLIFE PHARMA LLC	<u>300MG;15MG</u>
<b>AA</b>		<u>300MG;30MG</u>
<b>AA</b>		<u>300MG;60MG</u>
<b>AA</b> !	SPECGX LLC	<u>300MG;15MG</u>
<b>AA</b>		<u>300MG;30MG</u>
<b>AA</b>		<u>300MG;60MG</u>
<b>AA</b>	SUN PHARM INDS LTD	<u>300MG;30MG</u>
<b>AA</b>		<u>300MG;60MG</u>
<b>AA</b>	TEVA	<u>300MG;15MG</u>
<b>AA</b>		<u>300MG;30MG</u>
<b>AA</b> !		<u>300MG;60MG</u>
<b>AA</b>	VINTAGE	<u>300MG;15MG</u>
<b>AA</b>		<u>300MG;30MG</u>
<b>AA</b>	VINTAGE PHARMS	<u>300MG;60MG</u>
<b>TYLENOL W/ CODEINE NO. 3</b>		
<b>AA</b> !	JANSSEN PHARMS	<u>300MG;30MG</u>
<b>TYLENOL W/ CODEINE NO. 4</b>		
<b>AA</b>	JANSSEN PHARMS	<u>300MG;60MG</u>

<b>A040779</b>	<b>001</b>	May 29, 2008
<b>A202800</b>	<b>001</b>	Apr 15, 2013
<b>A202800</b>	<b>002</b>	Apr 15, 2013
<b>A202800</b>	<b>003</b>	Apr 15, 2013
<b>A040419</b>	<b>001</b>	May 31, 2001
<b>A040419</b>	<b>002</b>	May 31, 2001
<b>A040419</b>	<b>003</b>	May 31, 2001
<b>A085868</b>	<b>001</b>	
<b>A087083</b>	<b>001</b>	
<b>A088627</b>	<b>001</b>	Mar 06, 1985
<b>A088628</b>	<b>001</b>	Mar 06, 1985
<b>A088629</b>	<b>001</b>	Mar 06, 1985
<b>A089990</b>	<b>001</b>	Sep 30, 1988
<b>A089805</b>	<b>001</b>	Sep 30, 1988
<b>A089828</b>	<b>001</b>	Sep 30, 1988
<b>A085055</b>	<b>003</b>	
<b>A085055</b>	<b>004</b>	

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<b>AA</b> !	MIKART	<u>325MG/15ML;7.5MG/15ML</u>
<b>AA</b>	PHARM ASSOC	<u>325MG/15ML;7.5MG/15ML</u>
<b>AA</b>	VINTAGE PHARMS	<u>325MG/15ML;7.5MG/15ML</u>
<b>AA</b>	VISTAPHARM	<u>325MG/15ML;7.5MG/15ML</u>
!	MIKART	300MG/15ML;10MG/15ML
!	PHARM ASSOC	325MG/15ML;10MG/15ML

<b>A040482</b>	<b>001</b>	Sep 25, 2003
<b>A040838</b>	<b>001</b>	May 10, 2013
<b>A040894</b>	<b>001</b>	Jul 19, 2011
<b>A200343</b>	<b>001</b>	Jan 25, 2012
A040881	001	Feb 25, 2010
A040834	001	Apr 18, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-4 (of 436)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET;ORAL

ANEXSIA 5/325

AA SPECGX LLC 325MG;5MG A040409 001 Oct 20, 2000

ANEXSIA 7.5/325

AA SPECGX LLC 325MG;7.5MG A040405 001 Sep 08, 2000

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA ABHAI LLC 300MG;5MG A209036 001 Jun 21, 2017

AA 300MG;7.5MG A209036 002 Jun 21, 2017

AA 300MG;10MG A209036 003 Jun 21, 2017

AA 325MG;5MG A209037 001 Jun 21, 2017

AA 325MG;7.5MG A209037 002 Jun 21, 2017

AA 325MG;10MG A209037 003 Jun 21, 2017

AA ACTAVIS LABS FL INC 300MG;5MG A206470 001 Jun 02, 2016

AA 300MG;7.5MG A206470 002 Jun 02, 2016

AA 300MG;10MG A206470 003 Jun 02, 2016

AA AMNEAL PHARMS 300MG;10MG A207137 001 Nov 29, 2016

AA AMNEAL PHARMS NY 300MG;5MG A206869 001 Jun 23, 2017

AA 325MG;5MG A040736 001 Aug 25, 2006

AA 325MG;7.5MG A040746 002 May 10, 2016

AA 325MG;10MG A040746 001 Aug 25, 2006

AA AUROLIFE PHARMA LLC 325MG;5MG A201013 001 Apr 11, 2012

AA 325MG;7.5MG A201013 002 Apr 11, 2012

AA 325MG;10MG A201013 003 Apr 11, 2012

AA LANNETT HOLDINGS INC 300MG;5MG A207171 001 Jun 20, 2017

AA 300MG;7.5MG A207171 002 Jun 20, 2017

AA 300MG;10MG A207171 003 Jun 20, 2017

AA 325MG;5MG A207172 001 Jun 22, 2017

AA 325MG;7.5MG A207172 002 Jun 22, 2017

AA 325MG;10MG A207172 003 Jun 22, 2017

AA LARKEN LABS INC 325MG;5MG A202935 002 Jun 15, 2016

AA 325MG;7.5MG A202935 003 Jun 15, 2016

AA 325MG;10MG A202935 004 Jun 15, 2016

AA ! MIKART 300MG;5MG A040658 001 Jan 19, 2006

AA ! 300MG;7.5MG A040658 002 Mar 24, 2006

AA ! 300MG;10MG A040658 003 Jun 23, 2004

AA ! 325MG;2.5MG A040846 001 Jun 09, 2010

AA ! 325MG;7.5MG A040432 001 Jan 22, 2003

AA NOVEL LABS INC 300MG;5MG A206142 001 Nov 14, 2016

AA 300MG;7.5MG A206142 002 Nov 14, 2016

AA 300MG;10MG A206142 003 Nov 14, 2016

AA 325MG;5MG A206245 001 Dec 01, 2016

AA 325MG;7.5MG A206245 002 Dec 01, 2016

AA 325MG;10MG A206245 003 Dec 01, 2016

AA PAR PHARM 300MG;5MG A205001 001 Jul 05, 2016

AA 300MG;7.5MG A205001 002 Jul 05, 2016

AA 300MG;10MG A205001 003 Jul 05, 2016

AA RHODES PHARMS 325MG;5MG A202991 001 Apr 12, 2016

AA 325MG;7.5MG A202991 002 Apr 12, 2016

AA 325MG;10MG A202991 003 Apr 12, 2016

AA SPECGX LLC 300MG;5MG A206718 001 Mar 31, 2017

AA 300MG;7.5MG A206718 002 Mar 31, 2017

AA 300MG;10MG A206718 003 Mar 31, 2017

AA 325MG;10MG A040400 001 Jul 26, 2000

AA SUN PHARM IND'S INC 325MG;5MG A090118 001 Dec 23, 2008

AA 325MG;7.5MG A090118 002 Dec 23, 2008

AA 325MG;10MG A090118 003 Dec 23, 2008

AA TRIS PHARMA INC 300MG;5MG A202214 004 Mar 15, 2016

AA 300MG;7.5MG A202214 005 Mar 15, 2016

AA 300MG;10MG A202214 006 Mar 15, 2016

AA 325MG;5MG A202214 001 Mar 27, 2013

AA 325MG;7.5MG A202214 002 Mar 27, 2013

AA 325MG;10MG A202214 003 Mar 27, 2013

AA UPSHER-SMITH LABS 325MG;5MG A206484 001 Mar 24, 2017

AA 325MG;7.5MG A206484 002 Mar 24, 2017

AA 325MG;10MG A206484 003 Mar 24, 2017

AA VINTAGE PHARMS 300MG;5MG A090415 001 Jan 24, 2011

AA 300MG;7.5MG A090415 002 Jan 24, 2011

AA 300MG;10MG A090415 003 Jan 24, 2011

AA 325MG;5MG A040655 001 Jan 19, 2006

AA 325MG;7.5MG A040656 001 Jan 19, 2006

AA 325MG;10MG A040355 001 May 31, 2000

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-5 (of 436)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	WES PHARMA INC	<u>325MG;5MG</u>	<u>A210211 001</u>	Oct 30, 2017
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A210211 002</u>	Oct 30, 2017
<u>AA</u>		<u>325MG;10MG</u>	<u>A210211 003</u>	Oct 30, 2017
	<u>NORCO</u>			
<u>AA</u>	APIL	<u>325MG;2.5MG</u>	<u>A040148 004</u>	Jul 07, 2014
<u>AA</u>	!	<u>325MG;5MG</u>	<u>A040099 001</u>	Jun 25, 1997
<u>AA</u>	!	<u>325MG;5MG</u>	<u>A040148 005</u>	Jul 07, 2014
<u>AA</u>	!	<u>325MG;7.5MG</u>	<u>A040148 003</u>	Sep 12, 2000
<u>AA</u>	!	<u>325MG;10MG</u>	<u>A040148 001</u>	Feb 14, 1997

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

SOLUTION;ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	! SPECGX LLC	<u>325MG/5ML;5MG/5ML</u>	<u>A040680 001</u>	Sep 29, 2006
<u>AA</u>	VINTAGE PHARMS	<u>325MG/5ML;5MG/5ML</u>	<u>A203573 001</u>	Dec 18, 2014

TABLET;ORAL

OXYCET

<u>AA</u>	SPECGX LLC	<u>325MG;5MG</u>	<u>A087463 001</u>	Dec 07, 1983
-----------	------------	------------------	--------------------	--------------

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>	ACTAVIS ELIZABETH	<u>325MG;2.5MG</u>	<u>A201447 001</u>	Apr 12, 2013
<u>AA</u>		<u>325MG;5MG</u>	<u>A201447 002</u>	Apr 12, 2013
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A201447 003</u>	Apr 12, 2013
<u>AA</u>		<u>325MG;10MG</u>	<u>A201447 004</u>	Apr 12, 2013
<u>AA</u>	ALVOGEN MALTA	<u>325MG;5MG</u>	<u>A202677 003</u>	Mar 08, 2016
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A202677 001</u>	Jul 26, 2012
<u>AA</u>		<u>325MG;10MG</u>	<u>A202677 002</u>	Jul 26, 2012
<u>AA</u>	AMNEAL PHARMS	<u>325MG;5MG</u>	<u>A040777 001</u>	Nov 27, 2007
<u>AA</u>	AMNEAL PHARMS NY	<u>325MG;7.5MG</u>	<u>A040778 002</u>	Jun 27, 2014
<u>AA</u>		<u>325MG;10MG</u>	<u>A040778 001</u>	Nov 27, 2007
<u>AA</u>	ASCENT PHARMS INC	<u>325MG;2.5MG</u>	<u>A207419 001</u>	Mar 22, 2017
<u>AA</u>		<u>325MG;5MG</u>	<u>A207419 002</u>	Mar 22, 2017
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A207419 003</u>	Mar 22, 2017
<u>AA</u>		<u>325MG;10MG</u>	<u>A207419 004</u>	Mar 22, 2017
<u>AA</u>	AUROLIFE PHARMA LLC	<u>325MG;2.5MG</u>	<u>A201972 001</u>	Jul 15, 2013
<u>AA</u>		<u>325MG;5MG</u>	<u>A201972 002</u>	Jul 15, 2013
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A201972 003</u>	Jul 15, 2013
<u>AA</u>		<u>325MG;10MG</u>	<u>A201972 004</u>	Jul 15, 2013
<u>AA</u>	CHEMO RESEARCH SL	<u>325MG;5MG</u>	<u>A207574 001</u>	Dec 13, 2016
<u>AA</u>	LANNETT HOLDINGS INC	<u>325MG;5MG</u>	<u>A207333 001</u>	Sep 25, 2017
<u>AA</u>		<u>325MG;10MG</u>	<u>A207333 002</u>	Sep 25, 2017
<u>AA</u>	MAYNE PHARMA INC	<u>325MG;2.5MG</u>	<u>A090177 001</u>	Oct 20, 2008
<u>AA</u>		<u>325MG;5MG</u>	<u>A090177 002</u>	Oct 20, 2008
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A090177 003</u>	Oct 20, 2008
<u>AA</u>		<u>325MG;10MG</u>	<u>A090177 004</u>	Oct 20, 2008
<u>AA</u>	NESHER PHARMS	<u>325MG;2.5MG</u>	<u>A210079 001</u>	Dec 28, 2017
<u>AA</u>		<u>325MG;5MG</u>	<u>A210079 002</u>	Dec 28, 2017
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A210079 003</u>	Dec 28, 2017
<u>AA</u>		<u>325MG;10MG</u>	<u>A210079 004</u>	Dec 28, 2017
<u>AA</u>	NOVEL LABS INC	<u>325MG;2.5MG</u>	<u>A204407 001</u>	Feb 24, 2017
<u>AA</u>		<u>325MG;5MG</u>	<u>A204407 002</u>	Feb 24, 2017
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A204407 003</u>	Feb 24, 2017
<u>AA</u>		<u>325MG;10MG</u>	<u>A204407 004</u>	Feb 24, 2017
<u>AA</u>	RHODES PHARMS	<u>325MG;5MG</u>	<u>A201278 001</u>	Aug 28, 2014
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A201278 002</u>	Aug 28, 2014
<u>AA</u>		<u>325MG;10MG</u>	<u>A201278 003</u>	Aug 28, 2014
<u>AA</u>	SPECGX LLC	<u>325MG;7.5MG</u>	<u>A040545 001</u>	Jun 30, 2004
<u>AA</u>		<u>325MG;10MG</u>	<u>A040545 002</u>	Jun 30, 2004
<u>AA</u>	SUN PHARM INDNS INC	<u>325MG;2.5MG</u>	<u>A090535 001</u>	Dec 26, 2013
<u>AA</u>		<u>325MG;5MG</u>	<u>A090535 002</u>	Dec 26, 2013
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A090535 003</u>	Dec 26, 2013
<u>AA</u>		<u>325MG;10MG</u>	<u>A090535 004</u>	Dec 26, 2013
<u>AA</u>	VINTAGE PHARMS	<u>325MG;2.5MG</u>	<u>A090733 001</u>	Jul 11, 2013
<u>AA</u>		<u>325MG;5MG</u>	<u>A040105 001</u>	Jul 30, 1996
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A090734 001</u>	Jul 11, 2013
<u>AA</u>		<u>325MG;10MG</u>	<u>A090734 002</u>	Jul 11, 2013
<u>AA</u>	WATSON LABS	<u>325MG;5MG</u>	<u>A040171 001</u>	Oct 30, 1997
<u>AA</u>		<u>325MG;7.5MG</u>	<u>A040535 001</u>	Sep 05, 2003
<u>AA</u>		<u>325MG;10MG</u>	<u>A040535 002</u>	Sep 05, 2003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-6 (of 436)

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

**PERCOSET**

<b>AA</b>	!	VINTAGE PHARMS LLC	<u>325MG;2.5MG</u>	<b>A040330</b>	<b>001</b>	Jun 25, 1999
<b>AA</b>	!		<u>325MG;5MG</u>	<b>A040330</b>	<b>002</b>	Jun 25, 1999
<b>AA</b>	!		<u>325MG;7.5MG</u>	<b>A040330</b>	<b>003</b>	Nov 23, 2001
<b>AA</b>	!		<u>325MG;10MG</u>	<b>A040330</b>	<b>004</b>	Nov 23, 2001

**ROXICET**

<b>AA</b>	WEST-WARD PHARMS INT	<u>325MG;5MG</u>	<b>A087003</b>	<b>001</b>	
	OXYCODONE AND ACETAMINOPHEN				
	! MIKART	300MG;2.5MG	A040608	001	Dec 30, 2005
	!	300MG;5MG	A040608	002	Dec 30, 2005
	!	300MG;7.5MG	A040608	003	Dec 30, 2005
	!	300MG;10MG	A040608	004	Dec 30, 2005

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET;ORAL

**TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN**

<b>AB</b>	ALKEM LABS LTD	<u>325MG;37.5MG</u>	<b>A202076</b>	<b>001</b>	Mar 30, 2012
<b>AB</b>	AMNEAL PHARMS	<u>325MG;37.5MG</u>	<b>A090485</b>	<b>001</b>	Dec 09, 2009
<b>AB</b>	APOTEX INC	<u>325MG;37.5MG</u>	<b>A078778</b>	<b>001</b>	Apr 07, 2014
<b>AB</b>	AUROBINDO PHARMA LTD	<u>325MG;37.5MG</u>	<b>A207152</b>	<b>001</b>	Mar 22, 2017
<b>AB</b>	MACLEODS PHARMS LTD	<u>325MG;37.5MG</u>	<b>A206885</b>	<b>001</b>	May 02, 2017
<b>AB</b>	MICRO LABS LTD INDIA	<u>325MG;37.5MG</u>	<b>A201952</b>	<b>001</b>	Dec 14, 2012
<b>AB</b>	MYLAN	<u>325MG;37.5MG</u>	<b>A077858</b>	<b>001</b>	Sep 26, 2008
<b>AB</b>	PAR PHARM	<u>325MG;37.5MG</u>	<b>A076475</b>	<b>001</b>	Apr 21, 2005
<b>AB</b>	SUN PHARM INDNS INC	<u>325MG;37.5MG</u>	<b>A077184</b>	<b>001</b>	Dec 16, 2005
<b>AB</b>	ZYDUS PHARMS USA INC	<u>325MG;37.5MG</u>	<b>A090460</b>	<b>001</b>	Sep 06, 2012

**ULTRACET**

<b>AB</b>	++! JANSSEN PHARMS	<u>325MG;37.5MG</u>	<b>N021123</b>	<b>001</b>	Aug 15, 2001
-----------	--------------------	---------------------	----------------	------------	--------------

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE;ORAL

**ACETAZOLAMIDE**

<b>AB</b>	HERITAGE PHARMS INC	<u>500MG</u>	<b>A090779</b>	<b>001</b>	Jul 14, 2011
<b>AB</b>	NOSTRUM LABS INC	<u>500MG</u>	<b>A204691</b>	<b>001</b>	Mar 29, 2016
<b>AB</b>	NOVAST LABS LTD	<u>500MG</u>	<b>A203434</b>	<b>001</b>	Sep 30, 2016
<b>AB</b>	ZYDUS PHARMS USA INC	<u>500MG</u>	<b>A040904</b>	<b>001</b>	Dec 10, 2008

**DIAMOX**

<b>AB</b>	++! TEVA BRANDED PHARM	<u>500MG</u>	<b>N012945</b>	<b>001</b>	
-----------	------------------------	--------------	----------------	------------	--

TABLET;ORAL

**ACETAZOLAMIDE**

<b>AB</b>	HERITAGE PHARMA	<u>125MG</u>	<b>A205530</b>	<b>001</b>	Oct 27, 2016
<b>AB</b>		<u>250MG</u>	<b>A205530</b>	<b>002</b>	Oct 27, 2016
<b>AB</b>	LANNETT	<u>250MG</u>	<b>A084840</b>	<b>001</b>	
<b>AB</b>	STRIDES PHARMA	<u>125MG</u>	<b>A209734</b>	<b>001</b>	Nov 20, 2017
<b>AB</b>		<u>250MG</u>	<b>A209734</b>	<b>002</b>	Nov 20, 2017
<b>AB</b>	SUN PHARM INDUSTRIES	<u>125MG</u>	<b>A089753</b>	<b>002</b>	Jun 22, 1988
<b>AB</b>	TARO	<u>125MG</u>	<b>A040195</b>	<b>001</b>	May 28, 1997
<b>AB</b>	!	<u>250MG</u>	<b>A040195</b>	<b>002</b>	May 28, 1997

ACETAZOLAMIDE SODIUM

INJECTABLE;INJECTION

**ACETAZOLAMIDE SODIUM**

<b>AP</b>	PAR STERILE PRODUCTS	<u>EQ 500MG BASE/VIAL</u>	<b>A205358</b>	<b>001</b>	Jun 20, 2017
<b>AP</b>	SAGENT AGILA	<u>EQ 500MG BASE/VIAL</u>	<b>A200880</b>	<b>001</b>	May 09, 2012
<b>AP</b>	WEST-WARD PHARMS INT	<u>EQ 500MG BASE/VIAL</u>	<b>A040089</b>	<b>001</b>	Feb 28, 1995
<b>AP</b>	! X GEN PHARMS	<u>EQ 500MG BASE/VIAL</u>	<b>A040784</b>	<b>001</b>	Dec 10, 2008
<b>AP</b>	<b>ACETAZOLAMIDE SODIUM</b>	<u>EQ 500MG BASE/VIAL</u>	<b>A202693</b>	<b>001</b>	Dec 19, 2014

ACETIC ACID, GLACIAL

SOLUTION;IRRIGATION, URETHRALL

**ACETIC ACID 0.25% IN PLASTIC CONTAINER**

<b>AT</b>	B BRAUN	<u>250MG/100ML</u>	<b>N018161</b>	<b>001</b>	
<b>AT</b>	BAXTER HLTHCARE	<u>250MG/100ML</u>	<b>N018523</b>	<b>001</b>	Feb 19, 1982
<b>AT</b>	ICU MEDICAL INC	<u>250MG/100ML</u>	<b>N017656</b>	<b>001</b>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-7 (of 436)

ACETIC ACID, GLACIAL

SOLUTION/DROPS;OTIC

ACETIC ACID

<u>AT</u>	TARO	<u>2%</u>	<u>A088638 001</u>	Sep 06, 1984
<u>AT</u>	VINTAGE	<u>2%</u>	<u>A040607 001</u>	Feb 24, 2005
<u>AT</u>	! WOCKHARDT BIO AG	<u>2%</u>	<u>A040166 001</u>	Jul 26, 1996
	<u>VOSOL</u>			
<u>AT</u>	HI TECH PHARMA	<u>2%</u>	<u>N012179 001</u>	

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

ACETASOL HC

<u>AT</u>	ACTAVIS MID	<u>2%:1%</u>	<u>A087143 001</u>	Jan 13, 1982
	ATLANTIC			
	<u>HYDROCORTISONE AND ACETIC ACID</u>			
<u>AT</u>	TARO	<u>2%:1%</u>	<u>A088759 001</u>	Mar 04, 1985

VINTAGE

VOSOL HC

+! HI TECH PHARMA

2%:1%

A040609 001 Feb 06, 2006

N012770 001

ACETOHYDROXAMIC ACID

TABLET;ORAL

LITHOSTAT

+! MISSION PHARMA

250MG

N018749 001 May 31, 1983

ACETYLCHOLINE CHLORIDE

FOR SOLUTION;OPHTHALMIC

MIOCHOL-E

+! BAUSCH AND LOMB

20MG/VIAL

N020213 001 Sep 22, 1993

ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETADOTE

AP +! CUMBERLAND PHARMS

6GM/30ML (200MG/ML)

N021539 001 Jan 23, 2004

ACETYLCYSTEINE

AP AKORN INC

6GM/30ML (200MG/ML)

A203173 001 Mar 24, 2015

AP AUROBINDO PHARMA LTD

6GM/30ML (200MG/ML)

A207358 001 Feb 29, 2016

AP FRESENIUS KABI USA

6GM/30ML (200MG/ML)

A200644 001 Nov 07, 2012

AP MYLAN INSTITUTIONAL

6GM/30ML (200MG/ML)

A203624 001 Jun 19, 2015

AP SAGENT PHARMS

6GM/30ML (200MG/ML)

A091684 001 Oct 31, 2017

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

AN ALVOGEN INC

10%

A204674 001 Feb 11, 2014

AN 20%

A203853 001 Jun 21, 2012

AN HOSPIRA

10%

A073664 001 Aug 30, 1994

AN 20%

A074037 001 Aug 30, 1994

AN ! LUITPOLD

10%

A072489 001 Jul 28, 1995

AN ! 20%

A072547 001 Jul 28, 1995

TABLET, EFFERVESCENT;ORAL

CETYLEV

+ ARBOR PHARMS LLC

500MG

N207916 001 Jan 29, 2016

+!

2.5GM

N207916 002 Jan 29, 2016

ACITRETIN

CAPSULE;ORAL

ACITRETIN

AB BARR LABS INC

10MG

A091455 001 Apr 04, 2013

AB 25MG

A091455 002 Apr 04, 2013

AB IMPAX LABS INC

10MG

A202552 001 Dec 23, 2015

AB 17.5MG

A202552 002 Dec 23, 2015

AB 22.5MG

A202552 003 Dec 23, 2015

AB 25MG

A202552 004 Dec 23, 2015

AB MYLAN PHARMS INC

10MG

A202148 001 Sep 10, 2015

AB 25MG

A202148 002 Sep 10, 2015

AB SIGMAPHARM LABS LLC

10MG

A204633 001 May 22, 2015

AB 17.5MG

A204633 002 May 22, 2015

AB 22.5MG

A204633 003 May 22, 2015

AB 25MG

A204633 004 May 22, 2015

AB TEVA PHARMS USA

17.5MG

A202897 001 Apr 04, 2013

AB 22.5MG

A202897 002 Apr 04, 2013

SORIATANE

AB + STIEFEL LABS INC

10MG

N019821 001 Oct 28, 1996

AB +

17.5MG

N019821 003 Aug 06, 2009

AB +

22.5MG

N019821 004 Aug 06, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-8 (of 436)

ACITRETIN

CAPSULE;ORAL

SORIATANE

AB +!

25MG

N019821 002 Oct 28, 1996

ACLIDINIUM BROMIDE

POWDER, METERED;INHALATION

TUDORZA PRESSAIR

+! ASTRAZENECA PHARMS 0.4MG/INH

N202450 001 Jul 23, 2012

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE;ORAL

SEMPREX-D

+! AUXILIUM PHARMS LLC 8MG;60MG

N019806 001 Mar 25, 1994

ACYCLOVIR

CAPSULE;ORAL

ACYCLOVIR

AB ! APOTEX INC 200MG

A075677 001 Sep 28, 2005

AB BOSCOGEN 200MG

A075090 001 Jan 26, 1999

AB CADILA PHARMS LTD 200MG

A201445 001 Mar 06, 2014

AB CARLSBAD TECHNOLOGY 200MG

A206261 001 Aug 16, 2017

AB DAVA PHARMS INC 200MG

A074833 001 Apr 22, 1997

AB TEVA 200MG

A074578 001 Apr 22, 1997

AB ZYDUS PHARMS USA 200MG

A204313 001 Mar 25, 2016

INC

ZOVIRAX

AB + MYLAN PHARMS INC 200MG

N018828 001 Jan 25, 1985

CREAM;TOPICAL

ZOVIRAX

+! VIB 5%

N021478 001 Dec 30, 2002

OINTMENT;TOPICAL

ACYCLOVIR

AB AMNEAL PHARMS 5%

A204605 001 Jun 18, 2014

AB FOUGERA PHARMS INC 5%

A206633 001 May 11, 2016

AB G AND W LABS INC 5%

A205591 001 Nov 13, 2017

AB GLENMARK PHARMS LTD 5%

A205510 001 Jul 31, 2017

AB MYLAN PHARMS INC 5%

A202459 001 Apr 03, 2013

AB TARO 5%

A205469 001 Dec 21, 2016

AB TOLMAR 5%

A206437 001 Jul 31, 2017

ZOVIRAX

AB +! VALEANT BERMUDA 5%

N018604 001 Mar 29, 1982

SUSPENSION;ORAL

ACYCLOVIR

AB ACTAVIS MID 200MG/5ML

A074738 001 Apr 28, 1997

ATLANTIC

AB HI TECH PHARMA 200MG/5ML

A077026 001 Jun 07, 2005

ZOVIRAX

AB +! MYLAN PHARMS INC 200MG/5ML

N019909 001 Dec 22, 1989

TABLET;BUCCAL

SITAVIG

+! EPI HLTH 50MG

N203791 001 Apr 12, 2013

TABLET;ORAL

ACYCLOVIR

AB APOTEX INC 400MG

A077309 001 Sep 29, 2005

AB 800MG

A077309 002 Sep 29, 2005

AB CADILA PHARMS LTD 400MG

A202168 001 Nov 15, 2013

AB 800MG

A202168 002 Nov 15, 2013

AB CARLSBAD 400MG

A075382 001 Apr 30, 1999

AB 800MG

A075382 002 Apr 30, 1999

AB DAVA PHARMS INC 400MG

A074946 001 Nov 19, 1997

AB 800MG

A074946 002 Nov 19, 1997

AB HERITAGE PHARMS INC 400MG

A074891 001 Oct 31, 1997

AB 800MG

A074891 002 Oct 31, 1997

AB HETERO LABS LTD V 400MG

A203834 001 Oct 29, 2013

AB 800MG

A203834 002 Oct 29, 2013

AB MYLAN PHARMS INC 400MG

A075211 001 Sep 28, 1998

AB 800MG

A075211 002 Sep 28, 1998

AB TEVA 400MG

A074556 002 Apr 22, 1997

AB 800MG

A074556 003 Apr 22, 1997

AB ZYDUS PHARMS USA 400MG

A204314 001 Aug 19, 2014

AB INC 800MG

A204314 002 Aug 19, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-9 (of 436)

ACYCLOVIR

TABLET;ORAL

ZOVIRAX

<u>AB</u>	+	MYLAN PHARMS INC	<u>400MG</u>	<u>N020089 001</u>	Apr 30, 1991
<u>AB</u>	+!		<u>800MG</u>	<u>N020089 002</u>	Apr 30, 1991

ACYCLOVIR SODIUM

INJECTABLE;INJECTION

ACYCLOVIR SODIUM

<u>AP</u>		AUROBINDO PHARMA LTD	<u>EQ 50MG BASE/ML</u>	<u>A203701 001</u>	Oct 11, 2013
<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 50MG BASE/ML</u>	<u>A074930 001</u>	May 13, 1998
<u>AP</u>	!		<u>EQ 500MG BASE/VIAL</u>	<u>A075015 001</u>	Apr 30, 1998
<u>AP</u>		HIKMA PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A205771 001</u>	Feb 29, 2016
<u>AP</u>	!		<u>EQ 1GM BASE/VIAL</u>	<u>A205771 002</u>	Feb 29, 2016
<u>AP</u>		ZYDUS PHARMS USA INC	<u>EQ 500MG BASE/VIAL</u>	<u>A206606 001</u>	Jun 13, 2017
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>A206606 002</u>	Jun 13, 2017

ACYCLOVIR; HYDROCORTISONE

CREAM;TOPICAL

XERESE

+! VALEANT BERMUDA 5%;1%

N022436 001 Jul 31, 2009

ADAPALENE

CREAM;TOPICAL

ADAPALENE

<u>AB</u>		FOUGERA PHARMS	<u>0.1%</u>	<u>A090824 001</u>	Jun 30, 2010
<u>AB</u>	!	GALDERMA LABS LP	<u>0.1%</u>	<u>N020748 001</u>	May 26, 2000

GEL;TOPICAL

ADAPALENE

<u>AB</u>		ACTAVIS MID ATLANTIC	<u>0.3%</u>	<u>A201000 001</u>	Oct 27, 2014
<u>AB</u>		GLENMARK GENERICS	<u>0.1%</u>	<u>A091314 001</u>	Jul 01, 2010
<u>AB</u>		PLIVA HRVATSKA DOO	<u>0.1%</u>	<u>A090962 001</u>	Jun 02, 2010
<u>AB</u>		TARO	<u>0.3%</u>	<u>A208322 001</u>	Jun 23, 2016
<u>AB</u>		TOLMAR	<u>0.3%</u>	<u>A200298 001</u>	Jun 14, 2012

DIFFERIN

<u>AB</u>	+	GALDERMA LABS LP	<u>0.3%</u>	<u>N021753 001</u>	Jun 19, 2007
	LOTION;TOPICAL				

DIFFERIN

+! GALDERMA LABS LP 0.1%

N022502 001 Mar 17, 2010

SOLUTION;TOPICAL

ADAPALENE

<u>AB</u>		CALL INC	<u>0.1%</u>	<u>A203981 001</u>	Sep 23, 2016
<u>AB</u>	!		<u>0.1%</u>	<u>A204593 001</u>	Jan 05, 2016

ADAPALENE; BENZOYL PEROXIDE

GEL;TOPICAL

ADAPALENE AND BENZOYL PEROXIDE

<u>AB</u>		ACTAVIS MID ATLANTIC	<u>0.1%:2.5%</u>	<u>A203790 001</u>	Sep 30, 2015
<u>AB</u>	!	GALDERMA LABS LP	<u>0.1%:2.5%</u>	<u>N022320 001</u>	Dec 08, 2008
	EPIDUO FORTE				

+! GALDERMA LABS LP 0.3%:2.5%

N207917 001 Jul 15, 2015

ADEFOVIR DIPIVOXIL

TABLET;ORAL

ADEFOVIR DIPIVOXIL

<u>AB</u>		SIGMAPHARM LABS LLC	<u>10MG</u>	<u>A202051 001</u>	Aug 29, 2013
<u>AB</u>	!	GILEAD	<u>10MG</u>	<u>N0211449 001</u>	Sep 20, 2002

ADENOSINE

INJECTABLE;INJECTION

ADENOCARD

<u>AP</u>	+	ASTELLAS	<u>3MG/ML</u>	<u>N019937 002</u>	Oct 30, 1989
<u>AP</u>			<u>3MG/ML</u>	<u>A078076 001</u>	Oct 31, 2008

ADENOSINE

<u>AP</u>		AKORN	<u>3MG/ML</u>	<u>A077133 001</u>	Apr 27, 2005
<u>AP</u>		FRESENIUS KABI USA	<u>3MG/ML</u>	<u>A077283 001</u>	Jun 14, 2007
<u>AP</u>		GLAND PHARMA LTD	<u>3MG/ML</u>	<u>A090010 001</u>	Apr 28, 2009
<u>AP</u>		LUITPOLD	<u>3MG/ML</u>	<u>A078640 001</u>	Mar 21, 2014
<u>AP</u>		MYLAN LABS LTD	<u>3MG/ML</u>	<u>A078686 001</u>	May 13, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-10 (of 436)

**ADENOSINE**

INJECTABLE; INJECTION

**ADENOSINE**

<b>AP</b>	WEST-WARD PHARMS INT	<b><u>3MG/ML</u></b>	<b>A076404 001</b>	Jun 16, 2004
<b>AP</b>	SOLUTION; IV (INFUSION)	<b><u>3MG/ML</u></b>	<b>A076500 001</b>	Jun 16, 2004
<b>AP</b>	AKORN	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A090450 001</b>	Oct 02, 2014
<b>AP</b>		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A090450 002</b>	Oct 02, 2014
<b>AP</b>	AUROBINDO PHARMA LTD	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A205331 001</b>	Nov 02, 2017
<b>AP</b>		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A205331 002</b>	Nov 02, 2017
<b>AP</b>	EMCURE PHARMS LTD	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A202313 001</b>	Sep 15, 2014
<b>AP</b>		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A202313 002</b>	Sep 15, 2014
<b>AP</b>	FRESENIUS KABI USA	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A077897 001</b>	Nov 27, 2017
<b>AP</b>		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A077897 002</b>	Nov 27, 2017
<b>AP</b>	HOSPIRA INC	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A203883 001</b>	Mar 24, 2014
<b>AP</b>		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A203883 002</b>	Mar 24, 2014
<b>AP</b>	SAGENT STRIDES	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A090212 001</b>	Mar 28, 2014
<b>AP</b>		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A090212 002</b>	Mar 28, 2014
<b>AP</b> !	TEVA PHARMS USA	<b><u>60MG/20ML (3MG/ML)</u></b>	<b>A077425 001</b>	Aug 29, 2013
<b>AP</b> !		<b><u>90MG/30ML (3MG/ML)</u></b>	<b>A077425 002</b>	Aug 29, 2013

**AFATINIB DIMALEATE**

TABLET; ORAL

GILOTRIF

+	BOEHRINGER INGELHEIM	EQ 20MG BASE	N201292 001	Jul 12, 2013
+		EQ 30MG BASE	N201292 002	Jul 12, 2013
+		EQ 40MG BASE	N201292 003	Jul 12, 2013

**ALBENDAZOLE**

TABLET; ORAL

ALBENZA

+	IMPAK LABS INC	200MG	N020666 001	Jun 11, 1996
---	----------------	-------	-------------	--------------

**ALBUMIN HUMAN**

INJECTABLE; INJECTION

OPTISON

+	GE HEALTHCARE	10MG/ML	N020899 001	Dec 31, 1997
---	---------------	---------	-------------	--------------

**ALBUMIN IODINATED I-125 SERUM**

INJECTABLE; INJECTION

JEANATOPE

+	ISO TEX	100uCi/10ML (10uCi/ML)	N017836 003	Jun 08, 2004
+		500uCi/0.5ML	N017836 001	
+		1,000uCi/ML	N017836 002	

**ALBUMIN IODINATED I-131 SERUM**

INJECTABLE; INJECTION

MEGATOPE

+	ISO TEX	0.5mCi/VIAL	N017837 001	
+		1mCi/VIAL	N017837 002	

**ALBUTEROL SULFATE**

AEROSOL, METERED; INHALATION

PROAIR HFA

BX	! TEVA BRANDED PHARM	EQ 0.09MG BASE/INH	N021457 001	Oct 29, 2004
BX	! 3M DRUG DELIVERY VENTOLIN HFA	EQ 0.09MG BASE/INH	N020503 001	Aug 15, 1996

BX	! GLAXOSMITHKLINE POWDER, METERED; INHALATION PROAIR RESPICLICK	EQ 0.09MG BASE/INH	N020983 001	Apr 19, 2001
			N205636 001	Mar 31, 2015

SOLUTION; INHALATION

ACCUNEB

<b>AN</b>	! MYLAN SPECIALITY LP	<b><u>EQ 0.021% BASE</u></b>	<b>N020949 002</b>	Apr 30, 2001
<b>AN</b>	!	<b><u>EQ 0.042% BASE</u></b>	<b>N020949 001</b>	Apr 30, 2001

**ALBUTEROL SULFATE**

<b>AN</b>	AUROBINDO PHARMA LTD	<b><u>EQ 0.083% BASE</u></b>	<b>A206224 001</b>	Oct 17, 2017
<b>AN</b>	BAUSCH AND LOMB	<b><u>EQ 0.5% BASE</u></b>	<b>A075050 001</b>	Jun 18, 1998
<b>AN</b>	HI TECH PHARMA	<b><u>EQ 0.5% BASE</u></b>	<b>A074543 001</b>	Jan 15, 1998
<b>AN</b>	NEPHRON	<b><u>EQ 0.021% BASE</u></b>	<b>A076355 002</b>	Mar 31, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-11 (of 436)

ALBUTEROL SULFATE

SOLUTION; INHALATION

**ALBUTEROL SULFATE**

<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A076355 001</u>	Jun 28, 2004
<u>AN</u>	!	<u>EQ 0.083% BASE</u>	<u>A074880 001</u>	Sep 17, 1997
<u>AN</u>		<u>EQ 0.5% BASE</u>	<u>A075664 001</u>	Jun 26, 2001
<u>AN</u>	RITEDOSE CORP	<u>EQ 0.083% BASE</u>	<u>A077839 001</u>	Dec 16, 2008
<u>AN</u>	SUN PHARMA GLOBAL	<u>EQ 0.083% BASE</u>	<u>A207857 001</u>	Jul 21, 2017
<u>AN</u>	WATSON LABS	<u>EQ 0.021% BASE</u>	<u>A077772 001</u>	Sep 25, 2007
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A077772 002</u>	Sep 25, 2007

SYRUP; ORAL

**ALBUTEROL SULFATE**

<u>AA</u>	AMNEAL PHARMS	<u>EQ 2MG BASE/5ML</u>	<u>A079241 001</u>	May 12, 2010
<u>AA</u>	G AND W LABS INC	<u>EQ 2MG BASE/5ML</u>	<u>A074454 001</u>	Sep 25, 1995
<u>AA</u>	HI TECH PHARMA	<u>EQ 2MG BASE/5ML</u>	<u>A074749 001</u>	Jan 30, 1998
<u>AA</u>	! TEVA	<u>EQ 2MG BASE/5ML</u>	<u>A073419 001</u>	Mar 30, 1992
<u>AA</u>	VINTAGE	<u>EQ 2MG BASE/5ML</u>	<u>A078105 001</u>	Dec 27, 2006
<u>AA</u>	VISTAPHARM	<u>EQ 2MG BASE/5ML</u>	<u>A077788 001</u>	Jun 26, 2007

TABLET; ORAL

**ALBUTEROL SULFATE**

<u>AB</u>	MYLAN	<u>EQ 2MG BASE</u>	<u>A072894 002</u>	Jan 17, 1991
<u>AB</u>	!	<u>EQ 4MG BASE</u>	<u>A072894 001</u>	Jan 17, 1991
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 2MG BASE</u>	<u>A072637 002</u>	Dec 05, 1989
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A072637 001</u>	Dec 05, 1989

TABLET, EXTENDED RELEASE; ORAL

**ALBUTEROL SULFATE**

<u>AB</u>	MYLAN	<u>EQ 4MG BASE</u>	<u>A078092 002</u>	Jan 29, 2007
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A078092 001</u>	Jan 29, 2007

**VOSPIRE ER**

<u>AB</u>	DAVA PHARMS INC	<u>EQ 4MG BASE</u>	<u>A076130 002</u>	Sep 26, 2002
<u>AB</u>	!	<u>EQ 8MG BASE</u>	<u>A076130 003</u>	Sep 26, 2002

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

**ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE**

<u>AN</u>	CIPLA LTD	<u>EQ 0.083% BASE;0.017%</u>	<u>A077559 001</u>	Dec 31, 2007
<u>AN</u>	NEPHRON	<u>EQ 0.083% BASE;0.017%</u>	<u>A076749 001</u>	Dec 31, 2007
<u>AN</u>	RITEDOSE CORP	<u>EQ 0.083% BASE;0.017%</u>	<u>A202496 001</u>	Oct 01, 2012
<u>AN</u>	SUN PHARMA GLOBAL	<u>EQ 0.083% BASE;0.017%</u>	<u>A207875 001</u>	Aug 07, 2017
<u>AN</u>	TEVA PHARMS	<u>EQ 0.083% BASE;0.017%</u>	<u>A076724 001</u>	Dec 31, 2007
<u>AN</u>	WATSON LABS TEVA	<u>EQ 0.083% BASE;0.017%</u>	<u>A077063 001</u>	Dec 31, 2007

SPRAY, METERED; INHALATION

COMBIVENT RESPIMAT

+!	BOEHRINGER INGELHEIM	EQ 0.1MG BASE/INH;0.02MG/INH	N021747 001	Oct 07, 2011
----	----------------------	------------------------------	-------------	--------------

ALCAFTADINE

SOLUTION/DROPS; OPHTHALMIC

LASTACRAFT

+!	ALLERGAN	0.25%	N022134 001	Jul 28, 2010
----	----------	-------	-------------	--------------

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

**ALCLOMETASONE DIPROPIONATE**

<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A076973 001</u>	Jul 12, 2005
<u>AB</u>	GLENMARK GENERICS	<u>0.05%</u>	<u>A079061 001</u>	Jun 23, 2009
<u>AB</u>	TARO	<u>0.05%</u>	<u>A076587 001</u>	Sep 15, 2005

OINTMENT; TOPICAL

**ALCLOMETASONE DIPROPIONATE**

<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A076884 001</u>	Jul 18, 2005
<u>AB</u>	GLENMARK GENERICS	<u>0.05%</u>	<u>A079227 001</u>	Jul 30, 2009
<u>AB</u>	TARO	<u>0.05%</u>	<u>A076730 001</u>	Jul 29, 2004

ALECTINIB HYDROCHLORIDE

CAPSULE; ORAL

ALECENSA

+!	HOFFMANN-LA ROCHE	EQ 150MG BASE	N208434 001	Dec 11, 2015
----	-------------------	---------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-12 (of 436)

ALENDRONATE SODIUM

SOLUTION;ORAL

ALENDRONATE SODIUM

! WEST-WARD PHARMS  
INT

TABLET;ORAL

ALENDRONATE SODIUM

<u>AB</u>	APOTEX	<u>EQ 5MG BASE</u>	<u>A077982 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077982 002</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A077982 003</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A077982 004</u>	Aug 04, 2008
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A090124 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A090124 002</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A090124 003</u>	Aug 04, 2008
<u>AB</u>	AUSTARPHARMA LLC	<u>EQ 5MG BASE</u>	<u>A090258 001</u>	Sep 24, 2009
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090258 002</u>	Sep 24, 2009
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A090258 003</u>	Sep 24, 2009
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A090258 004</u>	Sep 24, 2009
<u>AB</u>	CIPLA LTD	<u>EQ 5MG BASE</u>	<u>A076768 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076768 002</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A076768 003</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076768 004</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A076768 005</u>	Aug 04, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 5MG BASE</u>	<u>A079049 003</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079049 004</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A079049 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A079049 002</u>	Aug 04, 2008
<u>AB</u>	IMPAX LABS INC	<u>EQ 5MG BASE</u>	<u>A075710 001</u>	Feb 06, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075710 002</u>	Feb 06, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A075710 003</u>	Feb 06, 2008
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A075710 004</u>	Feb 06, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A075710 005</u>	Feb 06, 2008
<u>AB</u>	JUBILANT CADISTA	<u>EQ 5MG BASE</u>	<u>A090557 001</u>	Feb 18, 2010
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090557 002</u>	Feb 18, 2010
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A090557 003</u>	Feb 18, 2010
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A090557 004</u>	Feb 18, 2010
<u>AB</u>	MYLAN	<u>EQ 5MG BASE</u>	<u>A076584 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076584 002</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A076584 003</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A076584 004</u>	Aug 04, 2008
<u>AB</u>	SUN PHARMA GLOBAL	<u>EQ 5MG BASE</u>	<u>A090022 001</u>	Sep 10, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090022 002</u>	Sep 10, 2008
<u>AB</u>		<u>EQ 35MG BASE</u>	<u>A090022 003</u>	Sep 10, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A090022 004</u>	Sep 10, 2008
<u>AB</u>	WATSON LABS	<u>EQ 35MG BASE</u>	<u>A076984 001</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076984 002</u>	Aug 04, 2008
<u>AB</u>		<u>EQ 70MG BASE</u>	<u>A076984 003</u>	Aug 04, 2008

**FOSAMAX**

<u>AB</u>	+! MERCK AND CO INC	<u>EQ 70MG BASE</u>	<u>N020560 005</u>	Oct 20, 2000
-----------	---------------------	---------------------	--------------------	--------------

TABLET, EFFERVESCENT;ORAL

BINOSTO

+! MISSION PHARMA

EQ 70MG BASE

N202344 001 Mar 12, 2012

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET;ORAL

FOSAMAX PLUS D

+ MERCK

EQ 70MG BASE;2,800 IU

N021762 001 Apr 07, 2005

+!

EQ 70MG BASE;5,600 IU

N021762 002 Apr 26, 2007

ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTA

<u>AP</u>	+! AKORN	<u>EQ 0.5MG BASE/ML</u>	<u>N019353 001</u>	Dec 29, 1986
-----------	----------	-------------------------	--------------------	--------------

ALFENTANIL

<u>AP</u>	HOSPIRA	<u>EQ 0.5MG BASE/ML</u>	<u>A075221 001</u>	Oct 28, 1999
-----------	---------	-------------------------	--------------------	--------------

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALFUZOSIN HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A079013 001</u>	Jul 18, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A079060 001</u>	Aug 30, 2012
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A090284 001</u>	Jan 17, 2012
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A079014 001</u>	Jul 18, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-13 (of 436)

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALFUZOSIN HYDROCHLORIDE

<u>AB</u>	SUN PHARMA GLOBAL	<u>10MG</u>	<u>A079057 001</u>	Jul 18, 2011
<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>A079056 001</u>	Jul 18, 2011
<u>AB</u>	TORRENT PHARMS	<u>10MG</u>	<u>A079054 001</u>	Jul 18, 2011
<u>AB</u>	UNICHEM LABS LTD	<u>10MG</u>	<u>A203192 001</u>	Jan 28, 2016

UROXATRAL

<u>AB</u>	+!	CONCORDIA PHARMS INC	<u>10MG</u>	<u>N021287 001</u>	Jun 12, 2003
-----------	----	----------------------	-------------	--------------------	--------------

ALISKIREN HEMIFUMARATE

CAPSULE, PELLET; ORAL  
 TEKTURNA

+!	NODEN PHARMA	EQ 37.5MG BASE	N210709 001	Nov 14, 2017
TABLET; ORAL TEKTURNA				
+!	NODEN PHARMA	EQ 150MG BASE	N021985 001	Mar 05, 2007

+!	NODEN PHARMA	EQ 300MG BASE	N021985 002	Mar 05, 2007
----	--------------	---------------	-------------	--------------

ALISKIREN HEMIFUMARATE; HYDROCHLORTIAZIDE

TABLET; ORAL

TEKTURNA HCT				
+!	NODEN PHARMA	EQ 150MG BASE;12.5MG	N022107 001	Jan 18, 2008
+		EQ 150MG BASE;25MG	N022107 002	Jan 18, 2008
+		EQ 300MG BASE;12.5MG	N022107 003	Jan 18, 2008
+		EQ 300MG BASE;25MG	N022107 004	Jan 18, 2008

ALITRETINOIN

GEL; TOPICAL  
 PANRETIN

+!	EISAI INC	EQ 0.1% BASE	N020886 001	Feb 02, 1999
----	-----------	--------------	-------------	--------------

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

<u>AB</u>	ACCORD HLTHCARE	<u>100MG</u>	<u>A203154 001</u>	May 06, 2013
<u>AB</u>		<u>300MG</u>	<u>A203154 002</u>	May 06, 2013
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077353 001</u>	Sep 08, 2005
<u>AB</u>		<u>300MG</u>	<u>A077353 002</u>	Sep 08, 2005
<u>AB</u>	INDOCO REMEDIES	<u>100MG</u>	<u>A204467 001</u>	Jul 28, 2016
<u>AB</u>		<u>300MG</u>	<u>A204467 002</u>	Jul 28, 2016
<u>AB</u>	IPCA LABS LTD	<u>100MG</u>	<u>A090637 001</u>	Mar 16, 2011
<u>AB</u>		<u>300MG</u>	<u>A090637 002</u>	Mar 16, 2011
<u>AB</u>	MYLAN	<u>100MG</u>	<u>A018659 001</u>	Oct 24, 1986
<u>AB</u>		<u>300MG</u>	<u>A018659 002</u>	Oct 24, 1986
<u>AB</u>	NORTHSTAR HLTHCARE	<u>100MG</u>	<u>A078253 001</u>	Sep 11, 2007
<u>AB</u>		<u>300MG</u>	<u>A078253 002</u>	Sep 11, 2007
<u>AB</u>	SUN PHARM INDNS INC	<u>100MG</u>	<u>A078390 001</u>	Aug 30, 2007
<u>AB</u>		<u>300MG</u>	<u>A078390 002</u>	Aug 30, 2007
<u>AB</u>	SUN PHARM INDUSTRIES	<u>100MG</u>	<u>A071450 002</u>	Jan 09, 1987
<u>AB</u>		<u>300MG</u>	<u>A071450 001</u>	Jan 09, 1987
<u>AB</u>	VINTAGE PHARMS	<u>100MG</u>	<u>A075798 001</u>	Jun 27, 2003
<u>AB</u>		<u>300MG</u>	<u>A075798 002</u>	Jun 27, 2003
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N018832 002</u>	Sep 28, 1984
<u>AB</u>		<u>300MG</u>	<u>N018877 001</u>	Sep 28, 1984
<u>AB</u>	ZYDUS PHARMS USA INC	<u>100MG</u>	<u>A210117 001</u>	Oct 12, 2017
<u>AB</u>		<u>300MG</u>	<u>A210117 002</u>	Oct 12, 2017
<u>AB</u>	<u>LOPURIN</u>			
<u>AB</u>	DR REDDYS LA	<u>100MG</u>	<u>A071586 001</u>	Apr 02, 1987
<u>AB</u>		<u>300MG</u>	<u>A071587 001</u>	Apr 02, 1987
<u>AB</u>	<u>ZYOPRIM</u>			
<u>AB</u>	+ CASPER PHARMA LLC	<u>100MG</u>	<u>N016084 001</u>	
<u>AB</u>	+!	<u>300MG</u>	<u>N016084 002</u>	

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALLOPURINOL SODIUM

<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 500MG BASE/VIAL</u>	<u>A076870 001</u>	Aug 26, 2004
<u>AP</u>	+!	MYLAN INSTITUTIONAL	<u>EQ 500MG BASE/VIAL</u>	<u>N020298 001</u>

May 17, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-14 (of 436)

ALLOPURINOL; LESINURAD

TABLET;ORAL

DUZALLO

+ IRONWOOD PHARMS INC 200MG;200MG  
+! 300MG;200MG

N209203 001 Aug 18, 2017  
N209203 002 Aug 18, 2017

ALMOTRIPTAN MALATE

TABLET;ORAL

**ALMOTRIPTAN MALATE**

<b>AB</b>	AJANTA PHARMA LTD	<b>EQ 6.25MG BASE</b>
<b>AB</b>		<b>EQ 12.5MG BASE</b>
<b>AB</b>	MYLAN PHARMS INC	<b>EQ 6.25MG BASE</b>
<b>AB</b>		<b>EQ 12.5MG BASE</b>
<b>AB</b>	TEVA PHARMS USA	<b>EQ 6.25MG BASE</b>
<b>AB</b>		<b>EQ 12.5MG BASE</b>

**A205523 001** Mar 03, 2016  
**A205523 002** Mar 03, 2016  
**A205171 001** Nov 09, 2015  
**A205171 002** Nov 09, 2015  
**A078027 001** Jul 07, 2015  
**A078027 002** Jul 07, 2015

**AXERT**

<b>AB</b> +	JANSSEN PHARMS	<b>EQ 6.25MG BASE</b>
<b>AB</b> +!		<b>EQ 12.5MG BASE</b>

**N021001 001** May 07, 2001  
**N021001 002** May 07, 2001

ALOGLIPTIN BENZOATE

TABLET;ORAL

NESINA

+ TAKEDA PHARMS USA EQ 6.25MG BASE  
+ EQ 12.5MG BASE  
+! EQ 25MG BASE

N022271 001 Jan 25, 2013  
N022271 002 Jan 25, 2013  
N022271 003 Jan 25, 2013

ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE

TABLET;ORAL

KAZANO

+ TAKEDA PHARMS USA EQ 12.5MG BASE;500MG  
+! EQ 12.5MG BASE;1GM

N203414 001 Jan 25, 2013  
N203414 002 Jan 25, 2013

ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL

OSENI

+ TAKEDA PHARMS USA EQ 12.5MG BASE;EQ 15MG BASE  
+ EQ 12.5MG BASE;EQ 30MG BASE  
+ EQ 12.5MG BASE;EQ 45MG BASE  
+ EQ 25MG BASE;EQ 15MG BASE  
+ EQ 25MG BASE;EQ 30MG BASE  
+! EQ 25MG BASE;EQ 45MG BASE

N022426 004 Jan 25, 2013  
N022426 005 Jan 25, 2013  
N022426 006 Jan 25, 2013  
N022426 001 Jan 25, 2013  
N022426 002 Jan 25, 2013  
N022426 003 Jan 25, 2013

ALOSETRON HYDROCHLORIDE

TABLET;ORAL

**ALOSETRON HYDROCHLORIDE**

<b>AB</b>	AMNEAL PHARMS	<b>EQ 0.5MG BASE</b>
<b>AB</b>		<b>EQ 1MG BASE</b>
<b>AB</b>	WEST-WARD PHARMS INT	<b>EQ 0.5MG BASE</b>
<b>AB</b>		<b>EQ 1MG BASE</b>
<b>AB</b>	<b><u>IOTRONEX</u></b>	
<b>AB</b> +	SEBELA IRELAND LTD	<b>EQ 0.5MG BASE</b>
<b>AB</b> +!		<b>EQ 1MG BASE</b>

**A206647 001** Dec 22, 2016  
**A206647 002** Dec 22, 2016  
**A200652 001** May 04, 2015  
**A200652 002** May 04, 2015  
**N021107 002** Dec 23, 2003  
**N021107 001** Feb 09, 2000

ALPHA-TOCOPHEROL ACETATE; ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN K

INJECTABLE;INJECTION

INFUVITE ADULT

+! SANDOZ INC 2 IU/ML;40MG/ML;12MCG/ML;40 IU/ML;1MCG/ML;3MG/ML;120MCG/ML;8MG/ML;1 .2MG/ML;0.72MG/ML;1.2MG/ML;660 IU/ML;0.03MG/ML

N021163 001 May 18, 2000

INJECTABLE;IV (INFUSION)

INFUVITE ADULT

+! SANDOZ INC 2 IU/ML;40MG/ML;12MCG/ML;40 IU/ML;1MCG/ML;3MG/ML;120MCG/ML;8MG/ML;1 .2MG/ML;0.72MG/ML;1.2MG/ML;660 IU/ML;30MCG/ML

N021559 001 Jun 16, 2003

ALPRAZOLAM

CONCENTRATE;ORAL

ALPRAZOLAM

! WEST-WARD PHARMS INT 1MG/ML

A074312 001 Oct 31, 1993

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-15 (of 436)

**ALPRAZOLAM**

TABLET;ORAL

**ALPRAZOLAM**

<b><u>AB</u></b>	ACTAVIS ELIZABETH	<b><u>0 .25MG</u></b>	<b><u>A074342 001</u></b>	Oct 31, 1993
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A074342 002</u></b>	Oct 31, 1993
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A074342 003</u></b>	Oct 31, 1993
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A074342 004</u></b>	Oct 31, 1993
<b><u>AB</u></b>	APOTEX INC	<b><u>0 .25MG</u></b>	<b><u>A077741 001</u></b>	Jan 19, 2007
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A077741 002</u></b>	Jan 19, 2007
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A077741 003</u></b>	Jan 19, 2007
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A077741 004</u></b>	Jan 19, 2007
<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>0 .25MG</u></b>	<b><u>A203346 001</u></b>	Jul 31, 2015
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A203346 002</u></b>	Jul 31, 2015
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A203346 003</u></b>	Jul 31, 2015
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A203346 004</u></b>	Jul 31, 2015
<b><u>AB</u></b>	DAVA INTL INC	<b><u>0 .25MG</u></b>	<b><u>A074174 001</u></b>	Oct 19, 1993
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A074174 002</u></b>	Oct 19, 1993
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A074174 003</u></b>	Oct 19, 1993
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A074174 004</u></b>	Oct 19, 1993
<b><u>AB</u></b>	MYLAN	<b><u>0 .25MG</u></b>	<b><u>A074215 001</u></b>	Jan 27, 1994
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A074215 002</u></b>	Jan 27, 1994
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A074215 003</u></b>	Jan 27, 1994
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A074215 004</u></b>	Jan 27, 1994
<b><u>AB</u></b>	NATCO PHARMA LTD	<b><u>0 .25MG</u></b>	<b><u>A200739 001</u></b>	Apr 15, 2015
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A200739 002</u></b>	Apr 15, 2015
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A200739 003</u></b>	Apr 15, 2015
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A200739 004</u></b>	Apr 15, 2015
<b><u>AB</u></b>	SANDOZ	<b><u>0 .25MG</u></b>	<b><u>A074112 001</u></b>	Dec 29, 1995
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A074112 002</u></b>	Dec 29, 1995
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A074112 003</u></b>	Dec 29, 1995
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A074909 001</u></b>	Mar 25, 1998
<b><u>AB</u></b>	SUN PHARMA GLOBAL	<b><u>0 .25MG</u></b>	<b><u>A090082 001</u></b>	Jun 17, 2010
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A090082 002</u></b>	Jun 17, 2010
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A090082 003</u></b>	Jun 17, 2010
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A090082 004</u></b>	Jun 17, 2010
<b><u>AB</u></b>	VINTAGE	<b><u>0 .25MG</u></b>	<b><u>A078491 001</u></b>	Sep 25, 2008
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A078491 002</u></b>	Sep 25, 2008
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A078491 003</u></b>	Sep 25, 2008
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A078491 004</u></b>	Dec 12, 2008
<b><u>AB</u></b>	VINTAGE PHARMS	<b><u>0 .25MG</u></b>	<b><u>A090248 001</u></b>	Sep 17, 2010
<b><u>AB</u></b>		<b><u>0 .5MG</u></b>	<b><u>A090248 002</u></b>	Sep 17, 2010
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A090248 003</u></b>	Sep 17, 2010
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A090248 004</u></b>	Sep 17, 2010

**XANAX**

<b><u>AB</u></b>	+	PHARMACIA AND UPJOHN	<b><u>0 .25MG</u></b>	<b><u>N018276 001</u></b>
<b><u>AB</u></b>	+		<b><u>0 .5MG</u></b>	<b><u>N018276 002</u></b>
<b><u>AB</u></b>	+!		<b><u>1MG</u></b>	<b><u>N018276 003</u></b>
<b><u>AB</u></b>	+		<b><u>2MG</u></b>	<b><u>N018276 004</u></b>

TABLET, EXTENDED RELEASE;ORAL

**ALPRAZOLAM**

<b><u>AB</u></b>	ACTAVIS ELIZABETH	<b><u>0 .5MG</u></b>	<b><u>A078056 001</u></b>	Feb 13, 2007
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A078056 002</u></b>	Feb 13, 2007
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A078056 003</u></b>	Feb 13, 2007
<b><u>AB</u></b>		<b><u>3MG</u></b>	<b><u>A078056 004</u></b>	Feb 13, 2007
<b><u>AB</u></b>	AMNEAL PHARMS NY	<b><u>0 .5MG</u></b>	<b><u>A078387 001</u></b>	May 30, 2008
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A078387 002</u></b>	May 30, 2008
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A078387 003</u></b>	May 30, 2008
<b><u>AB</u></b>		<b><u>3MG</u></b>	<b><u>A078387 004</u></b>	May 30, 2008
<b><u>AB</u></b>	ANCHEM PHARMS	<b><u>0 .5MG</u></b>	<b><u>A078469 001</u></b>	Sep 29, 2011
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A078469 002</u></b>	Sep 29, 2011
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A078469 003</u></b>	Sep 29, 2011
<b><u>AB</u></b>		<b><u>3MG</u></b>	<b><u>A078469 004</u></b>	Sep 29, 2011
<b><u>AB</u></b>	ANI PHARMS INC	<b><u>0 .5MG</u></b>	<b><u>A077725 001</u></b>	Jul 31, 2006
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A077725 002</u></b>	Jul 31, 2006
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A077725 004</u></b>	Jul 31, 2006
<b><u>AB</u></b>		<b><u>3MG</u></b>	<b><u>A077725 003</u></b>	Jul 31, 2006
<b><u>AB</u></b>	APOTEX INC	<b><u>0 .5MG</u></b>	<b><u>A078449 001</u></b>	Nov 12, 2008
<b><u>AB</u></b>		<b><u>1MG</u></b>	<b><u>A078449 004</u></b>	Dec 23, 2015
<b><u>AB</u></b>		<b><u>2MG</u></b>	<b><u>A078449 002</u></b>	Nov 12, 2008
<b><u>AB</u></b>		<b><u>3MG</u></b>	<b><u>A078449 003</u></b>	Nov 12, 2008
<b><u>AB</u></b>	AUROBINDO PHARMA	<b><u>0 .5MG</u></b>	<b><u>A090871 001</u></b>	Jun 07, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-16 (of 436)

**ALPRAZOLAM**

TABLET, EXTENDED RELEASE; ORAL

**ALPRAZOLAM**  
LTD

<b><u>AB</u></b>	<b><u>1MG</u></b>	<b><u>A090871 002</u></b>	Jun 07, 2011
<b><u>AB</u></b>	<b><u>2MG</u></b>	<b><u>A090871 003</u></b>	Jun 07, 2011
<b><u>AB</u></b>	<b><u>3MG</u></b>	<b><u>A090871 004</u></b>	Jun 07, 2011
<b><u>AB</u></b> HERITAGE PHARMS INC	<b><u>0 . 5MG</u></b>	<b><u>A078489 001</u></b>	Oct 17, 2008
<b><u>AB</u></b>	<b><u>1MG</u></b>	<b><u>A078489 002</u></b>	Oct 17, 2008
<b><u>AB</u></b>	<b><u>2MG</u></b>	<b><u>A078489 003</u></b>	Oct 17, 2008
<b><u>AB</u></b>	<b><u>3MG</u></b>	<b><u>A078489 004</u></b>	Oct 17, 2008

**XANAX XR**

<b><u>AB</u></b> + PHARMACIA AND UPJOHN	<b><u>0 . 5MG</u></b>	<b><u>N021434 001</u></b>	Jan 17, 2003
<b><u>AB</u></b> +	<b><u>1MG</u></b>	<b><u>N021434 002</u></b>	Jan 17, 2003
<b><u>AB</u></b> +	<b><u>2MG</u></b>	<b><u>N021434 003</u></b>	Jan 17, 2003
<b><u>AB</u></b> +!	<b><u>3MG</u></b>	<b><u>N021434 004</u></b>	Jan 17, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

**ALPRAZOLAM**

<b><u>AB</u></b> ACTAVIS ELIZABETH	<b><u>0 . 25MG</u></b>	<b><u>A078561 001</u></b>	Mar 16, 2010
<b><u>AB</u></b>	<b><u>0 . 5MG</u></b>	<b><u>A078561 002</u></b>	Mar 16, 2010
<b><u>AB</u></b>	<b><u>1MG</u></b>	<b><u>A078561 003</u></b>	Mar 16, 2010
<b><u>AB</u></b>	<b><u>2MG</u></b>	<b><u>A078561 004</u></b>	Mar 16, 2010
<b><u>AB</u></b> PAR PHARM	<b><u>0 . 25MG</u></b>	<b><u>A078088 001</u></b>	Jan 09, 2009
<b><u>AB</u></b>	<b><u>0 . 5MG</u></b>	<b><u>A078088 002</u></b>	Jan 09, 2009
<b><u>AB</u></b> !	<b><u>1MG</u></b>	<b><u>A078088 003</u></b>	Jan 09, 2009
<b><u>AB</u></b>	<b><u>2MG</u></b>	<b><u>A078088 004</u></b>	Jan 09, 2009

**ALPROSTADIL**

INJECTABLE; INJECTION

**ALPROSTADIL**

<b><u>AP</u></b> TEVA PHARMS USA	<b><u>0 . 5MG/ML</u></b>	<b><u>A075196 001</u></b>	Apr 30, 1999
<b><u>AP</u></b> WEST-WARD PHARMS INT	<b><u>0 . 5MG/ML</u></b>	<b><u>A074815 001</u></b>	Jan 20, 1998

**CAVERJECT**

<b><u>AP</u></b> + PHARMACIA AND UPJOHN	<b><u>0 . 01MG/VIAL</u></b>	<b><u>N020379 001</u></b>	Jul 06, 1995
<b><u>AP</u></b> +!	<b><u>0 . 02MG/VIAL</u></b>	<b><u>N020379 002</u></b>	Jul 06, 1995
<b><u>AP</u></b> +!	<b><u>0 . 04MG/VIAL</u></b>	<b><u>N020379 004</u></b>	May 19, 1997

**EDEX**

<b><u>AP</u></b> + AUXILIUM PHARMS LLC	<b><u>0 . 01MG/VIAL</u></b>	<b><u>N020649 002</u></b>	Jun 12, 1997
<b><u>AP</u></b> +	<b><u>0 . 02MG/VIAL</u></b>	<b><u>N020649 003</u></b>	Jun 12, 1997
<b><u>AP</u></b> +!	<b><u>0 . 04MG/VIAL</u></b>	<b><u>N020649 004</u></b>	Jun 12, 1997

**PROSTIN VR PEDIATRIC**

<b><u>AP</u></b> +! PHARMACIA AND UPJOHN	<b><u>0 . 5MG/ML</u></b>	<b><u>N018484 001</u></b>	
CAVERJECT			

+ PHARMACIA AND UPJOHN

0.005MG/VIAL

N020379 003 Jun 27, 1996

CAVERJECT IMPULSE

PHARMACIA AND UPJOHN

0.01MG/VIAL

N021212 001 Jun 11, 2002

0.02MG/VIAL

N021212 002 Jun 11, 2002

**EDEX**

+! AUXILIUM PHARMS LLC

0.01MG/VIAL

N020649 005 Jul 30, 1998

+!

0.02MG/VIAL

N020649 006 Jul 30, 1998

+!

0.04MG/VIAL

N020649 007 Jul 30, 1998

SUPPOSITORY; URETHRAL

MUSE

+! MYLAN SPECIALITY LP

0.125MG

N020700 001 Nov 19, 1996

+!

0.25MG

N020700 002 Nov 19, 1996

+!

0.5MG

N020700 003 Nov 19, 1996

+!

1MG

N020700 004 Nov 19, 1996

**ALTRETAMINE**

CAPSULE; ORAL

HEXALEN

+! EISAI INC

50MG

N019926 001 Dec 26, 1990

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-17 (of 436)

ALVIMOPAN

CAPSULE;ORAL  
 ENTEREG  
 +! CUBIST PHARMS 12MG N021775 001 May 20, 2008

AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

**AMANTADINE HYDROCHLORIDE**

<b>AB</b>	ALEMBIC PHARMS LTD	<b><u>100MG</u></b>	<b><u>A208966 001</u></b>	Jun 21, 2017
<b>AB</b>	BIONPHARMA INC	<b><u>100MG</u></b>	<b><u>A078720 001</u></b>	May 29, 2008
<b>AB</b>	HERITAGE PHARMA	<b><u>100MG</u></b>	<b><u>A209171 001</u></b>	Jun 12, 2017
<b>AB</b>	LANNETT HOLDINGS INC	<b><u>100MG</u></b>	<b><u>A209221 001</u></b>	Jun 15, 2017
<b>AB</b>	NEWGEN PHARMS LLC	<b><u>100MG</u></b>	<b><u>A207570 001</u></b>	Sep 30, 2016
<b>AB</b> !	SANDOZ	<b><u>100MG</u></b>	<b><u>A071293 001</u></b>	Feb 18, 1987
<b>AB</b>	STRIDES PHARMA	<b><u>100MG</u></b>	<b><u>A209047 001</u></b>	Jun 07, 2017
<b>AB</b>	USL PHARMA	<b><u>100MG</u></b>	<b><u>A070589 001</u></b>	Aug 05, 1986
<b>AB</b>	WATSON LABS INC	<b><u>100MG</u></b>	<b><u>A208107 001</u></b>	Dec 06, 2016
<b>AB</b>	ZYDUS PHARMS USA INC	<b><u>100MG</u></b>	<b><u>A208278 001</u></b>	May 31, 2016

CAPSULE, EXTENDED RELEASE;ORAL

GOCOVRI

+ ADAMAS PHARMA	EQ 68.5MG BASE	N208944 001 Aug 24, 2017
+!	EQ 137MG BASE	N208944 002 Aug 24, 2017

SYRUP;ORAL

**AMANTADINE HYDROCHLORIDE**

<b>AA</b> !	CMP PHARMA INC	<b><u>50MG/5ML</u></b>	<b><u>A075819 001</u></b>	Sep 11, 2002
<b>AA</b> !	HI TECH PHARMA	<b><u>50MG/5ML</u></b>	<b><u>A074170 001</u></b>	Oct 28, 1994
<b>AA</b> !	MIKART	<b><u>50MG/5ML</u></b>	<b><u>A074028 001</u></b>	Jun 28, 1993
<b>AA</b> !	PHARM ASSOC	<b><u>50MG/5ML</u></b>	<b><u>A074509 001</u></b>	Jul 17, 1995
<b>AA</b> !	WOCHARDT BIO AG	<b><u>50MG/5ML</u></b>	<b><u>A075060 001</u></b>	Dec 24, 1998

TABLET;ORAL

**AMANTADINE HYDROCHLORIDE**

<b>AB</b>	NEWGEN PHARMS LLC	<b><u>100MG</u></b>	<b><u>A207571 001</u></b>	Jan 31, 2017
<b>AB</b>	STRIDES PHARMA	<b><u>100MG</u></b>	<b><u>A209035 001</u></b>	Jun 09, 2017
<b>AB</b> !	USL PHARMA	<b><u>100MG</u></b>	<b><u>A076186 001</u></b>	Dec 16, 2002
<b>AB</b>	WATSON LABS INC	<b><u>100MG</u></b>	<b><u>A208096 001</u></b>	Dec 15, 2016

AMBRISENTAN

TABLET;ORAL

LETAIRIS

+ GILEAD	5MG	N022081 001 Jun 15, 2007
+!	10MG	N022081 002 Jun 15, 2007

AMCINONIDE

CREAM;TOPICAL

**AMCINONIDE**

<b>AB</b> !	FOUGERA PHARMS	<b><u>0.1%</u></b>	<b><u>A076065 001</u></b>	May 15, 2003
<b>AB</b>	TARO PHARM INDS	<b><u>0.1%</u></b>	<b><u>A076229 001</u></b>	May 31, 2002

LOTION;TOPICAL

AMCINONIDE

! FOUGERA PHARMS	0.1%	A076329 001 Nov 06, 2002
------------------	------	--------------------------

OINTMENT;TOPICAL

**AMCINONIDE**

<b>AB</b> !	FOUGERA PHARMS	<b><u>0.1%</u></b>	<b><u>A076096 001</u></b>	Nov 19, 2002
<b>AB</b>	TARO PHARM INDS	<b><u>0.1%</u></b>	<b><u>A076367 001</u></b>	Mar 19, 2003

AMIFOSTINE

INJECTABLE;INJECTION

**AMIFOSTINE**

<b>AP</b>	MYLAN LABS LTD	<b><u>500MG/VIAL</u></b>	<b><u>A204363 001</u></b>	Jul 17, 2017
<b>AP</b>	SUN PHARMA GLOBAL	<b><u>500MG/VIAL</u></b>	<b><u>A077126 001</u></b>	Mar 14, 2008
<b>ETHYOL</b>				
<b>AP</b> +!	CLINIGEN HLTHCARE	<b><u>500MG/VIAL</u></b>	<b><u>N020221 001</u></b>	Dec 08, 1995

AMIKACIN SULFATE

INJECTABLE;INJECTION

**AMIKACIN SULFATE**

<b>AP</b> !	EMCURE PHARMS LTD	<b><u>EQ 250MG BASE/ML</u></b>	<b><u>A204040 001</u></b>	Dec 12, 2013
<b>AP</b>	FRESENIUS KABI USA	<b><u>EQ 50MG BASE/ML</u></b>	<b><u>A205605 001</u></b>	Dec 09, 2015
<b>AP</b>		<b><u>EQ 250MG BASE/ML</u></b>	<b><u>A205604 001</u></b>	Dec 09, 2015
<b>AP</b>	SAGENT PHARMS	<b><u>EQ 250MG BASE/ML</u></b>	<b><u>A203323 001</u></b>	May 12, 2016
<b>AP</b>	TEVA PHARMS USA	<b><u>EQ 250MG BASE/ML</u></b>	<b><u>A064045 002</u></b>	Sep 28, 1993
<b>AP</b> !	WEST-WARD PHARMS INT	<b><u>EQ 50MG BASE/ML</u></b>	<b><u>A063313 001</u></b>	Apr 11, 1994

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-18 (of 436)

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

AP

EQ 250MG BASE/ML

A063315 001 Apr 11, 1994

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

AB ! PAR PHARM

5MG

A070346 001 Jan 22, 1986

AB SIGMAPHARM LABS LLC

5MG

A079133 001 Jan 30, 2009

AB WINDLAS HLTHCARE

5MG

A204180 001 Aug 07, 2015

MIDAMOR

AB + PADDOCK LLC

5MG

N018200 001

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB BARR

EQ 5MG ANHYDROUS:50MG

A071111 001 May 10, 1988

AB ! MYLAN

EQ 5MG ANHYDROUS:50MG

A073209 001 Oct 31, 1991

AMINO ACIDS

INJECTABLE; INJECTION

AMINO ACIDS

B BRAUN

15% (150GM/1000ML)

A091112 001 Apr 13, 2012

15% (300GM/2000ML)

A091112 002 Apr 13, 2012

AMINOSYN 10%

ICU MEDICAL INC

10% (10GM/100ML)

N017673 003

AMINOSYN 10% (PH6)

ICU MEDICAL INC

10% (10GM/100ML)

N017673 008 Nov 18, 1985

AMINOSYN 3.5%

ICU MEDICAL INC

3.5% (3.5GM/100ML)

N017789 004

AMINOSYN 5%

ICU MEDICAL INC

5% (5GM/100ML)

N017673 001

AMINOSYN 7%

ICU MEDICAL INC

7% (7GM/100ML)

N017673 002

AMINOSYN 7% (PH6)

ICU MEDICAL INC

7% (7GM/100ML)

N017673 006 Nov 18, 1985

AMINOSYN 8.5%

ICU MEDICAL INC

8.5% (8.5GM/100ML)

N017673 004

AMINOSYN 8.5% (PH6)

ICU MEDICAL INC

8.5% (8.5GM/100ML)

N017673 007 Nov 18, 1985

AMINOSYN II 10%

ICU MEDICAL INC

10% (10GM/100ML)

N019438 005 Apr 03, 1986

AMINOSYN II 10% IN PLASTIC CONTAINER

ICU MEDICAL INC

10% (10GM/100ML)

N020015 001 Dec 19, 1991

AMINOSYN II 15% IN PLASTIC CONTAINER

ICU MEDICAL INC

15% (15GM/100ML)

N020041 001 Dec 19, 1991

AMINOSYN II 7%

ICU MEDICAL INC

7% (7GM/100ML)

N019438 003 Apr 03, 1986

AMINOSYN II 8.5%

ICU MEDICAL INC

8.5% (8.5GM/100ML)

N019438 004 Apr 03, 1986

AMINOSYN-HBC 7%

ICU MEDICAL INC

7% (7GM/100ML)

N019374 001 Jul 12, 1985

AMINOSYN-HF 8%

ICU MEDICAL INC

8% (8GM/100ML)

A020345 001 Apr 04, 1996

AMINOSYN-PF 10%

ICU MEDICAL INC

10% (10GM/100ML)

N019492 002 Oct 17, 1986

AMINOSYN-PF 7%

ICU MEDICAL INC

7% (7GM/100ML)

N019398 001 Sep 06, 1985

AMINOSYN-RF 5.2%

ICU MEDICAL INC

5.2% (5.2GM/100ML)

N018429 001

CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER

BAXTER HLTHCARE

15% (15GM/100ML)

A020512 001 Aug 30, 1996

FREAMINE HBC 6.9%

B BRAUN

6.9% (6.9GM/100ML)

N016822 006 May 17, 1983

FREAMINE III 10%

B BRAUN

10% (10GM/100ML)

N016822 005

FREAMINE III 8.5%

B BRAUN

8.5% (8.5GM/100ML)

N016822 004

HEPATAMINE 8%

B BRAUN

8% (8GM/100ML)

N018676 001 Aug 03, 1982

NEPHRAMINE 5.4%

B BRAUN

5.4% (5.4GM/100ML)

N017766 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-19 (of 436)

AMINO ACIDS

INJECTABLE; INJECTION

PREMASOL 10% IN PLASTIC CONTAINER		A075880 002 Jun 19, 2003
BAXTER HLTHCARE 10% (10GM/100ML)		
PREMASOL 6% IN PLASTIC CONTAINER		A075880 001 Jun 19, 2003
BAXTER HLTHCARE 6% (6GM/100ML)		
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER		N020849 001 Aug 26, 1998
+! BAXTER HLTHCARE 20% (20GM/100ML)		
TRAVASOL 10% IN PLASTIC CONTAINER		N018931 003 Aug 23, 1984
BAXTER HLTHCARE 10% (10MG/100ML)		
TRAVASOL 5.5% IN PLASTIC CONTAINER		N018931 001 Aug 23, 1984
BAXTER HLTHCARE 5.5% (5.5GM/100ML)		
TRAVASOL 8.5% IN PLASTIC CONTAINER		N018931 002 Aug 23, 1984
BAXTER HLTHCARE 8.5% (8.5GM/100ML)		
TROPHAMINE		
+! B BRAUN 6% (6GM/100ML)		N019018 001 Jul 20, 1984
TROPHAMINE 10%		
+! B BRAUN 10% (10GM/100ML)		N019018 003 Sep 07, 1988

AMINO ACIDS; CALCIUM ACETATE; GLYCERIN; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PROCALAMINE		
B BRAUN	3%;26MG/100ML;3GM/100ML;54MG/100ML;41MG/100ML;150MG/100ML;200MG/100ML;120MG/100ML	N018582 001 May 08, 1982

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 2.75%;33MG/100ML;10GM/100ML;51MG/100ML; 261MG/100ML;217MG/100ML;112MG/100ML		N020678 002 Mar 26, 1997
CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 2.75%;33MG/100ML;25GM/100ML;51MG/100ML; 261MG/100ML;217MG/100ML;112MG/100ML		N020678 005 Mar 26, 1997
CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 2.75%;33MG/100ML;5GM/100ML;51MG/100ML; 261MG/100ML;217MG/100ML;112MG/100ML		N020678 001 Mar 26, 1997
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 4.25%;33MG/100ML;10GM/100ML;51MG/100ML; 261MG/100ML;297MG/100ML;77MG/100ML		N020678 009 Mar 26, 1997
CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 4.25%;33MG/100ML;20GM/100ML;51MG/100ML; 261MG/100ML;297MG/100ML;77MG/100ML		N020678 011 Mar 26, 1997
CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 4.25%;33MG/100ML;25GM/100ML;51MG/100ML; 261MG/100ML;297MG/100ML;77MG/100ML		N020678 012 Mar 26, 1997
CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 4.25%;33MG/100ML;5GM/100ML;51MG/100ML; 261MG/100ML;297MG/100ML;77MG/100ML		N020678 008 Mar 26, 1997
CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 5%;33MG/100ML;10GM/100ML;51MG/100ML;261MG/100ML;340MG/100ML;59MG/100ML		N020678 016 Mar 26, 1997
CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 5%;33MG/100ML;15GM/100ML;51MG/100ML;261MG/100ML;340MG/100ML;59MG/100ML		N020678 017 Mar 26, 1997
CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 5%;33MG/100ML;20GM/100ML;51MG/100ML;261MG/100ML;340MG/100ML;59MG/100ML		N020678 018 Mar 26, 1997
CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 5%;33MG/100ML;25GM/100ML;51MG/100ML;261MG/100ML;340MG/100ML;59MG/100ML		N020678 019 Mar 26, 1997
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER		
+! BAXTER HLTHCARE 5%;33MG/100ML;35GM/100ML;51MG/100ML;261MG/100ML;340MG/100ML;59MG/100ML		N020678 021 Mar 26, 1997

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM GLYCEROPHOSPHATE; SOYBEAN OIL

EMULSION; IV (INFUSION)

KABIVEN IN PLASTIC CONTAINER		
+ FRESENIUS KABI USA 3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML; 174MG/100ML;239MG/100ML;147MG/100ML;3.9GM/100ML (1026ML)		N200656 004 Aug 25, 2014
+ 3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML; 174MG/100ML;239MG/100ML;147MG/100ML;3.9GM/100ML (1540ML)		N200656 005 Aug 25, 2014
+ 3.3%;29MG/100ML;9.8GM/100ML;96MG/100ML; 174MG/100ML;239MG/100ML;147MG/100ML;3.9GM/100ML (1540ML)		N200656 006 Aug 25, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-20 (of 436)

**AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM GLYCEROPHOSPHATE; SOYBEAN OIL**

EMULSION; IV (INFUSION)

KABIVEN IN PLASTIC CONTAINER

174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9  
 GM/100ML (2053ML)

+! 3.3%; 29MG/100ML; 9.8GM/100ML; 96MG/100ML;  
 174MG/100ML; 239MG/100ML; 147MG/100ML; 3.9  
 GM/100ML (2566ML)

PERIKABIVEN IN PLASTIC CONTAINER

+ FRESENIUS KABI USA 2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML;  
 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5  
 GM/100ML (1440ML)

+ 2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML;  
 124MG/100ML; 170MG/100ML; 105MG/100ML; 3.5  
 GM/100ML (1920ML)

+ 2.4%; 20MG/100ML; 6.8GM/100ML; 68MG/100ML;  
 124MG/100ML; 170MG/100ML;  
 ; 105MG/100ML; 3.5GM/100ML (2400ML)

**AMINO ACIDS; DEXTROSE**

INJECTABLE; INJECTION

CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 2.75%; 10GM/100ML

N020734 002 Sep 29, 1997

CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 2.75%; 25GM/100ML

N020734 005 Sep 29, 1997

CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%; 5GM/100ML

N020734 001 Sep 29, 1997

CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 4.25%; 10GM/100ML

N020734 008 Sep 29, 1997

CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 4.25%; 20GM/100ML

N020734 010 Sep 29, 1997

CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 4.25%; 25GM/100ML

N020734 011 Sep 29, 1997

CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%; 5GM/100ML

N020734 007 Sep 29, 1997

CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5%; 10GM/100ML

N020734 014 Sep 29, 1997

CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5%; 15GM/100ML

N020734 015 Sep 29, 1997

CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5%; 20GM/100ML

N020734 016 Sep 29, 1997

CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5%; 25GM/100ML

N020734 017 Sep 29, 1997

CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5%; 35GM/100ML

N020734 018 Sep 29, 1997

**AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE**

INJECTABLE; INJECTION

FREAMINE III 8.5% W/ ELECTROLYTES  
 B BRAUN 8.5%; 110MG/100ML; 230MG/100ML; 10MG/100ML  
 ; 440MG/100ML; 690MG/100ML

N016822 007 Jul 01, 1988

**AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE**

INJECTABLE; INJECTION

AMINOSYN 3.5% M  
 ICU MEDICAL INC 3.5%; 21MG/100ML; 40MG/100ML; 128MG/100ML;  
 234MG/100ML

N017789 003

**AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE**

INJECTABLE; INJECTION

FREAMINE III 3% W/ ELECTROLYTES  
 B BRAUN 3%; 54MG/100ML; 40MG/100ML; 150MG/100ML; 20  
 0MG/100ML; 120MG/100ML

N016822 003

**AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE**

INJECTABLE; INJECTION

AMINOSYN II 10% W/ ELECTROLYTES  
 ICU MEDICAL INC 10%; 102MG/100ML; 45MG/100ML; 522MG/100ML;  
 410MG/100ML

N019437 004 Apr 03, 1986

AMINOSYN II 8.5% W/ ELECTROLYTES  
 ICU MEDICAL INC 8.5%; 102MG/100ML; 45MG/100ML; 522MG/100ML;  
 410MG/100ML

N019437 005 Apr 03, 1986

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-21 (of 436)

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 7% W/ ELECTROLYTES

ICU MEDICAL INC 7%;102MG/100ML;522MG/100ML;410MG/100ML N017789 002

AMINOSYN 8.5% W/ ELECTROLYTES

ICU MEDICAL INC 8.5%;102MG/100ML;522MG/100ML;410MG/100ML N017673 005  
L

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMINOCAPROIC ACID

AP LUITPOLD 250MG/ML A071192 001 Dec 01, 1987

AMINOCAPROIC ACID IN PLASTIC CONTAINER

AP ! HOSPIRA 250MG/ML A070010 001 Mar 09, 1987

SYRUP; ORAL

AMICAR

AA +! CLOVER PHARMS 1.25GM/5ML N015230 002

AMINOCAPROIC ACID

AA AKORN 1.25GM/5ML A074759 001 Sep 02, 1998

TABLET; ORAL

AMICAR

AB + CLOVER PHARMS 500MG N015197 001

AMINOCAPROIC

AB AKORN 500MG A075602 001 May 24, 2001

AMICAR

+! CLOVER PHARMS 1GM N015197 002 Jun 24, 2004

AMINOLEVULINIC ACID HYDROCHLORIDE

FOR SOLUTION; ORAL

GLEOLAN  
+! NXDC 1.5GM/VIAL N208630 001 Jun 06, 2017

GEL; TOPICAL

AMELUZ  
+! BIOFRONTERA 10% N208081 001 May 10, 2016

SOLUTION; TOPICAL

LEVULAN  
+! DUSA 20% N020965 001 Dec 03, 1999

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

AP ! HOSPIRA 25MG/ML A087242 001 Oct 26, 1983

AP LUITPOLD 25MG/ML A087600 001

AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER  
! JACOBUS 4GM/PACKET A074346 001 Jun 30, 1994

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

AP ! AKORN 50MG/ML A076232 001 Jul 05, 2006

AP AUROBINDO PHARMA LTD 50MG/ML A204550 001 Oct 25, 2017

AP ! FRESENIUS KABI USA 50MG/ML A075761 001 Oct 15, 2002

AP ! GLAND PHARMA LTD 50MG/ML A077161 001 Apr 20, 2005

AP HIKMA FARMACEUTICA 50MG/ML A077234 001 Feb 25, 2008

AP ! HOSPIRA 50MG/ML A075955 001 Oct 18, 2002

AP HOSPIRA INC 50MG/ML A203884 001 Nov 25, 2013

AP 50MG/ML A203885 001 Nov 25, 2013

AP ! MYLAN INSTITUTIONAL 50MG/ML A076217 001 Oct 15, 2002

AP WOCKHARDT 50MG/ML A077610 001 Oct 30, 2008

AP 50MG/ML A077834 001 Oct 30, 2008

NEXTERONE

AP + BAXTER HLTHCARE 50MG/ML N022325 001 Dec 24, 2008

+! 150MG/100ML (1.5MG/ML) N022325 002 Nov 16, 2010

+! 360MG/200ML (1.8MG/ML) N022325 003 Nov 16, 2010

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

AB APOTEX INC 200MG A078578 001 Nov 06, 2008

AB AUROBINDO PHARMA LTD 200MG A204742 001 Jun 03, 2016

AB MAYNE PHARMA INC 100MG A075389 002 Dec 28, 2017

AB 200MG A075389 001 Jan 25, 2001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-22 (of 436)

AMIODARONE HYDROCHLORIDE

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

<u>AB</u>	MURTY PHARMS	<u>400MG</u>	<u>A075389 003</u>	Dec 28, 2017
<u>AB</u>		<u>100MG</u>	<u>A077069 003</u>	Oct 04, 2016
<u>AB</u>		<u>200MG</u>	<u>A077069 001</u>	Apr 08, 2005
<u>AB</u>		<u>400MG</u>	<u>A077069 002</u>	Apr 08, 2005
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A075315 001</u>	Dec 23, 1998
<u>AB</u>		<u>400MG</u>	<u>A075315 002</u>	Jun 30, 2000
<u>AB</u>	TARO PHARM	<u>100MG</u>	<u>A075424 002</u>	Dec 18, 2002
<u>AB</u>		<u>200MG</u>	<u>A075424 001</u>	Mar 30, 2001
<u>AB</u>		<u>400MG</u>	<u>A076362 001</u>	Nov 29, 2002
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>A074739 001</u>	Nov 30, 1998
<u>AB</u>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A079029 001</u>	Sep 16, 2008

PACERONE

<u>AB</u>	UPSHER-SMITH LABS	<u>100MG</u>	<u>A075135 002</u>	Apr 12, 2005
<u>AB</u>		<u>200MG</u>	<u>A075135 001</u>	Apr 30, 1998

AMIODARONE HYDROCHLORIDE

TARO PHARM 300MG

A076362 002 Dec 02, 2003

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>10MG</u>	<u>A202446 001</u>	Jun 04, 2014
<u>AB</u>		<u>25MG</u>	<u>A202446 002</u>	Jun 04, 2014
<u>AB</u>		<u>50MG</u>	<u>A202446 003</u>	Jun 04, 2014
<u>AB</u>		<u>75MG</u>	<u>A202446 004</u>	Jun 04, 2014
<u>AB</u>		<u>100MG</u>	<u>A202446 005</u>	Jun 04, 2014
<u>AB</u>		<u>150MG</u>	<u>A202446 006</u>	Jun 04, 2014
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A086009 002</u>	
<u>AB</u>		<u>25MG</u>	<u>A086009 003</u>	
<u>AB</u>		<u>50MG</u>	<u>A086009 001</u>	
<u>AB</u>		<u>75MG</u>	<u>A086009 004</u>	
<u>AB</u>		<u>100MG</u>	<u>A086009 005</u>	
<u>AB</u>		<u>150MG</u>	<u>A086009 006</u>	
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A085969 001</u>	
<u>AB</u>		<u>25MG</u>	<u>A085966 001</u>	
<u>AB</u>		<u>50MG</u>	<u>A085968 001</u>	
<u>AB</u>		<u>75MG</u>	<u>A085971 001</u>	
<u>AB</u>		<u>100MG</u>	<u>A085967 001</u>	
<u>AB</u>		<u>150MG</u>	<u>A085970 001</u>	
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A040218 001</u>	Sep 11, 1997
<u>AB</u>		<u>25MG</u>	<u>A040218 002</u>	Sep 11, 1997
<u>AB</u>		<u>50MG</u>	<u>A040218 003</u>	Sep 11, 1997
<u>AB</u>		<u>75MG</u>	<u>A040218 004</u>	Sep 11, 1997
<u>AB</u>		<u>100MG</u>	<u>A040218 005</u>	Sep 11, 1997
<u>AB</u>		<u>150MG</u>	<u>A040218 006</u>	Sep 11, 1997
<u>AB</u>	ZYDUS PHARMS USA INC	<u>10MG</u>	<u>A210086 001</u>	Oct 06, 2017
<u>AB</u>		<u>25MG</u>	<u>A210086 002</u>	Oct 06, 2017
<u>AB</u>		<u>50MG</u>	<u>A210086 003</u>	Oct 06, 2017
<u>AB</u>		<u>75MG</u>	<u>A210086 004</u>	Oct 06, 2017
<u>AB</u>		<u>100MG</u>	<u>A210086 005</u>	Oct 06, 2017
<u>AB</u>		<u>150MG</u>	<u>A210086 006</u>	Oct 06, 2017

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

MYLAN PHARMS INC	EQ 12.5MG BASE;5MG	A071297 002	Dec 10, 1986
!	EQ 25MG BASE;10MG	A071297 001	Dec 10, 1986

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

MYLAN	10MG;2MG	A071443 002	Nov 10, 1988
	10MG;4MG	A071443 003	Nov 10, 1988
!	25MG;2MG	A071443 004	Nov 10, 1988
!	25MG;4MG	A071443 005	Nov 10, 1988
!	50MG;4MG	A071443 001	Nov 10, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-23 (of 436)

AMLODIPINE BESYLATE

TABLET;ORAL

**AMLODIPINE BESYLATE**

<b>AB</b>	ACCORD HLTHCARE	<b>EQ 2.5MG BASE</b>	<b>A202553 001</b>	Apr 29, 2013
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A202553 002</b>	Apr 29, 2013
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A202553 003</b>	Apr 29, 2013
<b>AB</b>	ALKEM	<b>EQ 2.5MG BASE</b>	<b>A078925 001</b>	May 04, 2009
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078925 002</b>	May 04, 2009
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078925 003</b>	May 04, 2009
<b>AB</b>	AMNEAL PHARMS NY	<b>EQ 2.5MG BASE</b>	<b>A078477 001</b>	Jan 16, 2008
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078477 002</b>	Jan 16, 2008
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078477 003</b>	Jan 16, 2008
<b>AB</b>	APOTEX	<b>EQ 2.5MG BASE</b>	<b>A076719 001</b>	May 23, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A076719 002</b>	May 23, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A076719 003</b>	May 23, 2007
<b>AB</b>	AUROBINDO PHARMA	<b>EQ 2.5MG BASE</b>	<b>A078021 001</b>	Jul 17, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078021 002</b>	Jul 17, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078021 003</b>	Jul 17, 2007
<b>AB</b>	CHINA RESOURCES	<b>EQ 2.5MG BASE</b>	<b>A090752 003</b>	May 16, 2016
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A090752 001</b>	Apr 15, 2011
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A090752 002</b>	Apr 15, 2011
<b>AB</b>	CIPLA LTD	<b>EQ 2.5MG BASE</b>	<b>A077073 001</b>	Sep 26, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A077073 002</b>	Sep 26, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A076692 001</b>	Jul 20, 2007
<b>AB</b>	DR REDDYS LABS LTD	<b>EQ 2.5MG BASE</b>	<b>A076692 002</b>	Jul 20, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A076692 003</b>	Jul 20, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078552 001</b>	Apr 08, 2009
<b>AB</b>	EPIC PHARMA LLC	<b>EQ 2.5MG BASE</b>	<b>A078552 002</b>	Apr 08, 2009
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078552 003</b>	Apr 08, 2009
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A077771 001</b>	Apr 12, 2011
<b>AB</b>	HIKMA PHARMS	<b>EQ 2.5MG BASE</b>	<b>A077771 002</b>	Apr 12, 2011
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A077771 003</b>	Apr 12, 2011
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A077955 001</b>	Aug 28, 2007
<b>AB</b>	INVAGEN PHARMS	<b>EQ 2.5MG BASE</b>	<b>A206367 001</b>	Dec 10, 2015
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A077955 002</b>	Aug 28, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A206367 002</b>	Dec 10, 2015
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A077955 003</b>	Aug 28, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A206367 003</b>	Dec 10, 2015
<b>AB</b>	LUPIN	<b>EQ 2.5MG BASE</b>	<b>A078043 001</b>	Jul 12, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078043 002</b>	Jul 12, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078043 003</b>	Jul 12, 2007
<b>AB</b>	MACLEODS PHARMS LTD	<b>EQ 5MG BASE</b>	<b>A201380 001</b>	Apr 13, 2012
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A201380 002</b>	Apr 13, 2012
<b>AB</b>	MYLAN PHARMS INC	<b>EQ 2.5MG BASE</b>	<b>A076418 001</b>	Oct 03, 2005
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A076418 002</b>	Oct 03, 2005
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A076418 003</b>	Oct 03, 2005
<b>AB</b>	ORCHID HLTHCARE	<b>EQ 2.5MG BASE</b>	<b>A078453 001</b>	Jul 02, 2009
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078453 002</b>	Jul 02, 2009
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078453 003</b>	Jul 02, 2009
<b>AB</b>	POLYGEN PHARMS	<b>EQ 2.5MG BASE</b>	<b>A207821 001</b>	Jul 11, 2016
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A207821 002</b>	Jul 11, 2016
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A207821 003</b>	Jul 11, 2016
<b>AB</b>	SUN PHARM INDs INC	<b>EQ 2.5MG BASE</b>	<b>A078231 001</b>	Nov 30, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078231 002</b>	Nov 30, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078231 003</b>	Nov 30, 2007
<b>AB</b>	SUN PHARM INDs LTD	<b>EQ 2.5MG BASE</b>	<b>A077974 001</b>	Jul 09, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A077974 002</b>	Jul 09, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A077974 003</b>	Jul 09, 2007
<b>AB</b>	TEVA	<b>EQ 2.5MG BASE</b>	<b>A076846 001</b>	Jun 28, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A076846 002</b>	Jun 28, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A076846 003</b>	Jun 28, 2007
<b>AB</b>	TORRENT PHARMS	<b>EQ 2.5MG BASE</b>	<b>A078573 001</b>	Sep 22, 2008
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A078573 002</b>	Sep 22, 2008
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A078573 003</b>	Sep 22, 2008
<b>AB</b>	UNICHEM LABS LTD	<b>EQ 2.5MG BASE</b>	<b>A203245 001</b>	Oct 21, 2013
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A203245 002</b>	Oct 21, 2013
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A203245 003</b>	Oct 21, 2013
<b>AB</b>	UPSHER-SMITH LABS	<b>EQ 2.5MG BASE</b>	<b>A077759 001</b>	Jul 09, 2007
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A077759 002</b>	Jul 09, 2007
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A077759 003</b>	Jul 09, 2007
<b>AB</b>	VINTAGE	<b>EQ 2.5MG BASE</b>	<b>A078414 001</b>	Apr 07, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-24 (of 436)

AMLODIPINE BESYLATE

TABLET;ORAL

**AMLODIPINE BESYLATE**

<b>AB</b>		<u>EQ 5MG BASE</u>	<b>A078414 002</b>	Apr 07, 2010	
<b>AB</b>		<u>EQ 10MG BASE</u>	<b>A078414 003</b>	Apr 07, 2010	
<b>AB</b>	VIVIMED GLOBAL	<u>EQ 2.5MG BASE</u>	<b>A077516 001</b>	Jul 11, 2007	
<b>AB</b>		<u>EQ 5MG BASE</u>	<b>A077516 002</b>	Jul 11, 2007	
<b>AB</b>		<u>EQ 10MG BASE</u>	<b>A077516 003</b>	Jul 11, 2007	
<b>AB</b>	WATSON LABS	<u>EQ 2.5MG BASE</u>	<b>A077671 001</b>	Jul 19, 2007	
<b>AB</b>		<u>EQ 5MG BASE</u>	<b>A077671 002</b>	Jul 19, 2007	
<b>AB</b>		<u>EQ 10MG BASE</u>	<b>A077671 003</b>	Jul 19, 2007	
<b>AB</b>	WEST-WARD PHARMS INT	<u>EQ 2.5MG BASE</u>	<b>A077262 001</b>	Jul 09, 2007	
<b>AB</b>		<u>EQ 5MG BASE</u>	<b>A077262 002</b>	Jul 09, 2007	
<b>AB</b>		<u>EQ 10MG BASE</u>	<b>A077262 003</b>	Jul 09, 2007	
<b>AB</b>	WOCKHARDT	<u>EQ 2.5MG BASE</u>	<b>A078500 001</b>	Sep 06, 2007	
<b>AB</b>		<u>EQ 5MG BASE</u>	<b>A078500 002</b>	Sep 06, 2007	
<b>AB</b>		<u>EQ 10MG BASE</u>	<b>A078500 003</b>	Sep 06, 2007	
<b>AB</b>	ZYDUS PHARMS USA	<u>EQ 2.5MG BASE</u>	<b>A078226 001</b>	Jul 09, 2007	
<b>AB</b>		<u>EQ 5MG BASE</u>	<b>A078226 002</b>	Jul 09, 2007	
<b>AB</b>		<u>EQ 10MG BASE</u>	<b>A078226 003</b>	Jul 09, 2007	
<b>NORVASC</b>					
<b>AB</b>	+	Pfizer	<u>EQ 2.5MG BASE</u>	<b>N019787 001</b>	Jul 31, 1992
<b>AB</b>	+		<u>EQ 5MG BASE</u>	<b>N019787 002</b>	Jul 31, 1992
<b>AB</b>	+!		<u>EQ 10MG BASE</u>	<b>N019787 003</b>	Jul 31, 1992

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET;ORAL

**AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM**

<b>AB</b>	DR REDDYS LABS LTD	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<b>A203874 001</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<b>A203874 002</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<b>A203874 003</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<b>A203874 004</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<b>A203874 005</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<b>A203874 006</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<b>A203874 007</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<b>A203874 008</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<b>A203874 009</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<b>A203874 010</b>	Mar 07, 2014
<b>AB</b>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<b>A203874 011</b>	Mar 07, 2014
<b>AB</b>	MYLAN PHARMS INC	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<b>A200465 001</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<b>A200465 002</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<b>A200465 003</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<b>A200465 004</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<b>A200465 005</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<b>A200465 006</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<b>A200465 007</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<b>A200465 008</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<b>A200465 009</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<b>A200465 010</b>	Nov 29, 2013
<b>AB</b>		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<b>A200465 011</b>	Nov 29, 2013

**CADUET**

<b>AB</b>	+	Pfizer	<u>EQ 2.5MG BASE;EQ 10MG BASE</u>	<b>N021540 009</b>	Jul 29, 2004
<b>AB</b>	+		<u>EQ 2.5MG BASE;EQ 20MG BASE</u>	<b>N021540 010</b>	Jul 29, 2004
<b>AB</b>	+		<u>EQ 2.5MG BASE;EQ 40MG BASE</u>	<b>N021540 011</b>	Jul 29, 2004
<b>AB</b>	+		<u>EQ 5MG BASE;EQ 10MG BASE</u>	<b>N021540 001</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 5MG BASE;EQ 20MG BASE</u>	<b>N021540 002</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 5MG BASE;EQ 40MG BASE</u>	<b>N021540 003</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 5MG BASE;EQ 80MG BASE</u>	<b>N021540 004</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 10MG BASE;EQ 10MG BASE</u>	<b>N021540 005</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 10MG BASE;EQ 20MG BASE</u>	<b>N021540 006</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 10MG BASE;EQ 40MG BASE</u>	<b>N021540 007</b>	Jan 30, 2004
<b>AB</b>	+		<u>EQ 10MG BASE;EQ 80MG BASE</u>	<b>N021540 008</b>	Jan 30, 2004

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE;ORAL

**AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE**

<b>AB</b>	APOTEX INC	<u>EQ 2.5MG BASE;10MG</u>	<b>A091431 001</b>	Dec 30, 2013
<b>AB</b>		<u>EQ 5MG BASE;10MG</u>	<b>A091431 002</b>	Dec 30, 2013
<b>AB</b>		<u>EQ 5MG BASE;20MG</u>	<b>A091431 003</b>	Dec 30, 2013
<b>AB</b>		<u>EQ 5MG BASE;40MG</u>	<b>A091431 004</b>	Dec 30, 2013
<b>AB</b>		<u>EQ 10MG BASE;20MG</u>	<b>A091431 005</b>	Dec 30, 2013
<b>AB</b>		<u>EQ 10MG BASE;40MG</u>	<b>A091431 006</b>	Dec 30, 2013
<b>AB</b>	AUROBINDO PHARMA	<u>EQ 2.5MG BASE;10MG</u>	<b>A202239 001</b>	Sep 05, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-25 (of 436)

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

LTD

<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A202239 002</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A202239 003</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A202239 004</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A202239 005</u>	Sep 05, 2012
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202239 006</u>	Sep 05, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 2.5MG BASE;10MG</u>	<u>A077183 001</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A077183 002</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A077183 003</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A090149 001</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A077183 004</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A090149 002</u>	Jul 05, 2011
<u>AB</u>	LUPIN PHARMS	<u>EQ 2.5MG BASE;10MG</u>	<u>A078466 001</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A078466 002</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A078466 003</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A078466 005</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A078466 004</u>	Feb 05, 2010
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A078466 006</u>	Jul 05, 2011
<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE;10MG</u>	<u>A077375 001</u>	May 21, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A077375 002</u>	May 21, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A077375 003</u>	May 21, 2010
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A079047 001</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A077375 004</u>	May 21, 2010
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A079047 002</u>	Jul 05, 2011
<u>AB</u>	PAR PHARM	<u>EQ 2.5MG BASE;10MG</u>	<u>A078381 001</u>	Jul 29, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A078381 002</u>	Jul 29, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A078381 003</u>	Jul 29, 2010
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A078381 005</u>	Jul 29, 2010
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A078381 004</u>	Jul 29, 2010
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A078381 006</u>	Jul 29, 2010
<u>AB</u>	TEVA PHARMS	<u>EQ 2.5MG BASE;10MG</u>	<u>A077179 001</u>	May 18, 2007
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A077179 002</u>	May 18, 2007
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A077179 003</u>	May 18, 2007
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A077179 005</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A077179 004</u>	May 18, 2007
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A077179 006</u>	Jul 05, 2011
<u>AB</u>	WATSON LABS	<u>EQ 2.5MG BASE;10MG</u>	<u>A077890 001</u>	Oct 14, 2010
<u>AB</u>		<u>EQ 5MG BASE;10MG</u>	<u>A077890 002</u>	Oct 14, 2010
<u>AB</u>		<u>EQ 5MG BASE;20MG</u>	<u>A077890 003</u>	Oct 14, 2010
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A077890 004</u>	Oct 14, 2010
<u>AB</u>	WATSON LABS INC	<u>EQ 5MG BASE;40MG</u>	<u>A090364 001</u>	Jul 05, 2011
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A090364 002</u>	Jul 05, 2011

LOTREL

<u>AB</u>	+	NOVARTIS	<u>EQ 2.5MG BASE;10MG</u>	<u>N020364 002</u>	Mar 03, 1995
<u>AB</u>	+		<u>EQ 5MG BASE;10MG</u>	<u>N020364 003</u>	Mar 03, 1995
<u>AB</u>	+		<u>EQ 5MG BASE;20MG</u>	<u>N020364 004</u>	Mar 03, 1995
<u>AB</u>	+		<u>EQ 5MG BASE;40MG</u>	<u>N020364 007</u>	Apr 11, 2006
<u>AB</u>	+		<u>EQ 10MG BASE;20MG</u>	<u>N020364 005</u>	Jun 20, 2002
<u>AB</u>	!+		<u>EQ 10MG BASE;40MG</u>	<u>N020364 006</u>	Apr 11, 2006

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	PAR PHARM INC	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A206137 001</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A206137 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A206137 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A206137 004</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A206137 005</u>	Oct 26, 2016
<u>AB</u>	TEVA PHARMS USA	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A202491 001</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A202491 002</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A202491 003</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A202491 004</u>	Nov 03, 2016
<u>AB</u>		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A202491 005</u>	Nov 03, 2016
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>A203580 001</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>A203580 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;25MG;40MG</u>	<u>A203580 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>A203580 004</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;25MG;40MG</u>	<u>A203580 005</u>	Oct 26, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-26 (of 436)

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

**TRIBENZOR**

<u>AB</u>	+	DAIICHI SANKYO	<u>EQ 5MG BASE;12.5MG;20MG</u>	<u>N200175 001</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 5MG BASE;12.5MG;40MG</u>	<u>N200175 002</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 5MG BASE;25MG;40MG</u>	<u>N200175 003</u>	Jul 23, 2010
<u>AB</u>	+		<u>EQ 10MG BASE;12.5MG;40MG</u>	<u>N200175 004</u>	Jul 23, 2010
<u>AB</u>	+!		<u>EQ 10MG BASE;25MG;40MG</u>	<u>N200175 005</u>	Jul 23, 2010

AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET;ORAL

**AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE**

<u>AB</u>		AUROBINDO PHARMA LTD	<u>5MG;12.5MG;160MG</u>	<u>A206180 001</u>	Dec 19, 2017
<u>AB</u>			<u>5MG;25MG;160MG</u>	<u>A206180 002</u>	Dec 19, 2017
<u>AB</u>			<u>10MG;12.5MG;160MG</u>	<u>A206180 003</u>	Dec 19, 2017
<u>AB</u>			<u>10MG;25MG;160MG</u>	<u>A206180 004</u>	Dec 19, 2017
<u>AB</u>			<u>10MG;25MG;320MG</u>	<u>A206180 005</u>	Dec 19, 2017
<u>AB</u>		LUPIN LTD	<u>5MG;12.5MG;160MG</u>	<u>A200797 001</u>	Jun 03, 2015
<u>AB</u>			<u>5MG;25MG;160MG</u>	<u>A200797 002</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;12.5MG;160MG</u>	<u>A200797 003</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;25MG;160MG</u>	<u>A200797 004</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;25MG;320MG</u>	<u>A200797 005</u>	Jun 03, 2015
<u>AB</u>		PAR PHARM	<u>5MG;12.5MG;160MG</u>	<u>A201087 001</u>	Jun 01, 2015
<u>AB</u>			<u>5MG;25MG;160MG</u>	<u>A201087 002</u>	Jun 01, 2015
<u>AB</u>			<u>10MG;12.5MG;160MG</u>	<u>A201087 003</u>	Jun 01, 2015
<u>AB</u>			<u>10MG;25MG;160MG</u>	<u>A201087 004</u>	Jun 01, 2015
<u>AB</u>			<u>10MG;25MG;320MG</u>	<u>A201087 005</u>	Jun 01, 2015
<u>AB</u>		TEVA PHARMS	<u>5MG;12.5MG;160MG</u>	<u>A200435 001</u>	Sep 25, 2012
<u>AB</u>			<u>5MG;25MG;160MG</u>	<u>A200435 002</u>	Sep 25, 2012
<u>AB</u>			<u>10MG;12.5MG;160MG</u>	<u>A200435 003</u>	Sep 25, 2012
<u>AB</u>			<u>10MG;25MG;160MG</u>	<u>A200435 004</u>	Sep 25, 2012
<u>AB</u>			<u>10MG;25MG;320MG</u>	<u>A200435 005</u>	Sep 25, 2012
<u>AB</u>		TORRENT PHARMS LTD	<u>5MG;12.5MG;160MG</u>	<u>A200435 006</u>	Sep 25, 2012
<u>AB</u>			<u>5MG;25MG;160MG</u>	<u>A201593 001</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;12.5MG;160MG</u>	<u>A201593 002</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;25MG;160MG</u>	<u>A201593 003</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;25MG;320MG</u>	<u>A201593 004</u>	Jun 03, 2015
<u>AB</u>			<u>10MG;25MG;320MG</u>	<u>A201593 005</u>	Jun 03, 2015

**EXFORGE HCT**

<u>AB</u>	+	NOVARTIS	<u>5MG;12.5MG;160MG</u>	<u>N022314 001</u>	Apr 30, 2009
<u>AB</u>	+		<u>5MG;25MG;160MG</u>	<u>N022314 002</u>	Apr 30, 2009
<u>AB</u>	+		<u>10MG;12.5MG;160MG</u>	<u>N022314 003</u>	Apr 30, 2009
<u>AB</u>	+		<u>10MG;25MG;160MG</u>	<u>N022314 004</u>	Apr 30, 2009
<u>AB</u>	+!		<u>10MG;25MG;320MG</u>	<u>N022314 005</u>	Apr 30, 2009

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

**AMLODIPINE AND OLMESARTAN MEDOXOMIL**

<u>AB</u>		AJANTA PHARMA LTD	<u>EQ 5MG BASE;20MG</u>	<u>A207216 001</u>	Oct 28, 2016
<u>AB</u>			<u>EQ 5MG BASE;40MG</u>	<u>A207216 002</u>	Oct 28, 2016
<u>AB</u>			<u>EQ 10MG BASE;20MG</u>	<u>A207216 003</u>	Oct 28, 2016
<u>AB</u>			<u>EQ 10MG BASE;40MG</u>	<u>A207216 004</u>	Oct 28, 2016
<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A207073 001</u>	Jul 17, 2017
<u>AB</u>			<u>EQ 5MG BASE;40MG</u>	<u>A207073 002</u>	Jul 17, 2017
<u>AB</u>			<u>EQ 10MG BASE;20MG</u>	<u>A207073 003</u>	Jul 17, 2017
<u>AB</u>			<u>EQ 10MG BASE;40MG</u>	<u>A207073 004</u>	Jul 17, 2017
<u>AB</u>		ALKEM LABS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A209042 001</u>	Aug 14, 2017
<u>AB</u>			<u>EQ 5MG BASE;40MG</u>	<u>A209042 002</u>	Aug 14, 2017
<u>AB</u>			<u>EQ 10MG BASE;20MG</u>	<u>A209042 003</u>	Aug 14, 2017
<u>AB</u>			<u>EQ 10MG BASE;40MG</u>	<u>A209042 004</u>	Aug 14, 2017
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE;20MG</u>	<u>A206906 001</u>	May 15, 2017
<u>AB</u>			<u>EQ 5MG BASE;40MG</u>	<u>A206906 002</u>	May 15, 2017
<u>AB</u>			<u>EQ 10MG BASE;20MG</u>	<u>A206906 003</u>	May 15, 2017
<u>AB</u>			<u>EQ 10MG BASE;40MG</u>	<u>A206906 004</u>	May 15, 2017
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A207807 001</u>	Jul 05, 2017
<u>AB</u>			<u>EQ 5MG BASE;40MG</u>	<u>A207807 002</u>	Jul 05, 2017
<u>AB</u>			<u>EQ 10MG BASE;20MG</u>	<u>A207807 003</u>	Jul 05, 2017
<u>AB</u>			<u>EQ 10MG BASE;40MG</u>	<u>A207807 004</u>	Jul 05, 2017
<u>AB</u>		JUBILANT GENERICS	<u>EQ 5MG BASE;20MG</u>	<u>A207450 001</u>	May 15, 2017
<u>AB</u>			<u>EQ 5MG BASE;40MG</u>	<u>A207450 002</u>	May 15, 2017
<u>AB</u>			<u>EQ 10MG BASE;20MG</u>	<u>A207450 003</u>	May 15, 2017
<u>AB</u>			<u>EQ 10MG BASE;40MG</u>	<u>A207450 004</u>	May 15, 2017
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A206884 001</u>	Oct 26, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-27 (of 436)

AMLODIPINE BESYLATE; OLMESARTAN MEDOXOMIL

TABLET;ORAL

**AMLODIPINE AND OLMESARTAN MEDOXOMIL**

<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A206884 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A206884 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A206884 004</u>	Oct 26, 2016
<u>AB</u>	MICRO LABS	<u>EQ 5MG BASE;20MG</u>	<u>A207435 001</u>	Nov 02, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207435 002</u>	Nov 02, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207435 003</u>	Nov 02, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207435 004</u>	Nov 02, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 5MG BASE;20MG</u>	<u>A091154 001</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A091154 002</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A091154 003</u>	Oct 26, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A091154 004</u>	Oct 26, 2016
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE;20MG</u>	<u>A202933 001</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A202933 002</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A202933 003</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202933 004</u>	Nov 25, 2016
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 5MG BASE;20MG</u>	<u>A207771 001</u>	Sep 22, 2017
<u>AB</u>		<u>EQ 5MG BASE;40MG</u>	<u>A207771 002</u>	Sep 22, 2017
<u>AB</u>		<u>EQ 10MG BASE;20MG</u>	<u>A207771 003</u>	Sep 22, 2017
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A207771 004</u>	Sep 22, 2017
<b>AZOR</b>				
<u>AB</u>	+ DAIICHI SANKYO	<u>EQ 5MG BASE;20MG</u>	<u>N022100 001</u>	Sep 26, 2007
<u>AB</u>	+	<u>EQ 5MG BASE;40MG</u>	<u>N022100 002</u>	Sep 26, 2007
<u>AB</u>	+	<u>EQ 10MG BASE;20MG</u>	<u>N022100 003</u>	Sep 26, 2007
<u>AB</u>	++!	<u>EQ 10MG BASE;40MG</u>	<u>N022100 004</u>	Sep 26, 2007

AMLODIPINE BESYLATE; PERINDOPRIL ARGININE

TABLET;ORAL

PRESTALIA

+	MARINA BIOTECH	<u>EQ 2.5MG BASE;3.5MG</u>
+		<u>EQ 5MG BASE;7MG</u>
++!		<u>EQ 10MG BASE;14MG</u>

N205003 001 Jan 21, 2015  
 N205003 002 Jan 21, 2015  
 N205003 003 Jan 21, 2015

AMLODIPINE BESYLATE; TELMISARTAN

TABLET;ORAL

**TELMISARTAN AND AMLODIPINE**

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 5MG BASE;40MG</u>	<u>A205234 001</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A205234 003</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A205234 002</u>	Nov 17, 2016
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A205234 004</u>	Nov 17, 2016
<u>AB</u>	LUPIN LTD	<u>EQ 5MG BASE;40MG</u>	<u>A201586 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A201586 003</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A201586 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A201586 004</u>	Jan 08, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE;40MG</u>	<u>A202516 001</u>	Aug 26, 2014
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A202516 003</u>	Aug 26, 2014
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202516 002</u>	Aug 26, 2014
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A202516 004</u>	Aug 26, 2014
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE;40MG</u>	<u>A202517 001</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 5MG BASE;80MG</u>	<u>A202517 003</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;40MG</u>	<u>A202517 002</u>	Jan 08, 2014
<u>AB</u>		<u>EQ 10MG BASE;80MG</u>	<u>A202517 004</u>	Jan 08, 2014
<b>TWYNSTA</b>				
<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>EQ 5MG BASE;40MG</u>	<u>N022401 001</u>	Oct 16, 2009
<u>AB</u>	+	<u>EQ 5MG BASE;80MG</u>	<u>N022401 003</u>	Oct 16, 2009
<u>AB</u>	+	<u>EQ 10MG BASE;40MG</u>	<u>N022401 002</u>	Oct 16, 2009
<u>AB</u>	++!	<u>EQ 10MG BASE;80MG</u>	<u>N022401 004</u>	Oct 16, 2009

AMLODIPINE BESYLATE; VALSARTAN

TABLET;ORAL

**AMLODIPINE BESYLATE AND VALSARTAN**

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 5MG BASE;160MG</u>	<u>A202713 001</u>	Apr 03, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A202713 003</u>	Apr 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A202713 002</u>	Apr 03, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A202713 004</u>	Apr 03, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE;160MG</u>	<u>A206512 001</u>	Apr 22, 2016
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A206512 002</u>	Apr 22, 2016
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A206512 003</u>	Apr 22, 2016
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A206512 004</u>	Apr 22, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-28 (of 436)

AMLODIPINE BESYLATE; VALSARTAN

TABLET;ORAL

**AMLODIPINE BESYLATE AND VALSARTAN**

<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE;160MG</u>	<u>A205137 001</u>	Sep 16, 2016
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A205137 003</u>	Sep 16, 2016
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A205137 002</u>	Sep 16, 2016
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A205137 004</u>	Sep 16, 2016
<u>AB</u>	LUPIN	<u>EQ 5MG BASE;160MG</u>	<u>A090245 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090245 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090245 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090245 004</u>	Mar 30, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE;160MG</u>	<u>A090483 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090483 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090483 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090483 004</u>	Mar 30, 2015
<u>AB</u>	NOVEL LABS INC	<u>EQ 5MG BASE;160MG</u>	<u>A202829 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A202829 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A202829 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A202829 004</u>	Mar 30, 2015
<u>AB</u>	PAR PHARM INC	<u>EQ 5MG BASE;160MG</u>	<u>A090011 001</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A090011 003</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A090011 002</u>	Mar 28, 2013
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A090011 004</u>	Mar 28, 2013
<u>AB</u>	TEVA PHARMS USA	<u>EQ 5MG BASE;160MG</u>	<u>A091235 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A091235 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A091235 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A091235 004</u>	Mar 30, 2015
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE;160MG</u>	<u>A202377 001</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 5MG BASE;320MG</u>	<u>A202377 002</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;160MG</u>	<u>A202377 003</u>	Mar 30, 2015
<u>AB</u>		<u>EQ 10MG BASE;320MG</u>	<u>A202377 004</u>	Mar 30, 2015
<b>EXFORGE</b>				
<u>AB</u>	+ NOVARTIS	<u>EQ 5MG BASE;160MG</u>	<u>N021990 002</u>	Jun 20, 2007
<u>AB</u>	+	<u>EQ 5MG BASE;320MG</u>	<u>N021990 004</u>	Jun 20, 2007
<u>AB</u>	+!	<u>EQ 10MG BASE;160MG</u>	<u>N021990 003</u>	Jun 20, 2007
<u>AB</u>	+!	<u>EQ 10MG BASE;320MG</u>	<u>N021990 005</u>	Jun 20, 2007

AMMONIA N-13

INJECTABLE;INTRAVENOUS

**AMMONIA N 13**

<u>AP</u>	3D IMAGING DRUG	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203779 001</u>	Oct 19, 2015
<u>AP</u>	BIOMEDCL RES FDN	<u>48.75mCi-487.5mCi/13ML (3.75-37.5mCi/ML)</u>	<u>A204352 001</u>	May 01, 2015
<u>AP</u>	BRIGHAM WOMENS HOSP	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203783 001</u>	Oct 30, 2014
<u>AP</u>	CARDINAL HEALTH 414	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203700 001</u>	Feb 25, 2013
<u>AP</u>	+! FEINSTEIN	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>N022119 001</u>	Aug 23, 2007
<u>AP</u>	GEN HOSP	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A207025 001</u>	Feb 03, 2016
<u>AP</u>	GLOBAL ISOTOPES LLC	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204465 001</u>	Oct 23, 2014
<u>AP</u>	IBA MOLECULAR N AM	<u>18.8mCi-188mCi/5ML (3.75-37.5mCi/ML)</u>	<u>A204667 001</u>	Apr 22, 2015
<u>AP</u>	JOHNS HOPKINS UNIV	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204514 001</u>	Aug 19, 2014
<u>AP</u>	KREITCHMAN PET CTR	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203938 001</u>	Dec 09, 2013
<u>AP</u>	MCPRF	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203321 001</u>	Feb 25, 2013
<u>AP</u>	MIDWEST MEDCL	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204457 001</u>	Nov 18, 2015
<u>AP</u>	MIPS CRF	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204535 001</u>	Nov 20, 2014
<u>AP</u>	PETNET	<u>30mCi-300mCi (3.75-37.5mCi/ML)</u>	<u>A204510 001</u>	Nov 02, 2015
<u>AP</u>	SPECTRON MRC LLC	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204455 001</u>	Apr 23, 2015
<u>AP</u>	UCLA BIOMEDICAL	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203812 001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204496 001</u>	Mar 28, 2014
<u>AP</u>	UNIV TX MD ANDERSON	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A203933 001</u>	Jun 27, 2014
<u>AP</u>	WA UNIV SCH MED	<u>30mCi-300mCi/8ML (3.75-37.5mCi/ML)</u>	<u>A204506 001</u>	Feb 07, 2014
	ESSENTIAL ISOTOPES	3.75-260mCi/ML	A205687 001	Dec 17, 2015
	HOUSTON CYCLOTRON	3.75-260mCi/ML	A203543 001	Dec 14, 2012
	NCM USA BRONX LLC	3.75-260mCi/mL	A204515 001	Feb 04, 2015
	PRECISION NUCLEAR	3.75-260mCi/ML	A204547 001	Aug 14, 2015
	SHERTECH LABS LLC	3.75-260mCi/ML	A204366 001	Sep 19, 2014
	WI MEDCL CYCLOTRON	3.75-260mCi/ML	A204356 001	Dec 18, 2014

AMMONIUM CHLORIDE

INJECTABLE;INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

! HOSPIRA 5MEQ/ML

A088366 001 Jun 13, 1984

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-29 (of 436)

AMMONIUM LACTATE

CREAM;TOPICAL

**AMMONIUM LACTATE**

<b>AB</b>	!	PERRIGO NEW YORK	<u>EQ 12% BASE</u>	<b>A075774 001</b>	May 01, 2002
<b>AB</b>		TARO	<u>EQ 12% BASE</u>	<b>A075883 001</b>	Apr 10, 2003
<b>AB</b>		WATSON LABS INC	<u>EQ 12% BASE</u>	<b>A076829 001</b>	Feb 07, 2006

LOTION;TOPICAL

**AMMONIUM LACTATE**

<b>AB</b>	!	PERRIGO NEW YORK	<u>EQ 12% BASE</u>	<b>A075570 001</b>	Jun 23, 2004
<b>AB</b>		TARO	<u>EQ 12% BASE</u>	<b>A076216 001</b>	May 28, 2004
<b>AB</b>		WATSON LABS INC	<u>EQ 12% BASE</u>	<b>A075575 001</b>	Jun 11, 2002

AMOXAPINE

TABLET;ORAL

AMOXAPINE

WATSON LABS	25MG	A072688 001	Aug 28, 1992
	50MG	A072689 001	Aug 28, 1992
	100MG	A072690 001	Aug 28, 1992
!	150MG	A072691 001	Aug 28, 1992

AMOXICILLIN

CAPSULE;ORAL

**AMOXICILLIN**

<b>AB</b>	AM ANTIBIOTICS	<u>250MG</u>	<b>A062058 001</b>
<b>AB</b>		<u>500MG</u>	<b>A062058 002</b>
<b>AB</b>	AUROBINDO	<u>250MG</u>	<b>A065271 001</b>
<b>AB</b>		<u>500MG</u>	<b>A065271 002</b>
<b>AB</b>	DAVA PHARMS INC	<u>250MG</u>	<b>A062884 001</b>
<b>AB</b>		<u>500MG</u>	<b>A062881 001</b>
<b>AB</b>	HIKMA PHARMS	<u>250MG</u>	<b>A065291 001</b>
<b>AB</b>		<u>500MG</u>	<b>A065291 002</b>
<b>AB</b>	SANDOZ	<u>250MG</u>	<b>A064076 001</b>
<b>AB</b>		<u>500MG</u>	<b>A064076 002</b>
<b>AB</b>	TEVA	<u>250MG</u>	<b>A061926 001</b>
<b>AB</b>	!	<u>500MG</u>	<b>A061926 003</b>

**AMOXIL**

<b>AB</b>	DR REDDYS LABS INC	<u>250MG</u>	<b>A062216 001</b>
<b>AB</b>		<u>500MG</u>	<b>A062216 004</b>

FOR SUSPENSION;ORAL

**AMOXICILLIN**

<b>AB</b>	AUROBINDO	<u>200MG/5ML</u>	<b>A065334 001</b>
<b>AB</b>		<u>400MG/5ML</u>	<b>A065334 002</b>
<b>AB</b>	AUROBINDO PHARMA LTD	<u>125MG/5ML</u>	<b>A204030 001</b>
<b>AB</b>		<u>250MG/5ML</u>	<b>A204030 002</b>
<b>AB</b>	DAVA PHARMS INC	<u>125MG/5ML</u>	<b>A062927 001</b>
<b>AB</b>		<u>250MG/5ML</u>	<b>A062927 002</b>
<b>AB</b>	HIKMA	<u>125MG/5ML</u>	<b>A065322 002</b>
<b>AB</b>		<u>200MG/5ML</u>	<b>A065325 002</b>
<b>AB</b>		<u>250MG/5ML</u>	<b>A065322 001</b>
<b>AB</b>		<u>400MG/5ML</u>	<b>A065325 001</b>
<b>AB</b>	SANDOZ	<u>125MG/5ML</u>	<b>A065387 001</b>
<b>AB</b>		<u>200MG/5ML</u>	<b>A065387 002</b>
<b>AB</b>		<u>250MG/5ML</u>	<b>A065387 002</b>
<b>AB</b>		<u>400MG/5ML</u>	<b>A065387 002</b>
<b>AB</b>	TEVA	<u>125MG/5ML</u>	<b>A061931 001</b>
<b>AB</b>		<u>200MG/5ML</u>	<b>A065119 001</b>
<b>AB</b>	!	<u>250MG/5ML</u>	<b>A061931 002</b>
<b>AB</b>	!	<u>400MG/5ML</u>	<b>A065119 002</b>
<b>AB</b>	WOCKHARDT BIO AG	<u>400MG/5ML</u>	<b>A065319 002</b>

**AMOXICILLIN PEDIATRIC**

<b>AB</b>	TEVA	<u>50MG/ML</u>	<b>A061931 003</b>
-----------	------	----------------	--------------------

**AMOXIL**

<b>AB</b>	DR REDDYS LABS INC	<u>50MG/ML</u>	<b>A062226 005</b>
<b>AB</b>		<u>125MG/5ML</u>	<b>A062226 001</b>
<b>AB</b>		<u>250MG/5ML</u>	<b>A062226 002</b>

**LAROTID**

<b>AB</b>	DR REDDYS LABS INC	<u>125MG/5ML</u>	<b>A062226 003</b>
<b>AB</b>		<u>250MG/5ML</u>	<b>A062226 004</b>

TABLET;ORAL

**AMOXICILLIN**

<b>AB</b>	AUROBINDO	<u>500MG</u>	<b>A065256 001</b>
<b>AB</b>		<u>875MG</u>	<b>A065256 002</b>
<b>AB</b>	HIKMA	<u>875MG</u>	<b>A065255 001</b>

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-30 (of 436)

AMOXICILLIN

TABLET;ORAL

AMOXICILLIN

<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065228 001</u>	Jul 13, 2005
<u>AB</u>		<u>875MG</u>	<u>A065228 002</u>	Jul 13, 2005
<u>AB</u>	TEVA	<u>500MG</u>	<u>A065056 001</u>	Sep 18, 2000
<u>AB</u>	!	<u>875MG</u>	<u>A065056 002</u>	Sep 18, 2000

TABLET, CHEWABLE;ORAL

AMOXICILLIN

TEVA	125MG	A064013 002	Sep 11, 1995
!	250MG	A064013 001	Dec 22, 1992

TABLET, EXTENDED RELEASE;ORAL

MOXATAG

+!	VERNALIS R AND D LTD	775MG	N050813 001	Jan 23, 2008
----	----------------------	-------	-------------	--------------

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN

<u>AB</u>	RISING PHARMS INC	<u>500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG</u>	<u>A206006 001</u>	Oct 07, 2016
<u>AB</u>	SANDOZ INC	<u>500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG</u>	<u>A202588 001</u>	Mar 04, 2014
<u>AB</u>	TEVA PHARMS USA	<u>500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG</u>	<u>A200218 001</u>	Aug 30, 2013
<b>PREVPAC</b>				
<u>AB</u>	+! TAKEDA PHARMS USA	<u>500MG,N/A,N/A;N/A,500MG,N/A;N/A,N/A,30MG</u>	<u>N050757 001</u>	Dec 02, 1997

AMOXICILLIN; CLARITHROMYCIN; OMEPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED RELEASE;ORAL

OMEПRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN

+!	GASTROENTERO	500MG, N/A, N/A; N/A, 500MG, N/A; N/A, N/A, 20MG	N050824 001	Feb 08, 2011
		G		

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	AUROBINDO PHARMA LTD	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A201090 001</u>	Dec 20, 2011
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A201090 002</u>	Dec 20, 2011
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A201091 001</u>	Dec 20, 2011
<u>AB</u>	HIKMA PHARMS	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065191 002</u>	Jan 25, 2005
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065191 001</u>	Jan 25, 2005
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065373 001</u>	Nov 09, 2007
<u>AB</u>	SANDOZ	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065066 001</u>	Jun 05, 2002
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065066 002</u>	Jun 05, 2002
<u>AB</u>	SANDOZ INC	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065098 001</u>	Dec 16, 2002
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065098 002</u>	Dec 16, 2002
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065358 001</u>	Aug 13, 2007
<u>AB</u>	TEVA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>A065089 001</u>	May 25, 2004
<u>AB</u>	!	<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>A065089 002</u>	May 25, 2004
<u>AB</u>	!	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065162 001</u>	Mar 12, 2004
<u>AB</u>	WOCKHARDT BIO AG	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>A065431 001</u>	Nov 25, 2008
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>A065420 001</u>	Dec 02, 2013
<b>AUGMENTIN '250'</b>				
<u>AB</u>	+! DR REDDYS LABS INC	<u>250MG/5ML;EQ 62.5MG BASE/5ML</u>	<u>N050575 002</u>	Aug 06, 1984
	AUGMENTIN '125'			
	+ DR REDDYS LABS INC	125MG/5ML;EQ 31.25MG BASE/5ML	N050575 001	Aug 06, 1984

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG;EQ 125MG BASE</u>	<u>A091569 001</u>	Jan 20, 2012
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A091569 002</u>	Jan 20, 2012
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>A091568 001</u>	Jan 20, 2012
<u>AB</u>	HIKMA PHARMS	<u>875MG;EQ 125MG BASE</u>	<u>A203824 001</u>	Aug 23, 2016
<u>AB</u>	MICRO LABS LTD INDIA	<u>250MG;EQ 125MG BASE</u>	<u>A205707 001</u>	Dec 30, 2016
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A205707 002</u>	Dec 30, 2016
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>A204755 003</u>	Dec 30, 2016
<u>AB</u>	!	<u>250MG;EQ 125MG BASE</u>	<u>A065189 001</u>	Aug 23, 2005
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A065064 001</u>	Mar 15, 2002
<u>AB</u>	!	<u>875MG;EQ 125MG BASE</u>	<u>A065063 001</u>	Mar 14, 2002
<u>AB</u>	!	<u>500MG;EQ 125MG BASE</u>	<u>A065117 001</u>	Nov 27, 2002
<u>AB</u>	SANDOZ INC	<u>875MG;EQ 125MG BASE</u>	<u>A065093 001</u>	Nov 21, 2002
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>A065101 001</u>	Oct 30, 2002
<u>AB</u>	TEVA	<u>500MG;EQ 125MG BASE</u>		

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-31 (of 436)

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	TEVA PHARMS USA	<u>875MG;EQ 125MG BASE</u>	<u>A065096 001</u>	Oct 29, 2002
		<u>AUGMENTIN '875'</u>		
<u>AB</u>	+ DR REDDYS LABS INC	<u>875MG;EQ 125MG BASE</u>	<u>N050720 001</u>	Feb 13, 1996
	TABLET, CHEWABLE;ORAL			
	AMOXICILLIN AND CLAVULANATE POTASSIUM			
	TEVA	200MG;EQ 28.5MG BASE	A065205 001	Feb 09, 2005
	!	400MG;EQ 57MG BASE	A065205 002	Feb 09, 2005
	TABLET, EXTENDED RELEASE;ORAL			
	<u>AMOXICILLIN AND CLAVULANATE POTASSIUM</u>			
<u>AB</u>	SANDOZ	<u>1GM;EQ 62.5MG BASE</u>	<u>A090227 001</u>	Apr 21, 2010
		<u>AUGMENTIN XR</u>		
<u>AB</u>	+! DR REDDYS LABS INC	<u>1GM;EQ 62.5MG BASE</u>	<u>N050785 001</u>	Sep 25, 2002

AMPHETAMINE

SUSPENSION, EXTENDED RELEASE;ORAL

ADZENYS ER

+! NEOS THERAPS INC	EQ 1.25MG BASE/ML	N204325 001	Sep 15, 2017
DYANAVEL XR		N208147 001	Oct 19, 2015

+! TRIS PHARMA INC EQ 2.5MG BASE/ML

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL  
ADZENYS XR-ODT

+ NEOS THERAPS	EQ 3.1MG BASE	N204326 001	Jan 27, 2016
+	EQ 6.3MG BASE	N204326 002	Jan 27, 2016
+	EQ 9.4MG BASE	N204326 003	Jan 27, 2016
+	EQ 12.5MG BASE	N204326 004	Jan 27, 2016
+	EQ 15.7MG BASE	N204326 005	Jan 27, 2016
+	EQ 18.8MG BASE	N204326 006	Jan 27, 2016

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL

ADDERALL XR 10

<u>AB</u> + SHIRE	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N021303 001</u>	Oct 11, 2001
-------------------	--------------------------------	--------------------	--------------

ADDERALL XR 15

<u>AB</u> + SHIRE	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N021303 006</u>	May 22, 2002
-------------------	------------------------------------	--------------------	--------------

ADDERALL XR 20

<u>AB</u> + SHIRE	<u>5MG;5MG;5MG;5MG</u>	<u>N021303 002</u>	Oct 11, 2001
-------------------	------------------------	--------------------	--------------

ADDERALL XR 25

<u>AB</u> + SHIRE	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>N021303 004</u>	May 22, 2002
-------------------	------------------------------------	--------------------	--------------

ADDERALL XR 30

<u>AB</u> +! SHIRE	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N021303 003</u>	Oct 11, 2001
--------------------	--------------------------------	--------------------	--------------

ADDERALL XR 5

<u>AB</u> + SHIRE	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N021303 005</u>	May 22, 2002
-------------------	------------------------------------	--------------------	--------------

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u> ACTAVIS ELIZABETH	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A077302 001</u>	Jun 22, 2012
-----------------------------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A077302 002</u>	Jun 22, 2012
-----------	--------------------------------	--------------------	--------------

<u>AB</u>	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A077302 003</u>	Jun 22, 2012
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>5MG;5MG;5MG;5MG</u>	<u>A077302 004</u>	Jun 22, 2012
-----------	------------------------	--------------------	--------------

<u>AB</u>	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>A077302 005</u>	Jun 22, 2012
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A077302 006</u>	Jun 22, 2012
-----------	--------------------------------	--------------------	--------------

<u>AB</u> IMPAX LABS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A076852 001</u>	Feb 16, 2016
----------------------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A076852 002</u>	Feb 16, 2016
-----------	--------------------------------	--------------------	--------------

<u>AB</u>	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A076852 003</u>	Feb 16, 2016
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>5MG;5MG;5MG;5MG</u>	<u>A076852 004</u>	Feb 16, 2016
-----------	------------------------	--------------------	--------------

<u>AB</u>	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>A076852 005</u>	Feb 16, 2016
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A076852 006</u>	Feb 16, 2016
-----------	--------------------------------	--------------------	--------------

<u>AB</u> TEVA	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A077488 001</u>	Apr 29, 2013
----------------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A077488 002</u>	Apr 29, 2013
-----------	--------------------------------	--------------------	--------------

<u>AB</u>	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A077488 003</u>	Apr 29, 2013
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>5MG;5MG;5MG;5MG</u>	<u>A077488 004</u>	Apr 29, 2013
-----------	------------------------	--------------------	--------------

<u>AB</u>	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>A077488 005</u>	Apr 29, 2013
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A077488 006</u>	Apr 29, 2013
-----------	--------------------------------	--------------------	--------------

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u> BARR LABS INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A076536 001</u>	Feb 12, 2013
-------------------------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A076536 002</u>	Feb 12, 2013
-----------	--------------------------------	--------------------	--------------

<u>AB</u>	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A076536 003</u>	Feb 12, 2013
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>5MG;5MG;5MG;5MG</u>	<u>A076536 004</u>	Feb 12, 2013
-----------	------------------------	--------------------	--------------

<u>AB</u>	<u>6.25MG;6.25MG;6.25MG;6.25MG</u>	<u>A076536 005</u>	Feb 12, 2013
-----------	------------------------------------	--------------------	--------------

<u>AB</u>	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A076536 006</u>	Feb 12, 2013
-----------	--------------------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-32 (of 436)

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL  
 MYDAYIS

+	SHIRE DEV LLC	3.125MG;3.125MG;3.125MG;3.125MG	N022063 001	Jun 20, 2017
+		6.25MG;6.25MG;6.25MG;6.25MG	N022063 002	Jun 20, 2017
+		9.375MG;9.375MG;9.375MG;9.375MG	N022063 003	Jun 20, 2017
+		12.5MG;12.5MG;12.5MG;12.5MG	N022063 004	Jun 20, 2017

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040456 001</u>	May 06, 2003
<u>AB</u>		<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A206340 001</u>	Feb 05, 2016
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A206340 002</u>	Feb 05, 2016
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040456 002</u>	May 06, 2003
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A206340 003</u>	Feb 05, 2016
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A206340 004</u>	Feb 05, 2016
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A206340 005</u>	Feb 05, 2016
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A040456 003</u>	May 06, 2003
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A206340 006</u>	Feb 05, 2016
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040456 004</u>	May 06, 2003
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A206340 007</u>	Feb 05, 2016
<u>AB</u>	ALVOGEN MALTA	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A207388 001</u>	Jul 28, 2017
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A207388 002</u>	Jul 28, 2017
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A207388 003</u>	Jul 28, 2017
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A207388 004</u>	Jul 28, 2017
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A207388 005</u>	Jul 28, 2017
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A207388 006</u>	Jul 28, 2017
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A207388 007</u>	Jul 28, 2017
<u>AB</u>	AUROLIFE PHARMA LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A202424 001</u>	Nov 27, 2013
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A202424 002</u>	Nov 27, 2013
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A202424 003</u>	Nov 27, 2013
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A202424 004</u>	Nov 27, 2013
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A202424 005</u>	Nov 27, 2013
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A202424 006</u>	Nov 27, 2013
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A202424 007</u>	Nov 27, 2013
<u>AB</u>	BARR	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040422 001</u>	Feb 11, 2002
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040422 005</u>	Mar 19, 2003
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040422 002</u>	Feb 11, 2002
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040422 006</u>	Mar 19, 2003
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040422 007</u>	Mar 19, 2003
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A040422 003</u>	Feb 11, 2002
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040422 004</u>	Feb 11, 2002
<u>AB</u>	EPIC PHARMA LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040444 001</u>	Jun 19, 2002
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040444 005</u>	Nov 03, 2014
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040444 002</u>	Jun 19, 2002
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040444 006</u>	Nov 03, 2014
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040444 007</u>	Nov 03, 2014
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A040444 003</u>	Jun 19, 2002
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040444 004</u>	Jun 19, 2002
<u>AB</u>	MYLAN PHARMS INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A206721 001</u>	Nov 10, 2015
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A206721 002</u>	Nov 10, 2015
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A206721 003</u>	Nov 10, 2015
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A206721 004</u>	Nov 10, 2015
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A206721 005</u>	Nov 10, 2015
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A206721 006</u>	Nov 10, 2015
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A206721 007</u>	Nov 10, 2015
<u>AB</u>	NESHER PHARMS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A207340 001</u>	Oct 31, 2017
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A207340 002</u>	Oct 31, 2017
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A207340 003</u>	Oct 31, 2017
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A207340 004</u>	Oct 31, 2017
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A207340 005</u>	Oct 31, 2017
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A207340 006</u>	Oct 31, 2017
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A207340 007</u>	Oct 31, 2017
<u>AB</u>	SANDOZ	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040439 004</u>	Sep 27, 2002
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040439 001</u>	Jun 14, 2002
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A040439 002</u>	Jun 14, 2002
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040439 003</u>	Jun 14, 2002
<u>AB</u>	SPECGX LLC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040440 001</u>	Oct 07, 2003
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040440 002</u>	Oct 07, 2003
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040440 003</u>	Oct 07, 2003
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040440 004</u>	Oct 07, 2003
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040440 005</u>	Oct 07, 2003
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A040440 006</u>	Oct 07, 2003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-33 (of 436)

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

**DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE**

<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040440</u>	<u>007</u>	Oct 07, 2003
<u>AB</u>	SUN PHARM INDUSTRIES	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A040480</u>	<u>001</u>	Sep 09, 2003
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A040480</u>	<u>002</u>	Sep 09, 2003
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A040480</u>	<u>003</u>	Sep 09, 2003
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A040480</u>	<u>004</u>	Sep 09, 2003
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A040480</u>	<u>005</u>	Sep 09, 2003
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A040480</u>	<u>006</u>	Sep 09, 2003
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A040480</u>	<u>007</u>	Sep 09, 2003
<u>AB</u>	SUNRISE PHARM INC	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>A209799</u>	<u>001</u>	Dec 28, 2017
<u>AB</u>		<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>A209799</u>	<u>002</u>	Dec 28, 2017
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>A209799</u>	<u>003</u>	Dec 28, 2017
<u>AB</u>		<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>A209799</u>	<u>004</u>	Dec 28, 2017
<u>AB</u>		<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>A209799</u>	<u>005</u>	Dec 28, 2017
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>A209799</u>	<u>006</u>	Dec 28, 2017
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>A209799</u>	<u>007</u>	Dec 28, 2017

AMPHETAMINE SULFATE

TABLET; ORAL

EVEKEO

ARBOR PHARMS LLC  
!

5MG

10MG

A200166 001 Aug 09, 2012  
A200166 002 Aug 09, 2012

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

! X GEN PHARMS 50MG/VIAL

A063206 001 Apr 29, 1992

INJECTABLE, LIPID COMPLEX; INJECTION

ABELCET

+! LEADIANT BIOSCI INC 5MG/ML

N050724 001 Nov 20, 1995

INJECTABLE, LIPOSOMAL; INJECTION

AMBISOME

+! ASTELLAS 50MG/VIAL

N050740 001 Aug 11, 1997

AMPICILLIN SODIUM

INJECTABLE; INJECTION

**AMPICILLIN SODIUM**

<u>AP</u>	ACS DOBFAR SPA	<u>EQ 500MG BASE/VIAL</u>	<u>A090884</u>	<u>001</u>	Apr 03, 2013
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090884</u>	<u>002</u>	Apr 03, 2013
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090884</u>	<u>003</u>	Apr 03, 2013
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090889</u>	<u>001</u>	Apr 03, 2013
<u>AP</u>	ANTIBIOTICE	<u>EQ 250MG BASE/VIAL</u>	<u>A090354</u>	<u>001</u>	Dec 28, 2009
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A090354</u>	<u>002</u>	Dec 28, 2009
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090354</u>	<u>003</u>	Dec 28, 2009
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090354</u>	<u>004</u>	Dec 28, 2009
<u>AP</u>	AUROBINDO PHARMA	<u>EQ 125MG BASE/VIAL</u>	<u>A065499</u>	<u>001</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A065499</u>	<u>002</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065499</u>	<u>003</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065499</u>	<u>004</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065499</u>	<u>005</u>	Aug 17, 2010
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065493</u>	<u>001</u>	Aug 17, 2010
<u>AP</u>	HANFORD GC	<u>EQ 250MG BASE/VIAL</u>	<u>A062772</u>	<u>006</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A062772</u>	<u>007</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062772</u>	<u>001</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062772</u>	<u>003</u>	Apr 15, 1993
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A063142</u>	<u>001</u>	Apr 15, 1993
<u>AP</u>	HOSPIRA INC	<u>EQ 250MG BASE/VIAL</u>	<u>A202864</u>	<u>001</u>	Sep 04, 2015
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A202864</u>	<u>002</u>	Sep 04, 2015
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A202864</u>	<u>003</u>	Sep 04, 2015
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A202864</u>	<u>004</u>	Sep 04, 2015
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A202865</u>	<u>001</u>	Sep 04, 2015
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 10GM BASE/VIAL</u>	<u>A201404</u>	<u>001</u>	Dec 20, 2013
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A062797</u>	<u>001</u>	Jul 12, 1993
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A062719</u>	<u>001</u>	May 12, 1987
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A062719</u>	<u>003</u>	May 12, 1987
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062719</u>	<u>002</u>	May 12, 1987
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A062797</u>	<u>002</u>	Jul 12, 1993
<u>AP</u>	MYLAN LABS LTD	<u>EQ 250MG BASE/VIAL</u>	<u>A201025</u>	<u>001</u>	Apr 09, 2014
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A201025</u>	<u>002</u>	Apr 09, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-34 (of 436)

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A201025 003</u>	Apr 09, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A201025 004</u>	Apr 09, 2014
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A202198 001</u>	Apr 07, 2014
<u>AP</u>	SAGENT PHARMS	<u>EQ 125MG BASE/VIAL</u>	<u>A090583 001</u>	Nov 27, 2015
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>A090583 002</u>	Nov 27, 2015
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A090583 003</u>	Nov 27, 2015
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090583 004</u>	Nov 27, 2015
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090583 005</u>	Nov 27, 2015
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090581 001</u>	Oct 20, 2015
<u>AP</u>	! SANDOZ	<u>EQ 125MG BASE/VIAL</u>	<u>A061395 001</u>	
<u>AP</u>	!	<u>EQ 250MG BASE/VIAL</u>	<u>A061395 002</u>	
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A061395 003</u>	
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A061395 004</u>	
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A061395 005</u>	
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A061395 006</u>	

POWDER; INTRAVENOUS

AMPICILLIN SODIUM

<u>AP</u>	! SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A062738 001</u>	Feb 19, 1987
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062738 002</u>	Feb 19, 1987

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<u>AP</u>	ACS DOBFAR	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065406 001</u>	Dec 22, 2009
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065406 002</u>	Dec 22, 2009
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065403 001</u>	Dec 23, 2009
<u>AP</u>	ANTIBIOTICE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201406 001</u>	Dec 07, 2015
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201406 002</u>	Dec 07, 2015
<u>AP</u>	AUROBINDO PHARMA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090340 001</u>	Sep 20, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090349 001</u>	Sep 20, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090340 002</u>	Sep 20, 2010
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090349 002</u>	Sep 20, 2010
<u>AP</u>	HANFORD GC	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090339 001</u>	Sep 20, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065176 001</u>	Nov 30, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065176 002</u>	Nov 30, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065188 001</u>	Nov 25, 2005
<u>AP</u>	HOSPIRA INC	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090375 001</u>	Dec 21, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090653 001</u>	Dec 21, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090375 002</u>	Dec 21, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090653 002</u>	Dec 21, 2011
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090646 001</u>	Dec 21, 2011
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065222 001</u>	Nov 29, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065222 002</u>	Nov 29, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065314 001</u>	Nov 27, 2006
<u>AP</u>	MUSTAFA NEVZAT ILAC	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065316 001</u>	Jun 29, 2007
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065316 002</u>	Jun 29, 2007
<u>AP</u>	MYLAN LABS LTD	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A201024 001</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A201024 002</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A202197 001</u>	Apr 07, 2014
<u>AP</u>	SAGENT PHARMS	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A090579 001</u>	Jan 08, 2016
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A090579 002</u>	Jan 08, 2016
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A090578 001</u>	Jan 11, 2016
<u>AP</u>	SANDOZ	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065241 001</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065310 001</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065241 002</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065310 002</u>	Jul 25, 2006
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065240 001</u>	Jul 25, 2006
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065074 001</u>	Mar 19, 2002
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A065074 002</u>	Mar 19, 2002
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>A065076 001</u>	Mar 19, 2002
<b>UNASYN</b>				
<u>AP</u>	! PFIZER	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>A062901 002</u>	Feb 27, 1992
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A062901 001</u>	Nov 23, 1988
<u>AP</u>	+!	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N050608 002</u>	Dec 31, 1986
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N050608 001</u>	Dec 31, 1986
<u>AP</u>	+!	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N050608 005</u>	Dec 10, 1993

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-35 (of 436)

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE;ORAL

AMPICILLIN TRIHYDRATE

<u>AB</u>	DAVA PHARMS INC	<u>EQ 250MG BASE</u>	<u>A062883 001</u>	Feb 25, 1988
<u>AB</u>	!	<u>EQ 500MG BASE</u>	<u>A062882 001</u>	Feb 25, 1988
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>A064082 001</u>	Aug 29, 1995
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A064082 002</u>	Aug 29, 1995
FOR SUSPENSION;ORAL				
AMPICILLIN TRIHYDRATE				
	DAVA PHARMS INC	EQ 125MG BASE/5ML	A062982 001	Feb 10, 1989
!		EQ 250MG BASE/5ML	A062982 002	Feb 10, 1989

ANAGRELIDE HYDROCHLORIDE

CAPSULE;ORAL

AGRYLIN

<u>AB</u>	SHIRE LLC	<u>EQ 0.5MG BASE</u>	<u>N020333 001</u>	Mar 14, 1997
<u>AB</u>	<u>ANAGRELIDE HYDROCHLORIDE</u>			
<u>AB</u>	BARR	<u>EQ 0.5MG BASE</u>	<u>A076530 001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076530 002</u>	Apr 18, 2005
<u>AB</u>	IMPAX LABS	<u>EQ 0.5MG BASE</u>	<u>A076910 001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A076910 002</u>	Apr 18, 2005
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 0.5MG BASE</u>	<u>A076468 001</u>	Apr 18, 2005
<u>AB</u>	!	<u>EQ 1MG BASE</u>	<u>A076468 002</u>	Apr 18, 2005
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 0.5MG BASE</u>	<u>A209151 001</u>	Jun 30, 2017
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A209151 002</u>	Jun 30, 2017

ANASTROZOLE

TABLET;ORAL

ANASTROZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A090568 001</u>	Jun 28, 2010
<u>AB</u>	APOTEX INC	<u>1MG</u>	<u>A200654 001</u>	May 11, 2012
<u>AB</u>	CIPLA LTD	<u>1MG</u>	<u>A091164 001</u>	Jun 28, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A090732 001</u>	Jun 28, 2010
<u>AB</u>	FRESENIUS KABI ONCOL	<u>1MG</u>	<u>A090088 001</u>	Jun 28, 2010
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A091051 001</u>	Jun 28, 2010
<u>AB</u>	NATCO PHARMA LTD	<u>1MG</u>	<u>A079220 001</u>	Jun 28, 2010
<u>AB</u>	SANTOS BIOTECH	<u>1MG</u>	<u>A078944 001</u>	Jun 28, 2010
<u>AB</u>	TEVA PHARMS	<u>1MG</u>	<u>A078058 001</u>	Jun 28, 2010
<u>AB</u>	WEST-WARD PHARMS INT	<u>1MG</u>	<u>A078485 001</u>	Jun 28, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>1MG</u>	<u>A078921 001</u>	Jun 28, 2010

ARIMIDEX

<u>AB</u>	+!	ASTRAZENECA PHARMS	<u>1MG</u>	<u>N020541 001</u>	Dec 27, 1995
-----------	----	--------------------	------------	--------------------	--------------

ANGIOTENSIN II

SOLUTION;IV (INFUSION)

GIAPREZA

+!	LA JOLLA PHARM CO	2.5MG/ML (2.5MG/ML) 5MG/2ML (2.5MG/ML)	N209360 001	Dec 21, 2017
+			N209360 002	Dec 21, 2017

ANIDULAFUNGIN

INJECTABLE;IV (INFUSION)

ERAXIS

+	VICURON	50MG/VIAL	N021632 001	Feb 17, 2006
+		100MG/VIAL	N021632 002	Nov 14, 2006

APIXABAN

TABLET;ORAL

ELIQUIS

+	BRISTOL MYERS SQUIBB	2.5MG	N202155 001	Dec 28, 2012
+		5MG	N202155 002	Dec 28, 2012

APOMORPHINE HYDROCHLORIDE

INJECTABLE;SUBCUTANEOUS

APOKYN

+	US WORLDMEDS	30MG/3ML (10MG/ML)	N021264 002	Apr 20, 2004
---	--------------	--------------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-36 (of 436)

APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

APRACLONIDINE HYDROCHLORIDE

<u>AT</u>	AKORN INC	<u>EQ 0.5% BASE</u>	<u>A077764 001</u> Mar 12, 2009
		<u>IOPIDINE</u>	
<u>AT</u>	+! NOVARTIS PHARMS CORP	<u>EQ 0.5% BASE</u>	<u>N020258 001</u> Jul 30, 1993

+!

EQ 1% BASE

N019779 001 Dec 31, 1987

APREMILAST

TABLET;ORAL

OTEZLA

+ CELGENE CORP	10MG	N205437 001 Mar 21, 2014
+	20MG	N205437 002 Mar 21, 2014
+!	30MG	N205437 003 Mar 21, 2014

APREPITANT

CAPSULE;ORAL

APREPITANT

<u>AB</u>	GLENMARK PHARMS LTD	<u>40MG</u>	<u>A207777 001</u> Oct 12, 2017
<u>AB</u>		<u>80MG</u>	<u>A207777 002</u> Oct 12, 2017
<u>AB</u>		<u>125MG</u>	<u>A207777 003</u> Oct 12, 2017
<u>AB</u>	SANDOZ	<u>40MG</u>	<u>A090999 001</u> Sep 24, 2012
<u>AB</u>		<u>80MG</u>	<u>A090999 002</u> Sep 24, 2012
<u>AB</u>		<u>125MG</u>	<u>A090999 003</u> Sep 24, 2012

EMEND

<u>AB</u>	+ MERCK	<u>40MG</u>	<u>N021549 003</u> Jun 30, 2006
<u>AB</u>	+	<u>80MG</u>	<u>N021549 001</u> Mar 26, 2003
<u>AB</u>	+!	<u>125MG</u>	<u>N021549 002</u> Mar 26, 2003

EMULSION;IV (INFUSION)

CINVANTI

+! HERON THERAPS INC 130MG/18ML (7.2MG/ML)

FOR SUSPENSION;ORAL

EMEND

+! MSD MERCK CO 125MG/KIT

N209296 001 Nov 09, 2017

N207865 001 Dec 17, 2015

ARFORMOTEROL TARTRATE

SOLUTION;INHALATION

BROVANA

+! SUNOVION EQ 0.015MG BASE/2ML

N021912 001 Oct 06, 2006

ARGATROBAN

INJECTABLE;INJECTION

ARGATROBAN

<u>AP</u>	FRESENIUS KABI USA	<u>250MG/2.5ML (100MG/ML)</u>	<u>N201811 001</u> Mar 23, 2015
<u>AP</u>	HIKMA PHARM CO LTD	<u>250MG/2.5ML (100MG/ML)</u>	<u>N203049 001</u> Jan 05, 2012
<u>AP</u>	HOSPIRA INC	<u>250MG/2.5ML (100MG/ML)</u>	<u>A204120 001</u> Sep 21, 2016
<u>AP</u>	MYLAN INSTITUTIONAL	<u>250MG/2.5ML (100MG/ML)</u>	<u>A202626 001</u> Jun 30, 2014
<u>AP</u>	+! NOVARTIS PHARMS CORP	<u>250MG/2.5ML (100MG/ML)</u>	<u>N020883 001</u> Jun 30, 2000

<u>AP</u>	PAR STERILE PRODUCTS	<u>250MG/2.5ML (100MG/ML)</u>	<u>A091665 001</u> Jun 30, 2014
	+! HIKMA PHARM CO LTD	50MG/50ML (1MG/ML)	N203049 002 Sep 30, 2016

INJECTABLE;IV (INFUSION)

ARGATROBAN IN SODIUM CHLORIDE

<u>AP</u>	GLAND PHARMA LTD	<u>125MG/125ML (1MG/ML)</u>	<u>A205570 001</u> May 22, 2017
<u>AP</u>	+! SANDOZ	<u>125MG/125ML (1MG/ML)</u>	<u>N022485 001</u> May 09, 2011
	ARGATROBAN IN 0.9% SODIUM CHLORIDE		
	TEVA PHARMS USA	250MG/250ML (1MG/ML)	N206769 001 Dec 15, 2014
	ARGATROBAN IN SODIUM CHLORIDE		
	+! EAGLE PHARMS	50MG/50ML (1MG/ML)	N022434 001 Jun 29, 2011

ARGININE HYDROCHLORIDE

INJECTABLE;INJECTION

R-GENE 10

+! PHARMACIA AND UPJOHN 10GM/100ML

N016931 001

ARIPIPRAZOLE

FOR SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ABILIFY MAINTENA KIT

+ OTSUKA PHARM CO LTD	300MG/VIAL	N202971 001 Feb 28, 2013
+	300MG	N202971 003 Sep 29, 2014
+!	400MG/VIAL	N202971 002 Feb 28, 2013
+	400MG	N202971 004 Sep 29, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-37 (of 436)

**ARIPIPRAZOLE**

SOLUTION;ORAL

**ARIPIPRAZOLE**

<b>AA</b>	!	AMNEAL PHARMS	<u>1MG/ML</u>	<b>A203906</b>	<b>001</b>	Aug 14, 2015
<b>AA</b>		APOTEX INC	<u>1MG/ML</u>	<b>A204094</b>	<b>001</b>	Sep 30, 2015
<b>AA</b>		SILARX PHARMS INC	<u>1MG/ML</u>	<b>A204171</b>	<b>001</b>	Aug 14, 2015

TABLET;ORAL

**ABILIFY**

<b>AB</b>	+	OTSUKA	<u>2MG</u>	<b>N021436</b>	<b>006</b>	Nov 15, 2002
<b>AB</b>	+		<u>5MG</u>	<b>N021436</b>	<b>005</b>	Nov 15, 2002
<b>AB</b>	+		<u>10MG</u>	<b>N021436</b>	<b>001</b>	Nov 15, 2002
<b>AB</b>	+		<u>15MG</u>	<b>N021436</b>	<b>002</b>	Nov 15, 2002
<b>AB</b>	+		<u>20MG</u>	<b>N021436</b>	<b>003</b>	Nov 15, 2002
<b>AB</b>	+		<u>30MG</u>	<b>N021436</b>	<b>004</b>	Nov 15, 2002

**ARIPIPRAZOLE**

<b>AB</b>		ACCORD HLTHCARE	<u>2MG</u>	<b>A206251</b>	<b>001</b>	Dec 07, 2016
<b>AB</b>			<u>5MG</u>	<b>A206251</b>	<b>002</b>	Dec 07, 2016
<b>AB</b>			<u>10MG</u>	<b>A206251</b>	<b>003</b>	Dec 07, 2016
<b>AB</b>			<u>15MG</u>	<b>A206251</b>	<b>004</b>	Dec 07, 2016
<b>AB</b>			<u>20MG</u>	<b>A206251</b>	<b>005</b>	Dec 07, 2016
<b>AB</b>			<u>30MG</u>	<b>A206251</b>	<b>006</b>	Dec 07, 2016
<b>AB</b>		AJANTA PHARMA LTD	<u>2MG</u>	<b>A206174</b>	<b>001</b>	Sep 12, 2016
<b>AB</b>			<u>5MG</u>	<b>A206174</b>	<b>002</b>	Sep 12, 2016
<b>AB</b>			<u>10MG</u>	<b>A206174</b>	<b>003</b>	Sep 12, 2016
<b>AB</b>			<u>15MG</u>	<b>A206174</b>	<b>004</b>	Sep 12, 2016
<b>AB</b>			<u>20MG</u>	<b>A206174</b>	<b>005</b>	Sep 12, 2016
<b>AB</b>			<u>30MG</u>	<b>A206174</b>	<b>006</b>	Sep 12, 2016
<b>AB</b>		ALEMBIC PHARMS LTD	<u>2MG</u>	<b>A202101</b>	<b>001</b>	Apr 28, 2015
<b>AB</b>			<u>5MG</u>	<b>A202101</b>	<b>002</b>	Apr 28, 2015
<b>AB</b>			<u>10MG</u>	<b>A202101</b>	<b>003</b>	Apr 28, 2015
<b>AB</b>			<u>15MG</u>	<b>A202101</b>	<b>004</b>	Apr 28, 2015
<b>AB</b>			<u>20MG</u>	<b>A202101</b>	<b>005</b>	Apr 28, 2015
<b>AB</b>			<u>30MG</u>	<b>A202101</b>	<b>006</b>	Apr 28, 2015
<b>AB</b>		AMNEAL PHARMS	<u>2MG</u>	<b>A204838</b>	<b>001</b>	Jun 17, 2016
<b>AB</b>			<u>5MG</u>	<b>A204838</b>	<b>002</b>	Jun 17, 2016
<b>AB</b>			<u>10MG</u>	<b>A204838</b>	<b>003</b>	Jun 17, 2016
<b>AB</b>			<u>15MG</u>	<b>A204838</b>	<b>004</b>	Jun 17, 2016
<b>AB</b>			<u>20MG</u>	<b>A204838</b>	<b>005</b>	Jun 17, 2016
<b>AB</b>			<u>30MG</u>	<b>A204838</b>	<b>006</b>	Jun 17, 2016
<b>AB</b>		APOTEX INC	<u>2MG</u>	<b>A078583</b>	<b>001</b>	Jul 24, 2015
<b>AB</b>			<u>5MG</u>	<b>A078583</b>	<b>002</b>	Jul 24, 2015
<b>AB</b>			<u>10MG</u>	<b>A078583</b>	<b>003</b>	Jul 24, 2015
<b>AB</b>			<u>15MG</u>	<b>A078583</b>	<b>004</b>	Jul 24, 2015
<b>AB</b>			<u>20MG</u>	<b>A078583</b>	<b>005</b>	Jul 24, 2015
<b>AB</b>			<u>30MG</u>	<b>A078583</b>	<b>006</b>	Jul 24, 2015
<b>AB</b>		AUROBINDO PHARMA LTD	<u>2MG</u>	<b>A203908</b>	<b>001</b>	Oct 08, 2015
<b>AB</b>			<u>5MG</u>	<b>A203908</b>	<b>002</b>	Oct 08, 2015
<b>AB</b>			<u>10MG</u>	<b>A203908</b>	<b>003</b>	Oct 08, 2015
<b>AB</b>			<u>15MG</u>	<b>A203908</b>	<b>004</b>	Oct 08, 2015
<b>AB</b>			<u>20MG</u>	<b>A203908</b>	<b>005</b>	Oct 08, 2015
<b>AB</b>			<u>30MG</u>	<b>A203908</b>	<b>006</b>	Oct 08, 2015
<b>AB</b>		HETERO LABS LTD V	<u>2MG</u>	<b>A205064</b>	<b>001</b>	Apr 28, 2015
<b>AB</b>			<u>5MG</u>	<b>A205064</b>	<b>002</b>	Apr 28, 2015
<b>AB</b>			<u>10MG</u>	<b>A205064</b>	<b>003</b>	Apr 28, 2015
<b>AB</b>			<u>15MG</u>	<b>A205064</b>	<b>004</b>	Apr 28, 2015
<b>AB</b>			<u>20MG</u>	<b>A205064</b>	<b>005</b>	Apr 28, 2015
<b>AB</b>			<u>30MG</u>	<b>A205064</b>	<b>006</b>	Apr 28, 2015
<b>AB</b>		MACLEODS PHARMS LTD	<u>2MG</u>	<b>A204111</b>	<b>001</b>	Oct 07, 2016
<b>AB</b>			<u>5MG</u>	<b>A204111</b>	<b>002</b>	Oct 07, 2016
<b>AB</b>			<u>10MG</u>	<b>A204111</b>	<b>003</b>	Oct 07, 2016
<b>AB</b>			<u>15MG</u>	<b>A204111</b>	<b>004</b>	Oct 07, 2016
<b>AB</b>			<u>20MG</u>	<b>A204111</b>	<b>005</b>	Oct 07, 2016
<b>AB</b>			<u>30MG</u>	<b>A204111</b>	<b>006</b>	Oct 07, 2016
<b>AB</b>		ORCHID HLTHCARE	<u>2MG</u>	<b>A202683</b>	<b>001</b>	May 23, 2017
<b>AB</b>			<u>5MG</u>	<b>A202683</b>	<b>002</b>	May 23, 2017
<b>AB</b>			<u>10MG</u>	<b>A202683</b>	<b>003</b>	May 23, 2017
<b>AB</b>			<u>15MG</u>	<b>A202683</b>	<b>004</b>	May 23, 2017
<b>AB</b>			<u>20MG</u>	<b>A202683</b>	<b>005</b>	May 23, 2017
<b>AB</b>			<u>30MG</u>	<b>A202683</b>	<b>006</b>	May 23, 2017
<b>AB</b>		PRINSTON INC	<u>2MG</u>	<b>A205363</b>	<b>001</b>	Dec 04, 2017
<b>AB</b>			<u>5MG</u>	<b>A205363</b>	<b>002</b>	Dec 04, 2017
<b>AB</b>			<u>10MG</u>	<b>A205363</b>	<b>003</b>	Dec 04, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-38 (of 436)

ARIPIPRAZOLE

TABLET;ORAL

ARIPIPRAZOLE

<u>AB</u>		<u>15MG</u>	<u>A205363</u>	<u>004</u>	Dec 04, 2017
<u>AB</u>		<u>20MG</u>	<u>A205363</u>	<u>005</u>	Dec 04, 2017
<u>AB</u>		<u>30MG</u>	<u>A205363</u>	<u>006</u>	Dec 04, 2017
<u>AB</u>	SANTOS BIOTECH	<u>2MG</u>	<u>A091279</u>	<u>001</u>	Jan 09, 2017
<u>AB</u>		<u>5MG</u>	<u>A091279</u>	<u>002</u>	Jan 09, 2017
<u>AB</u>		<u>10MG</u>	<u>A091279</u>	<u>003</u>	Jan 09, 2017
<u>AB</u>		<u>15MG</u>	<u>A091279</u>	<u>004</u>	Jan 09, 2017
<u>AB</u>		<u>20MG</u>	<u>A091279</u>	<u>005</u>	Jan 09, 2017
<u>AB</u>		<u>30MG</u>	<u>A091279</u>	<u>006</u>	Jan 09, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>2MG</u>	<u>A206383</u>	<u>001</u>	Sep 29, 2016
<u>AB</u>		<u>5MG</u>	<u>A206383</u>	<u>002</u>	Sep 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A206383</u>	<u>003</u>	Sep 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A206383</u>	<u>004</u>	Sep 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A206383</u>	<u>005</u>	Sep 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A206383</u>	<u>006</u>	Sep 29, 2016
<u>AB</u>	TEVA PHARMS USA	<u>2MG</u>	<u>A078607</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A078607</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A078608</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A078708</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A078708</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A078708</u>	<u>003</u>	Apr 28, 2015
<u>AB</u>	TORRENT PHARMS LTD	<u>2MG</u>	<u>A201519</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>10MG</u>	<u>A201519</u>	<u>003</u>	Apr 28, 2015
<u>AB</u>		<u>5MG</u>	<u>A201519</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A201519</u>	<u>004</u>	Apr 28, 2015
<u>AB</u>		<u>20MG</u>	<u>A201519</u>	<u>005</u>	Apr 28, 2015
<u>AB</u>		<u>30MG</u>	<u>A201519</u>	<u>006</u>	Apr 28, 2015
ABILITY MYCITE KIT					
	+ OTSUKA PHARM CO LTD	<u>2MG</u>	N207202	<u>001</u>	Nov 13, 2017
	+!		N207202	<u>002</u>	Nov 13, 2017
	+		N207202	<u>003</u>	Nov 13, 2017
	+		N207202	<u>004</u>	Nov 13, 2017
	+		N207202	<u>005</u>	Nov 13, 2017
	+		N207202	<u>006</u>	Nov 13, 2017

TABLET, ORALLY DISINTEGRATING;ORAL

ARIPIPRAZOLE

<u>AB</u>	! ALEMBIC PHARMS LTD	<u>10MG</u>	<u>A202102</u>	<u>001</u>	Apr 28, 2015
<u>AB</u>		<u>15MG</u>	<u>A202102</u>	<u>002</u>	Apr 28, 2015
<u>AB</u>	ORCHID HLTHCARE	<u>10MG</u>	<u>A202547</u>	<u>001</u>	Dec 11, 2017
<u>AB</u>		<u>15MG</u>	<u>A202547</u>	<u>002</u>	Dec 11, 2017

ARIPIPRAZOLE LAUROXIL

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ARISTADA

+	ALKERMES INC	441MG/1.6ML (275.63MG/ML)	N207533	<u>001</u>	Oct 05, 2015
+		662MG/2.4ML (275.83MG/ML)	N207533	<u>002</u>	Oct 05, 2015
+!		882MG/3.2ML (275.63MG/ML)	N207533	<u>003</u>	Oct 05, 2015
+		1064MG/3.9ML (272.82MG/ML)	N207533	<u>004</u>	Jun 05, 2017

ARMODAFINIL

TABLET;ORAL

ARMODAFINIL

<u>AB</u>	LUPIN LTD	<u>50MG</u>	<u>A200751</u>	<u>001</u>	Nov 28, 2016
<u>AB</u>		<u>150MG</u>	<u>A200751</u>	<u>003</u>	Nov 28, 2016
<u>AB</u>		<u>200MG</u>	<u>A200751</u>	<u>004</u>	Nov 28, 2016
<u>AB</u>		<u>250MG</u>	<u>A200751</u>	<u>005</u>	Nov 28, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A200043</u>	<u>001</u>	Jun 01, 2012
<u>AB</u>		<u>150MG</u>	<u>A200043</u>	<u>002</u>	Jun 01, 2012
<u>AB</u>		<u>250MG</u>	<u>A200043</u>	<u>003</u>	Jun 01, 2012
<u>AB</u>	NATCO PHARMA LTD	<u>50MG</u>	<u>A202768</u>	<u>001</u>	Nov 28, 2016
<u>AB</u>		<u>150MG</u>	<u>A202768</u>	<u>002</u>	Nov 28, 2016
<u>AB</u>		<u>200MG</u>	<u>A202768</u>	<u>005</u>	Sep 28, 2017
<u>AB</u>		<u>250MG</u>	<u>A202768</u>	<u>003</u>	Nov 28, 2016

NUVIGIL

<u>AB</u>	+	CEPHALON	<u>50MG</u>	<u>N021875</u>	<u>001</u>	Jun 15, 2007
<u>AB</u>	+		<u>150MG</u>	<u>N021875</u>	<u>003</u>	Jun 15, 2007
<u>AB</u>	+		<u>200MG</u>	<u>N021875</u>	<u>005</u>	Mar 26, 2009
<u>AB</u>	+		<u>250MG</u>	<u>N021875</u>	<u>004</u>	Jun 15, 2007

ARMODAFINIL

NATCO PHARMA LTD	100MG	A202768	004	Sep 28, 2017
------------------	-------	---------	-----	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-39 (of 436)

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

TRISENOX

+! CEPHALON  
+!

1MG/ML  
2MG/ML

N021248 001 Sep 25, 2000  
N021248 002 Oct 13, 2017

ARTEMETHER; LUMEFANTRINE

TABLET; ORAL

COARTEM

+! NOVARTIS

20MG;120MG

N022268 001 Apr 07, 2009

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE

AP HOSPIRA

4%:EQ 0.017MG BASE/1.7ML (4%:EQ 0.01MG  
BASE/ML)

A079138 001 Jun 18, 2010

SEPTOCAINE

AP +! DEPROCO

4%:EQ 0.0085MG BASE/1.7ML (4%:EQ  
0.005MG BASE/ML)

N022010 001 Mar 30, 2006

AP +!

4%:EQ 0.017MG BASE/1.7ML (4%:EQ 0.01MG  
BASE/ML)

N020971 001 Apr 03, 2000

ULTACAN

AP HANSAMED INC

4%:EQ 0.0085MG BASE/1.7ML (4%:EQ  
0.005MG BASE/ML)

A201751 001 Jul 11, 2017

ULTACAN FORTE

AP HANSAMED INC

4%:EQ 0.017MG BASE/1.7ML (4%:EQ 0.01MG  
BASE/ML)

A201750 001 Jul 11, 2017

ORABLOC

+ PIERREL

4%;EQ 0.009MG BASE/1.8ML (EQ 0.005MG  
BASE/ML)

N022466 001 Feb 26, 2010

+!

4%;EQ 0.018MG BASE/1.8ML (EQ 0.01MG  
BASE/ML)

N022466 002 Feb 26, 2010

ASCORBIC ACID

SOLUTION; IV (INFUSION)

ASCOR

+! MCGUFF

25,000MG/50ML (500MG/ML)

N209112 001 Oct 02, 2017

ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE;  
PYRIDOXINE; RIBOFLAVIN; THIAMINE; TOCOPHEROL ACETATE; VITAMIN A; VITAMIN K

INJECTABLE; IV (INFUSION)

INFUVITE PEDIATRIC

+! SANDOZ INC

80MG/VIAL;0.02MG/VIAL;400  
IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VI  
AL;17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/  
VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL

N021265 001 Feb 21, 2001

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+! SANDOZ INC

80MG/VIAL;0.02MG/VIAL;400  
IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VI  
AL;17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/  
VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL

N021646 001 Jan 29, 2004

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE;  
PHYTONADIONE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE;  
VITAMIN A; VITAMIN E

FOR SOLUTION; IV (INFUSION)

M.V.I. PEDIATRIC

+! HOSPIRA

80MG/VIAL;0.02MG/VIAL;0.001MG/VIAL;5MG/  
VIAL;0.01MG/VIAL;0.14MG/VIAL;17MG/VIAL;  
0.2MG/VIAL;1MG/VIAL;1.4MG/VIAL;EQ 1.2MG  
BASE/VIAL;0.7MG/VIAL;7MG/VIAL

N018920 001 Sep 21, 2000

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE;  
PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A;  
VITAMIN E; VITAMIN K

INJECTABLE; IV (INFUSION)

M.V.I. ADULT

+! HOSPIRA

200MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15M  
G/VIAL;0.005MG/VIAL;0.6MG/VIAL;40MG/VIA  
L;6MG/VIAL;3.6MG/VIAL;6MG/VIAL;1MG/VIAL  
;10MG/VIAL;0.15MG/VIAL

N021625 001 Jan 30, 2004

M.V.I. ADULT (PHARMACY BULK PACKAGE)

+! HOSPIRA

200MG/5ML;0.06MG/5ML;0.005MG/5ML;15MG/5  
ML;0.005MG/5ML;0.6MG/5ML;40MG/5ML;6MG/5  
ML;3.6MG/5ML;6MG/5ML;1MG/5ML;10MG/5ML;0  
.15MG/5ML

N021643 001 Feb 18, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-40 (of 436)

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

MOVIPREP

+! SALIX PHARMS 4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM N021881 001 Aug 02, 2006

ASENAPINE MALEATE

TABLET; SUBLINGUAL

SAPHRIS

+! FOREST LABS LLC	EQ 2.5 BASE	N022117 003 Mar 12, 2015
+!	EQ 5MG BASE	N022117 001 Aug 13, 2009
+!	EQ 10MG BASE	N022117 002 Aug 13, 2009

ASPIRIN

CAPSULE, EXTENDED RELEASE; ORAL

DURLAZA

+! NEW HAVEN PHARMS 162.5MG N200671 001 Sep 04, 2015

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINAL

AA +! ALLERGAN SALES LLC 325MG;50MG;40MG

N017534 005 Apr 16, 1986

LANORTINAL

AA LANNETT 325MG;50MG;40MG

A086996 002 Oct 11, 1985

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

AA ! HIKMA INTL PHARMS 325MG;50MG;40MG

A086162 002 Feb 16, 1984

AA PII 325MG;50MG;40MG

A204195 001 Sep 22, 2016

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE AND CODEINE PHOSPHATE

AB MAYNE PHARMA INC 325MG;50MG;40MG;30MG

A203335 001 Oct 30, 2015

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

AB NEXGEN PHARMA INC 325MG;50MG;40MG;30MG

A075231 001 Nov 30, 2001

AB STEVENS J 325MG;50MG;40MG;30MG

A074951 001 Aug 31, 1998

FIORINAL W/CODEINE

AB +! ALLERGAN SALES LLC 325MG;50MG;40MG;30MG

N019429 003 Oct 26, 1990

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

SYNALGOS-DC

+! SUN PHARM 356.4MG;30MG;16MG N011483 004 Sep 06, 1983  
 INDUSTRIES

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

SANDOZ 385MG;30MG;25MG

A074654 001 Dec 31, 1996

! 770MG;60MG;50MG

A074654 002 Dec 31, 1996

ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

AB ! HERITAGE PHARMS INC 325MG;200MG

A089594 001 Mar 31, 1989

AB NOVAST LABS LTD 325MG;200MG

A040832 001 Jan 07, 2010

AB OXFORD PHARMS 325MG;200MG

A040252 001 Dec 10, 1997

AB SANDOZ 325MG;200MG

A040116 001 Apr 25, 1996

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

AB INGENUS PHARMS NJ 325MG;200MG;16MG

A040860 001 Jan 07, 2010

AB ! SANDOZ 325MG;200MG;16MG

A040118 001 Apr 16, 1996

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOX

AB +! BOEHRINGER 25MG;200MG

N020884 001 Nov 22, 1999

INGELHEIM

ASPIRIN AND DIPYRIDAMOLE

AB AMNEAL PHARMS 25MG;200MG

A206392 001 Mar 08, 2016

AB BARR 25MG;200MG

A078804 001 Aug 14, 2009

AB IMPAX LABS INC 25MG;200MG

A206964 001 Jan 18, 2017

AB PAR PHARM INC 25MG;200MG

A207944 001 Jan 18, 2017

AB SANDOZ INC 25MG;200MG

A206739 001 Jan 18, 2017

AB ZYDUS PHARMS USA 25MG;200MG

A206753 001 Aug 29, 2017

INC

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-41 (of 436)

ASPIRIN; METHOCARBAMOL

TABLET;ORAL

METHOCARBAMOL AND ASPIRIN

! STEVENS J

325MG;400MG

A081145 001 Jan 31, 1995

ASPIRIN; OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL

YOSPRALA

+ ARALEZ PHARMS

81MG;40MG

++!

325MG;40MG

N205103 001 Sep 14, 2016

N205103 002 Sep 14, 2016

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

OXYCODONE AND ASPIRIN

AA ACTAVIS LABS FL INC 325MG;4.8355MG

A090084 001 Mar 22, 2011

AA MAYNE PHARMA INC 325MG;4.8355MG

A091670 001 Mar 16, 2011

PERCODAN

AA +! ENDO PHARMS 325MG;4.8355MG

N007337 007 Aug 05, 2005

ATAZZANAVIR SULFATE

CAPSULE;ORAL

ATAZANAVIR SULFATE

AB TEVA PHARMS USA EQ 150MG BASE

A091673 002 Apr 22, 2014

AB EQ 200MG BASE

A091673 003 Apr 22, 2014

AB EQ 300MG BASE

A091673 004 Apr 22, 2014

REYATAZ

AB + BRISTOL MYERS SQUIBB EQ 150MG BASE

N021567 002 Jun 20, 2003

AB + EQ 200MG BASE

N021567 003 Jun 20, 2003

AB +! EQ 300MG BASE

N021567 004 Oct 16, 2006

ATAZZANAVIR SULFATE

TEVA PHARMS USA EQ 100MG BASE

A091673 001 Apr 22, 2014

POWDER;ORAL

REYATAZ

+! BRISTOL MYERS SQUIBB

EQ 50MG BASE/PACKET

N206352 001 Jun 02, 2014

ATAZZANAVIR SULFATE; COBICISTAT

TABLET;ORAL

EVOTAZ

+! BRISTOL-MYERS SQUIBB

EQ 300MG BASE;150MG

N206353 001 Jan 29, 2015

ATENOLOL

TABLET;ORAL

ATENOLOL

AB ALVOGEN MALTA 25MG

A073646 001 Jul 31, 1992

AB 50MG

A072303 001 Jul 15, 1988

AB 100MG

A072304 001 Jul 15, 1988

AB AUROBINDO PHARMA 25MG

A078512 001 Oct 31, 2007

AB 50MG

A078512 002 Oct 31, 2007

AB 100MG

A078512 003 Oct 31, 2007

AB DAVA PHARMS INC 50MG

A073542 001 Dec 19, 1991

AB 100MG

A073543 001 Dec 19, 1991

AB IPCA LABS LTD 25MG

A077877 001 Dec 27, 2006

AB 50MG

A077877 002 Dec 27, 2006

AB 100MG

A077877 003 Dec 27, 2006

AB MYLAN 25MG

A073457 002 Apr 26, 1999

AB 50MG

A073457 003 Jan 24, 1992

AB 100MG

A073457 001 Jan 24, 1992

AB SANDOZ 25MG

A074052 001 May 01, 1992

AB 50MG

A073025 001 Sep 17, 1991

AB 100MG

A073026 001 Sep 17, 1991

AB SUN PHARM INDNS INC 25MG

A078210 001 Jul 10, 2007

AB 50MG

A078210 002 Jul 10, 2007

AB 100MG

A078210 003 Jul 10, 2007

AB SUN PHARM INDUSTRIES 25MG

A074499 001 Jul 30, 1997

AB 50MG

A073475 001 Mar 30, 1993

AB 100MG

A073476 001 Mar 30, 1993

AB TEVA 25MG

A074056 003 Jul 19, 2004

AB 50MG

A074056 001 Jan 18, 1995

AB 100MG

A074056 002 Jan 18, 1995

AB UNIQUE PHARM LABS 25MG

A077443 001 Sep 13, 2006

AB 50MG

A077443 002 Sep 13, 2006

AB 100MG

A077443 003 Sep 13, 2006

#### ATENOLOL

**TABLET; ORAL**

### **ATENOLOL**

<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A076900</u>	<u>001</u>	Jan 28, 2005
<u>AB</u>		<u>50MG</u>	<u>A076900</u>	<u>002</u>	Jan 28, 2005
<u>AB</u>		<u>100MG</u>	<u>A076900</u>	<u>003</u>	Jan 28, 2005
<b>TENORMIN</b>					
<u>AB</u>	+ ALVOGEN MALTA	<u>25MG</u>	<u>N018240</u>	<u>004</u>	Apr 09, 1990
<u>AB</u>	+	<u>50MG</u>	<u>N018240</u>	<u>001</u>	
<u>AB</u>	+!	<u>100MG</u>	<u>N018240</u>	<u>002</u>	

#### ATENOLOL; CHLORTHALIDONE

**TABLET; ORAL**

## ATENOLOL AND CHLORTHALIDONE

<u>AB</u>	ALVOGEN MALTA	<u>50MG;25MG</u>	<u>A072301</u>	<u>001</u>	May 31, 1990	
<u>AB</u>		<u>100MG;25MG</u>	<u>A072302</u>	<u>001</u>	May 31, 1990	
<u>AB</u>	MYLAN	<u>50MG;25MG</u>	<u>A074203</u>	<u>001</u>	Oct 31, 1993	
<u>AB</u>		<u>100MG;25MG</u>	<u>A074203</u>	<u>002</u>	Oct 31, 1993	
<u>AB</u>	SUN PHARM INDUSTRIES	<u>50MG;25MG</u>	<u>A073582</u>	<u>002</u>	Apr 29, 1993	
<u>AB</u>		<u>100MG;25MG</u>	<u>A073582</u>	<u>001</u>	Apr 29, 1993	
<u>AB</u>	WATSON LABS	<u>50MG;25MG</u>	<u>A073665</u>	<u>001</u>	Jul 02, 1992	
<u>AB</u>		<u>100MG;25MG</u>	<u>A073665</u>	<u>002</u>	Jul 02, 1992	
<u>TENORETIC 100</u>						
<u>AB</u>	+!	ALVOGEN MALTA	<u>100MG;25MG</u>	<u>N018760</u>	<u>001</u>	Jun 08, 1984
<u>TENORETIC 50</u>						
<u>AB</u>	+	ALVOGEN MALTA	<u>50MG;25MG</u>	<u>N018760</u>	<u>002</u>	Jun 08, 1984

## ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

## ATOMOXETINE HYDROCHLORIDE

<u><b>AB</b></u>	APOTEX INC	<u><b>10MG</b></u>	<u><b>A078983</b></u>	<u><b>001</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>18MG</b></u>	<u><b>A078983</b></u>	<u><b>002</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>25MG</b></u>	<u><b>A078983</b></u>	<u><b>003</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>40MG</b></u>	<u><b>A078983</b></u>	<u><b>004</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>60MG</b></u>	<u><b>A078983</b></u>	<u><b>005</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>80MG</b></u>	<u><b>A078983</b></u>	<u><b>006</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A078983</b></u>	<u><b>007</b></u>	May 30, 2017
<u><b>AB</b></u>	AUROBINDO PHARMA LTD	<u><b>10MG</b></u>	<u><b>A079016</b></u>	<u><b>001</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>18MG</b></u>	<u><b>A079016</b></u>	<u><b>002</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>25MG</b></u>	<u><b>A079016</b></u>	<u><b>003</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>40MG</b></u>	<u><b>A079016</b></u>	<u><b>004</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>60MG</b></u>	<u><b>A079016</b></u>	<u><b>005</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>80MG</b></u>	<u><b>A079016</b></u>	<u><b>006</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A079016</b></u>	<u><b>007</b></u>	May 30, 2017
<u><b>AB</b></u>	GLENMARK PHARMS LTD	<u><b>10MG</b></u>	<u><b>A079019</b></u>	<u><b>001</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>18MG</b></u>	<u><b>A079019</b></u>	<u><b>002</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>25MG</b></u>	<u><b>A079019</b></u>	<u><b>003</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>40MG</b></u>	<u><b>A079019</b></u>	<u><b>004</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>60MG</b></u>	<u><b>A079019</b></u>	<u><b>005</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>80MG</b></u>	<u><b>A079019</b></u>	<u><b>006</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A079019</b></u>	<u><b>007</b></u>	May 30, 2017
<u><b>AB</b></u>	TEVA PHARMS USA	<u><b>10MG</b></u>	<u><b>A079022</b></u>	<u><b>001</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>18MG</b></u>	<u><b>A079022</b></u>	<u><b>002</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>25MG</b></u>	<u><b>A079022</b></u>	<u><b>003</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>40MG</b></u>	<u><b>A079022</b></u>	<u><b>004</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>60MG</b></u>	<u><b>A079022</b></u>	<u><b>005</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>80MG</b></u>	<u><b>A079022</b></u>	<u><b>006</b></u>	May 30, 2017
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A079022</b></u>	<u><b>007</b></u>	May 30, 2017

STRATTERA

<u>AB</u>	+	LILLY	<u>10MG</u>	<u>N021411</u>	<u>002</u>	Nov 26, 2002
<u>AB</u>	+		<u>18MG</u>	<u>N021411</u>	<u>003</u>	Nov 26, 2002
<u>AB</u>	+		<u>25MG</u>	<u>N021411</u>	<u>004</u>	Nov 26, 2002
<u>AB</u>	+		<u>40MG</u>	<u>N021411</u>	<u>005</u>	Nov 26, 2002
<u>AB</u>	+!		<u>60MG</u>	<u>N021411</u>	<u>006</u>	Nov 26, 2002
<u>AB</u>	+		<u>80MG</u>	<u>N021411</u>	<u>007</u>	Feb 14, 2005
<u>AB</u>	+		<u>100MG</u>	<u>N021411</u>	<u>008</u>	Feb 14, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-43 (of 436)

ATORVASTATIN CALCIUM

TABLET;ORAL

**ATORVASTATIN CALCIUM**

<b>AB</b>	APOTEX INC	<u>EQ 10MG BASE</u>	<b>A090548 001</b>	May 29, 2012
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A090548 002</b>	May 29, 2012
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A090548 003</b>	May 29, 2012
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A090548 004</b>	May 29, 2012
<b>AB</b>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<b>A091650 001</b>	Jul 17, 2012
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A091650 002</b>	Jul 17, 2012
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A091650 003</b>	Jul 17, 2012
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A202357 001</b>	Jul 17, 2012
<b>AB</b>	INVAGEN PHARMS	<u>EQ 10MG BASE</u>	<b>A204846 001</b>	Jan 09, 2017
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A204846 002</b>	Jan 09, 2017
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A204846 003</b>	Jan 09, 2017
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A204846 004</b>	Jan 09, 2017
<b>AB</b>	KREMERS URBAN PHARMS	<u>EQ 10MG BASE</u>	<b>A091624 001</b>	Apr 05, 2013
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A091624 002</b>	Apr 05, 2013
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A091624 003</b>	Apr 05, 2013
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A091624 004</b>	Apr 05, 2013
<b>AB</b>	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<b>A091226 001</b>	May 29, 2012
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A091226 002</b>	May 29, 2012
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A091226 003</b>	May 29, 2012
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A091226 004</b>	May 29, 2012
<b>AB</b>	SANDOZ INC	<u>EQ 10MG BASE</u>	<b>A077575 001</b>	May 29, 2012
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A077575 002</b>	May 29, 2012
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A077575 003</b>	May 29, 2012
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A077575 004</b>	May 29, 2012
<b>AB</b>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<b>A205519 001</b>	May 19, 2016
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A205519 002</b>	May 19, 2016
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A205519 003</b>	May 19, 2016
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A205519 004</b>	May 19, 2016
<b>AB</b>	SUN PHARM INDs LTD	<u>EQ 10MG BASE</u>	<b>A076477 001</b>	Nov 30, 2011
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A076477 002</b>	Nov 30, 2011
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A076477 003</b>	Nov 30, 2011
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A076477 004</b>	Nov 30, 2011
<b>AB</b>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<b>A205300 001</b>	Mar 27, 2017
<b>AB</b>		<u>EQ 20MG BASE</u>	<b>A205300 002</b>	Mar 27, 2017
<b>AB</b>		<u>EQ 40MG BASE</u>	<b>A205300 003</b>	Mar 27, 2017
<b>AB</b>		<u>EQ 80MG BASE</u>	<b>A205300 004</b>	Mar 27, 2017
<b>LIPITOR</b>				
<b>AB</b>	+ PFIZER	<u>EQ 10MG BASE</u>	<b>N020702 001</b>	Dec 17, 1996
<b>AB</b>	+	<u>EQ 20MG BASE</u>	<b>N020702 002</b>	Dec 17, 1996
<b>AB</b>	+	<u>EQ 40MG BASE</u>	<b>N020702 003</b>	Dec 17, 1996
<b>AB</b>	++!	<u>EQ 80MG BASE</u>	<b>N020702 004</b>	Apr 07, 2000

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET;ORAL

EZETIMIBE AND ATORVASTATIN CALCIUM

WATSON LABS TEVA	EQ 10MG BASE;10MG	A206084 001	Apr 26, 2017
	EQ 20MG BASE;10MG	A206084 002	Apr 26, 2017
	EQ 40MG BASE;10MG	A206084 003	Apr 26, 2017
!	EQ 80MG BASE;10MG	A206084 004	Apr 26, 2017

ATOVAQUONE

SUSPENSION;ORAL

**ATOVAQUONE**

<b>AB</b>	AMNEAL PHARMS	<u>750MG/5ML</u>	<b>A202960 001</b>	Mar 18, 2014
<b>AB</b>	APOTEX INC	<u>750MG/5ML</u>	<b>A209750 001</b>	Oct 11, 2017
<b>AB</b>	PADDOCK LLC	<u>750MG/5ML</u>	<b>A207833 001</b>	Apr 28, 2017

**MEPRON**

<b>AB</b>	++! GLAXOSMITHKLINE LLC	<u>750MG/5ML</u>	<b>N020500 001</b>	Feb 08, 1995
-----------	-------------------------	------------------	--------------------	--------------

ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET;ORAL

**ATOVAQUONE AND PROGUANIL HYDROCHLORIDE**

<b>AB</b>	GLENMARK GENERICS	<u>62.5MG;25MG</u>	<b>A091211 002</b>	Apr 06, 2015
<b>AB</b>		<u>250MG;100MG</u>	<b>A091211 001</b>	Jan 12, 2011
<b>AB</b>	MYLAN PHARMS INC	<u>62.5MG;25MG</u>	<b>A202362 001</b>	May 27, 2014
<b>AB</b>		<u>250MG;100MG</u>	<b>A202362 002</b>	May 27, 2014

**MALARONE**

<b>AB</b>	++! GLAXOSMITHKLINE	<u>250MG;100MG</u>	<b>N021078 001</b>	Jul 14, 2000
-----------	---------------------	--------------------	--------------------	--------------

**MALARONE PEDIATRIC**

<b>AB</b>	++ GLAXOSMITHKLINE	<u>62.5MG;25MG</u>	<b>N021078 002</b>	Jul 14, 2000
-----------	--------------------	--------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-44 (of 436)

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A206011 001</u>	Apr 08, 2015
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090761 001</u>	Oct 18, 2012
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A206096 001</u>	Jun 22, 2017
<u>AP</u>	NANJING KING-FRIEND	<u>10MG/ML</u>	<u>A091489 001</u>	Feb 17, 2012
<u>AP</u> !	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A074901 001</u>	Jul 18, 1997

ATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A206010 001</u>	Apr 08, 2015
<u>AP</u>	HOSPIRA INC	<u>10MG/ML</u>	<u>A090782 001</u>	Oct 18, 2012
<u>AP</u>	MYLAN LABS LTD	<u>10MG/ML</u>	<u>A206001 001</u>	Apr 07, 2017
<u>AP</u>	NANJING KING-FRIEND	<u>10MG/ML</u>	<u>A091488 001</u>	Feb 17, 2012
<u>AP</u> !	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A074900 001</u>	Jul 18, 1997

ATROPINE

INJECTABLE; INJECTION

ATROOPEN

+!	MERIDIAN MEDCL TECHN	EQ 0.25MG SULFATE/0.3ML	N017106 004	Sep 17, 2004
+!		EQ 0.5MG SULFATE/0.7ML	N017106 003	Jun 19, 2003
+!		EQ 1MG SULFATE/0.7ML	N017106 002	Jun 19, 2003
+!		EQ 2MG SULFATE/0.7ML	N017106 001	

ATROPINE SULFATE

SOLUTION; INTRAVENOUS

ATROPINE SULFATE LIFESHIELD ABBOBJECT SYRINGE				
+! HOSPIRA	0.5MG/5ML (0.1MG/ML)		N021146 004	Aug 17, 2017
+!	1MG/10ML (0.1MG/ML)		N021146 005	Aug 17, 2017

SOLUTION; INTRAVENOUS, INTRAMUSCULAR, SUBCUTANEOUS, ENDOTRACHEAL

ATROPINE SULFATE ANSYR PLASTIC SYRINGE				
+! HOSPIRA	0.25MG/5ML (0.05MG/ML)		N021146 002	Jul 09, 2001
+!	0.5MG/5ML (0.1MG/ML)		N021146 001	Jul 09, 2001
+!	1MG/10ML (0.1MG/ML)		N021146 003	Jul 09, 2001

SOLUTION/DROPS; OPHTHALMIC

ATROPINE SULFATE				
+! AKORN	1%		N206289 001	Jul 18, 2014
IISOPTO ATROPINE				
NOVARTIS PHARMS CORP	1%		N208151 001	Dec 01, 2016

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

+! SEBELA IRELAND LTD	0.025MG; 1MG	N017744 002
-----------------------	--------------	-------------

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE				
! WEST-WARD PHARMS	0.025MG/5ML; 2.5MG/5ML		A087708 001	May 03, 1982

INT

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

<u>AA</u>	ANI PHARMS INC	<u>0.025MG; 2.5MG</u>	<u>A086727 001</u>	
<u>AA</u>	LANNETT	<u>0.025MG; 2.5MG</u>	<u>A085372 001</u>	
<u>AA</u>	MYLAN	<u>0.025MG; 2.5MG</u>	<u>A085762 001</u>	
<u>AA</u>	PAR PHARM	<u>0.025MG; 2.5MG</u>	<u>A040357 001</u>	May 02, 2000

LOMOTIL

<u>AA</u> +! GD SEARLE LLC	<u>0.025MG; 2.5MG</u>	<u>N012462 001</u>
----------------------------	-----------------------	--------------------

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR

DUODOTE

+! MERIDIAN MEDCL	2.1MG/0.7ML; 600MG/2ML	N021983 001	Sep 28, 2006
-------------------	------------------------	-------------	--------------

AURANOFIN

CAPSULE; ORAL

RIDAURA

+! SEBELA IRELAND LTD	3MG	N018689 001	May 24, 1985
-----------------------	-----	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-45 (of 436)

**AVANAFIL**

TABLET;ORAL

STENDRA

+ METUCHEN PHARMS	50MG	N202276 001	Apr 27, 2012
+	100MG	N202276 002	Apr 27, 2012
+!	200MG	N202276 003	Apr 27, 2012

**AVIBACTAM SODIUM; CEFTAZIDIME**

POWDER;IV (INFUSION)

AVYCAZ

+! CEREXA	EQ 0.5GM BASE;2GM/VIAL	N206494 001	Feb 25, 2015
-----------	------------------------	-------------	--------------

**AXITINIB**

TABLET;ORAL

INLYTA

+ PF PRISM CV	1MG	N202324 001	Jan 27, 2012
+!	5MG	N202324 002	Jan 27, 2012

**AZACITIDINE**

POWDER;IV (INFUSION), SUBCUTANEOUS

**AZACITIDINE**

<b>AP</b>	ACTAVIS LLC	<b><u>100MG/VIAL</u></b>	<b>N208216 001</b>	Apr 29, 2016
<b>AP</b>	DR REDDYS LABS LTD	<b><u>100MG/VIAL</u></b>	<b>A201537 001</b>	Sep 16, 2013
<b>AP</b>	MYLAN INSTITUTIONAL	<b><u>100MG/VIAL</u></b>	<b>A204949 001</b>	Apr 28, 2016
<b>AP</b>	NATCO PHARMA LTD	<b><u>100MG/VIAL</u></b>	<b>A207234 001</b>	Jun 23, 2017
<b>AP</b>	SHILPA MEDICARE	<b><u>100MG/VIAL</u></b>	<b>A207518 001</b>	Sep 29, 2016

**VIDAZA**

<b>AP</b>	+! CELGENE	<b><u>100MG/VIAL</u></b>	<b>N050794 001</b>	May 19, 2004
-----------	------------	--------------------------	--------------------	--------------

**AZATHIOPRINE**

TABLET;ORAL

**AZASAN**

<b>AB</b>	AAIPHARMA LLC	<b><u>25MG</u></b>	<b>A075252 002</b>	Feb 03, 2003
<b>AB</b>		<b><u>50MG</u></b>	<b>A075252 001</b>	Jun 07, 1999
<b>AB</b>		<b><u>75MG</u></b>	<b>A075252 003</b>	Feb 03, 2003
<b>AB</b>		<b><u>100MG</u></b>	<b>A075252 004</b>	Feb 03, 2003

**AZATHIOPRINE**

<b>AB</b>	AMNEAL PHARMS LLC	<b><u>50MG</u></b>	<b>A074069 001</b>	Feb 16, 1996
<b>AB</b>	MYLAN	<b><u>50MG</u></b>	<b>A075568 001</b>	Dec 13, 1999
<b>AB</b>	ZYDUS PHARMS USA	<b><u>25MG</u></b>	<b>A077621 002</b>	Sep 05, 2008
<b>AB</b>		<b><u>50MG</u></b>	<b>A077621 001</b>	Mar 15, 2007
<b>AB</b>		<b><u>75MG</u></b>	<b>A077621 003</b>	Sep 05, 2008
<b>AB</b>		<b><u>100MG</u></b>	<b>A077621 004</b>	Sep 05, 2008

**IMURAN**

<b>AB</b>	+! SEBELA IRELAND LTD	<b><u>50MG</u></b>	<b>N016324 001</b>	
-----------	-----------------------	--------------------	--------------------	--

**AZATHIOPRINE SODIUM**

INJECTABLE;INJECTION

AZATHIOPRINE SODIUM

!	WEST-WARD PHARMS INT	EQ 100MG BASE/VIAL	A074419 001	Mar 31, 1995
---	----------------------	--------------------	-------------	--------------

**AZELAIC ACID**

AEROSOL, FOAM;TOPICAL

FINACEA

+! BAYER HLTHCARE	15%	N207071 001	Jul 29, 2015
-------------------	-----	-------------	--------------

CREAM;TOPICAL

AZELEX

+! ALLERGAN	20%	N020428 001	Sep 13, 1995
-------------	-----	-------------	--------------

GEL;TOPICAL

FINACEA

+! BAYER HLTHCARE	15%	N021470 001	Dec 24, 2002
-------------------	-----	-------------	--------------

**AZELASTINE HYDROCHLORIDE**

SOLUTION/DROPS;OPHTHALMIC

**AZELASTINE HYDROCHLORIDE**

<b>AT</b>	AKORN	<b><u>0.05%</u></b>	<b>A203660 001</b>	Nov 08, 2016
<b>AT</b>	APOTEX INC	<b><u>0.05%</u></b>	<b>A078621 001</b>	Aug 03, 2009
<b>AT</b>	! SANDOZ INC	<b><u>0.05%</u></b>	<b>A202305 001</b>	May 31, 2012
<b>AT</b>	SUN PHARMA GLOBAL	<b><u>0.05%</u></b>	<b>A078738 001</b>	Jun 21, 2010

SPRAY, METERED;NASAL

ASTELIN

<b>AB</b>	+! MYLAN SPECIALITY LP	<b><u>EQ 0.125MG BASE/SPRAY</u></b>	<b>N020114 001</b>	Nov 01, 1996
-----------	------------------------	-------------------------------------	--------------------	--------------

ASTEPERO

<b>AB</b>	+! MYLAN SPECIALITY LP	<b><u>EQ 0.1876MG BASE/SPRAY</u></b>	<b>N022203 002</b>	Aug 31, 2009
-----------	------------------------	--------------------------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-46 (of 436)

AZELASTINE HYDROCHLORIDE

SPRAY, METERED;NASAL

AZELASTINE HYDROCHLORIDE

<u>AB</u>	ALKEM LABS LTD	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A208156 001</u>	Aug 18, 2017
<u>AB</u>	AMNEAL PHARMS LLC	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A204660 001</u>	Aug 28, 2017
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A208199 001</u>	Dec 15, 2017
<u>AB</u>	APOTEX INC	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A077954 001</u>	Apr 30, 2009
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A201846 001</u>	Aug 31, 2012
<u>AB</u>	BRECKENRIDGE PHARM	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A090176 001</u>	Jul 28, 2015
<u>AB</u>	PERRIGO ISRAEL	<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A202743 001</u>	May 08, 2014
<u>AB</u>	SUN PHARMA GLOBAL	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A090423 001</u>	May 23, 2012
<u>AB</u>	UPSHER-SMITH LABS	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A202609 001</u>	Mar 17, 2017
<u>AB</u>	WEST-WARD PHARMS INT	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A091444 001</u>	Oct 24, 2014
<u>AB</u>		<u>EQ 0.1876MG BASE/SPRAY</u>	<u>A207243 001</u>	Sep 22, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 0.125MG BASE/SPRAY</u>	<u>A091409 001</u>	Aug 14, 2017

AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE

SPRAY, METERED;NASAL

AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE

<u>AB</u>	APOTEX INC	<u>EQ 0.125MG BASE/SPRAY;0.05MG/SPRAY</u>	<u>A207712 001</u>	Apr 28, 2017
<u>DYMISTA</u>			<u>N202236 001</u>	May 01, 2012

AZILSARTAN KAMEDOXOMIL

TABLET;ORAL

EDARBIA

+ ARBOR PHARMS LLC	EQ 40MG MEDOXOMIL
+!	EQ 80MG MEDOXOMIL

N200796 001 Feb 25, 2011  
N200796 002 Feb 25, 2011

AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE

TABLET;ORAL

EDARBYCLOR

+ ARBOR PHARMS LLC	EQ 40MG MEDOXOMIL;12.5MG
+!	EQ 40MG MEDOXOMIL;25MG

N202331 001 Dec 20, 2011  
N202331 002 Dec 20, 2011

AZITHROMYCIN

FOR SUSPENSION;ORAL

AZITHROMYCIN

<u>AB</u>	LUPIN LTD	<u>EQ 100MG BASE/5ML</u>	<u>A065488 001</u>	May 15, 2015
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A065488 002</u>	May 15, 2015
<u>AB</u>	PLIVA	<u>EQ 100MG BASE/5ML</u>	<u>A065246 002</u>	Jul 05, 2006
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A065246 001</u>	Jul 05, 2006
<u>AB</u>	TEVA PHARMS	<u>EQ 100MG BASE/5ML</u>	<u>A065419 001</u>	Jun 24, 2008
<u>AB</u>		<u>EQ 200MG BASE/5ML</u>	<u>A065419 002</u>	Jun 24, 2008

ZITHROMAX

<u>AB</u>	+ PFIZER	<u>EQ 100MG BASE/5ML</u>
<u>AB</u>	+!	<u>EQ 200MG BASE/5ML</u>
	+!	EQ 1GM BASE/PACKET

N050710 001 Oct 19, 1995  
N050710 002 Oct 19, 1995  
N050693 001 Sep 28, 1994

FOR SUSPENSION, EXTENDED RELEASE;ORAL

ZMAX

+! PF PRISM CV	EQ 2GM BASE/BOT
----------------	-----------------

N050797 001 Jun 10, 2005

INJECTABLE;INJECTION

AZITHROMYCIN

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A203294 001</u>	Jun 19, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A065179 001</u>	Dec 13, 2005
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065501 001</u>	Nov 09, 2009
<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>A065500 001</u>	Jun 26, 2009
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A065511 001</u>	Jun 26, 2009
<u>AP</u>	MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A204732 001</u>	Jan 26, 2017
<u>AP</u>	SAGENT STRIDES	<u>EQ 500MG BASE/VIAL</u>	<u>A065506 001</u>	Mar 24, 2009
<u>AP</u>	SUN PHARM IND LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A090923 001</u>	Apr 02, 2013

ZITHROMAX

<u>AP</u>	+! PFIZER	<u>EQ 500MG BASE/VIAL</u>
	SOLUTION/DROPS;OPHTHALMIC AZASITE	

N050733 001 Jan 30, 1997

+! OAK PHARMS INC	1%
-------------------	----

N050810 001 Apr 27, 2007

TABLET;ORAL

AZITHROMYCIN

<u>AB</u>	APOTEX CORP	<u>EQ 250MG BASE</u>	<u>A065507 001</u>	Jul 13, 2011
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065509 001</u>	Jul 13, 2011
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065508 001</u>	Jul 13, 2011
<u>AB</u>	LUPIN LTD	<u>EQ 250MG BASE</u>	<u>A065398 001</u>	May 15, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-47 (of 436)

AZITHROMYCIN

TABLET;ORAL

AZITHROMYCIN

<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065399 001</u>	May 15, 2015
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065400 001</u>	May 15, 2015
<u>AB</u>	MYLAN	<u>EQ 600MG BASE</u>	<u>A065360 001</u>	Jan 08, 2007
<u>AB</u>	PLIVA	<u>EQ 250MG BASE</u>	<u>A065225 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065223 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065218 001</u>	Nov 14, 2005
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>A065211 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065212 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065209 001</u>	Nov 14, 2005
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A065153 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065193 001</u>	Nov 14, 2005
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065150 001</u>	Nov 14, 2005
<u>AB</u>	WOCHARDT	<u>EQ 250MG BASE</u>	<u>A065404 001</u>	Feb 11, 2008
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065405 001</u>	Feb 11, 2008
<u>AB</u>		<u>EQ 600MG BASE</u>	<u>A065302 003</u>	Feb 11, 2008
<u>ZITHROMAX</u>				
<u>AB</u>	+ PFIZER	<u>EQ 250MG BASE</u>	<u>N050711 001</u>	Jul 18, 1996
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N050784 001</u>	May 24, 2002
<u>AB</u>	!+	<u>EQ 600MG BASE</u>	<u>N050730 001</u>	Jun 12, 1996

AZTREONAM

FOR SOLUTION;INHALATION

CAYSTON

+! GILEAD

75MG/VIAL

N050814 001 Feb 22, 2010

INJECTABLE;INJECTION

AZACTAM

<u>AP</u>	+! BRISTOL MYERS SQUIBB	<u>1GM/VIAL</u>	<u>N050580 002</u>	Dec 31, 1986
<u>AP</u>	+!	<u>2GM/VIAL</u>	<u>N050580 003</u>	Dec 31, 1986
<u>AZTREONAM</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>1GM/VIAL</u>	<u>A065439 002</u>	Jun 18, 2010
<u>AP</u>		<u>2GM/VIAL</u>	<u>A065439 003</u>	Jun 18, 2010
AZACTAM IN PLASTIC CONTAINER				
+!	BRISTOL MYERS SQUIBB	20MG/ML	N050632 002	May 24, 1989
+!		40MG/ML	N050632 001	May 24, 1989
AZTREONAM				
	FRESENIUS KABI USA	500MG/VIAL	A065439 001	Jun 18, 2010

BACITRACIN

INJECTABLE;INJECTION

BACIIM

<u>AP</u>	X GEN PHARMS	<u>50,000 UNITS/VIAL</u>	<u>A064153 001</u>	May 09, 1997
<u>BACITRACIN</u>				
<u>AP</u>	AKORN	<u>50,000 UNITS/VIAL</u>	<u>A206719 001</u>	Oct 20, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>50,000 UNITS/VIAL</u>	<u>A065116 001</u>	Dec 03, 2002
<u>AP</u>	! PHARMACIA AND UPJOHN	<u>50,000 UNITS/VIAL</u>	<u>A060733 002</u>	
<u>AP</u>	SAGENT STRIDES	<u>50,000 UNITS/VIAL</u>	<u>A090211 001</u>	May 11, 2010
<u>AP</u>	XELLIA PHARMS APS PHARMACIA AND UPJOHN	<u>50,000 UNITS/VIAL</u>	<u>A203177 001</u>	Aug 25, 2014
	OINTMENT;OPHTHALMIC	10,000 UNITS/VIAL	A060733 001	

BACITRACIN

! PERRIGO CO TENNESSEE

500 UNITS/GM

A061212 001

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

<u>AT</u>	AKORN	<u>400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A065213 001</u>	Jul 25, 2012
<u>AT</u>	! BAUSCH AND LOMB	<u>400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064068 001</u>	Oct 30, 1995
	OINTMENT;TOPICAL CORTISPORIN			
+!	MONARCH PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM	N050168 002	May 04, 1984

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-48 (of 436)

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

<u>AT</u>	AKORN	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A065088 001</u>	Feb 06, 2004
<u>AT</u>	BAUSCH AND LOMB	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A064064 001</u>	Oct 30, 1995
<u>AT</u>	PERRIGO CO TENNESSEE	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>A060764 002</u>	
	<u>NEOSPORIN</u>			
<u>AT</u>	CASPER PHARMA LLC	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N050417 001</u>	

BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

<u>AT</u>	AKORN	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064028 001</u>	Jan 30, 1995
<u>AT</u>	BAUSCH AND LOMB	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A064046 001</u>	Jan 26, 1995
<u>AT</u>	PERRIGO CO TENNESSEE	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>A065022 001</u>	Feb 27, 2002

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

!	PERRIGO CO TENNESSEE	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062166 002
---	-------------------------	--	-------------

BACLOFEN

INJECTABLE;INTRATHECAL

BACLOFEN

<u>AP</u>	EMERALD INTL LTD	<u>0.05MG/ML</u>	<u>A091193 001</u>	May 03, 2016
<u>AP</u>		<u>0.5MG/ML</u>	<u>A091193 002</u>	May 03, 2016
<u>AP</u>		<u>2MG/ML</u>	<u>A091193 003</u>	May 03, 2016

GABLOFEN

<u>AP</u>	PIRAMAL CRITICAL	<u>0.05MG/ML</u>	<u>N022462 001</u>	Nov 19, 2010
<u>AP</u>		<u>0.5MG/ML</u>	<u>N022462 002</u>	Nov 19, 2010
<u>AP</u>		<u>2MG/ML</u>	<u>N022462 003</u>	Nov 19, 2010

LIORESAL

<u>AP</u>	+! SAOL THERAPS RES LTD	<u>0.05MG/ML</u>	<u>N020075 003</u>	Nov 07, 1996
<u>AP</u>	+!	<u>0.5MG/ML</u>	<u>N020075 001</u>	Jun 17, 1992
<u>AP</u>	+!	<u>2MG/ML</u>	<u>N020075 002</u>	Jun 17, 1992

GABLOFEN

+!	PIRAMAL CRITICAL	1MG/ML
----	------------------	--------

TABLET;ORAL

BACLOFEN

<u>AB</u>	IMPAK LABS	<u>10MG</u>	<u>A077971 001</u>	Oct 26, 2007
<u>AB</u>		<u>20MG</u>	<u>A077971 002</u>	Oct 26, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>10MG</u>	<u>A072234 001</u>	Jul 21, 1988
<u>AB</u>	!	<u>20MG</u>	<u>A072235 001</u>	Jul 21, 1988
<u>AB</u>	LANNETT	<u>10MG</u>	<u>A077241 002</u>	Jul 06, 2007
<u>AB</u>		<u>20MG</u>	<u>A077241 001</u>	Dec 20, 2005
<u>AB</u>	MYLAN	<u>20MG</u>	<u>A077121 002</u>	Jul 29, 2005
<u>AB</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A090334 001</u>	Feb 18, 2010
<u>AB</u>		<u>20MG</u>	<u>A090334 002</u>	Feb 18, 2010
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078401 002</u>	Sep 18, 2009
<u>AB</u>		<u>20MG</u>	<u>A078401 001</u>	Sep 18, 2009
<u>AB</u>	OXFORD PHARMS	<u>10MG</u>	<u>A077088 002</u>	Oct 31, 2007
<u>AB</u>		<u>20MG</u>	<u>A077088 001</u>	Oct 31, 2007
<u>AB</u>	RUBICON RES PVT LTD	<u>10MG</u>	<u>A209102 002</u>	Nov 28, 2017
<u>AB</u>		<u>20MG</u>	<u>A209102 003</u>	Nov 28, 2017
<u>AB</u>	SUN PHARM INDs INC	<u>10MG</u>	<u>A077862 001</u>	Aug 14, 2006
<u>AB</u>		<u>20MG</u>	<u>A077862 002</u>	Aug 14, 2006
<u>AB</u>	USL PHARMA	<u>10MG</u>	<u>A074584 001</u>	Aug 19, 1996
<u>AB</u>		<u>20MG</u>	<u>A074584 002</u>	Aug 19, 1996
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A077068 002</u>	Aug 30, 2005
<u>AB</u>		<u>20MG</u>	<u>A077068 001</u>	Aug 30, 2005
	RUBICON RES PVT LTD	5MG	A209102 001	Nov 28, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-49 (of 436)

BALSALAZIDE DISODIUM

CAPSULE;ORAL

BALSALAZIDE DISODIUM

<u>AB</u>	APOTEX INC	<u>750MG</u>	<u>A077883 001</u>	Dec 28, 2007
<u>AB</u>	MYLAN	<u>750MG</u>	<u>A077807 001</u>	Dec 28, 2007
<u>AB</u>	WEST-WARD PHARMS INTL	<u>750MG</u>	<u>A077806 001</u>	Dec 28, 2007

COLAZAL

<u>AB</u>	+!	VALEANT PHARMS INTL	<u>750MG</u>	<u>N020610 001</u>	Jul 18, 2000
-----------	----	---------------------	--------------	--------------------	--------------

TABLET;ORAL

BALSALAZIDE DISODIUM

<u>AB</u>	PAR PHARM INC	<u>1.1GM</u>	<u>A206336 001</u>	Sep 08, 2015	
<u>AB</u>	+!	VALEANT PHARMS INTL	<u>1.1GM</u>	<u>N022205 001</u>	Feb 03, 2012

BARIUM SULFATE

FOR SUSPENSION;ORAL

E-Z-HD

+!	BRACCO	98% (334GM/BOTTLE)	N208036 001	Jan 11, 2016
E-Z-PAQUE		96% (169GM/BOTTLE)	N208036 002	Apr 07, 2017

PASTE;ORAL

VARIBAR

+!	BRACCO	40% (92GM/230ML)	N208844 001	Oct 14, 2016
----	--------	------------------	-------------	--------------

SUSPENSION;ORAL

LIQUID E-Z-PAQUE

+!	BRACCO	60% (213GM/355ML)	N208143 003	Mar 01, 2017
READI-CAT 2		2% (9GM/450ML)	N208143 001	Jan 15, 2016

READI-CAT 2 SMOOTHIES

+!	BRACCO	2% (9GM/450ML)	N208143 002	Jan 15, 2016
----	--------	----------------	-------------	--------------

TAGITOL V

+!	BRACCO	40% (8GM/20ML)	N208143 005	Aug 04, 2017
----	--------	----------------	-------------	--------------

VARIBAR NECTAR

+!	BRACCO	40% (96GM/240ML)	N208143 004	Jul 07, 2017
----	--------	------------------	-------------	--------------

BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED

TABLET;ORAL

DUAVEE

+!	WYETH PHARMS PFIZER	EQ 20MG BASE;0.45MG	N022247 001	Oct 03, 2013
----	---------------------	---------------------	-------------	--------------

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED;INHALATION

QVAR 40

+!	TEVA BRANDED PHARM	0.04MG/INH	N020911 002	Sep 15, 2000
----	--------------------	------------	-------------	--------------

QVAR 80

+!	TEVA BRANDED PHARM	0.08MG/INH	N020911 001	Sep 15, 2000
----	--------------------	------------	-------------	--------------

QVAR REDIHALER

+	NORTON WATERFORD	0.04MG/INH	N207921 001	Aug 03, 2017
---	------------------	------------	-------------	--------------

+!

0.08MG/INH	N207921 002	Aug 03, 2017
------------	-------------	--------------

AEROSOL, METERED;NASAL

QNASL

+	TEVA BRANDED PHARM	0.04MG/ACTUATION	N202813 002	Dec 17, 2014
---	--------------------	------------------	-------------	--------------

+!

0.08MG/ACTUATION	N202813 001	Mar 23, 2012
------------------	-------------	--------------

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED;NASAL

BECONASE AQ

+!	GLAXOSMITHKLINE	EQ 0.042MG DIPROP/SPRAY	N019389 001	Jul 27, 1987
----	-----------------	-------------------------	-------------	--------------

BEDAQUILINE FUMARATE

TABLET;ORAL

SIRTURO

+!	JANSSEN THERAP	EQ 100MG BASE	N204384 001	Dec 28, 2012
----	----------------	---------------	-------------	--------------

BELINOSTAT

POWDER;IV (INFUSION)

BELEODAQ

+!	SPECTRUM PHARMS	500MG/VIAL	N206256 001	Jul 03, 2014
----	-----------------	------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-50 (of 436)

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

**BENAZEPRIL HYDROCHLORIDE**

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A076820</u> <u>001</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A076820</u> <u>002</u>	Feb 03, 2006
<u>AB</u>		<u>20MG</u>	<u>A076820</u> <u>003</u>	Feb 03, 2006
<u>AB</u>		<u>40MG</u>	<u>A076820</u> <u>004</u>	Feb 03, 2006
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A077128</u> <u>001</u>	Mar 08, 2006
<u>AB</u>		<u>10MG</u>	<u>A077128</u> <u>002</u>	Mar 08, 2006
<u>AB</u>		<u>20MG</u>	<u>A077128</u> <u>003</u>	Mar 08, 2006
<u>AB</u>		<u>40MG</u>	<u>A077128</u> <u>004</u>	Mar 08, 2006
<u>AB</u>	AUROBINDO PHARMA	<u>10MG</u>	<u>A078212</u> <u>001</u>	May 22, 2008
<u>AB</u>		<u>20MG</u>	<u>A078212</u> <u>002</u>	May 22, 2008
<u>AB</u>		<u>40MG</u>	<u>A078212</u> <u>003</u>	May 22, 2008
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>5MG</u>	<u>A076333</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076333</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076333</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076333</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076430</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076430</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076430</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076430</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A076118</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076118</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076118</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076118</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A076402</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076402</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076402</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076402</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	SUN PHARM INDs LTD	<u>5MG</u>	<u>A076344</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076344</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076344</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076344</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076211</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>A076211</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>A076211</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>A076211</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	ZYDUS PHARMS USA	<u>5MG</u>	<u>A078848</u> <u>001</u>	May 23, 2008
<u>AB</u>		<u>10MG</u>	<u>A078848</u> <u>002</u>	May 23, 2008
<u>AB</u>		<u>20MG</u>	<u>A078848</u> <u>003</u>	May 23, 2008
<u>AB</u>		<u>40MG</u>	<u>A078848</u> <u>004</u>	May 23, 2008

**LOTENSIN**

<u>AB</u> +	US PHARMS HOLDINGS	<u>5MG</u>	<u>N019851</u> <u>001</u>	Jun 25, 1991
I				
<u>AB</u> +		<u>10MG</u>	<u>N019851</u> <u>002</u>	Jun 25, 1991
<u>AB</u> +		<u>20MG</u>	<u>N019851</u> <u>003</u>	Jun 25, 1991
<u>AB</u> +!		<u>40MG</u>	<u>N019851</u> <u>004</u>	Jun 25, 1991

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

**BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE**

<u>AB</u>	APOTEX INC	<u>5MG;6.25MG</u>	<u>A078794</u> <u>001</u>	Aug 21, 2014
<u>AB</u>		<u>10MG;12.5MG</u>	<u>A078794</u> <u>002</u>	Aug 21, 2014
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A078794</u> <u>003</u>	Aug 21, 2014
<u>AB</u>		<u>20MG;25MG</u>	<u>A078794</u> <u>004</u>	Aug 21, 2014
<u>AB</u>	MYLAN	<u>5MG;6.25MG</u>	<u>A076688</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG;12.5MG</u>	<u>A076688</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076688</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG;25MG</u>	<u>A076688</u> <u>004</u>	Feb 11, 2004
<u>AB</u>	SANDOZ	<u>5MG;6.25MG</u>	<u>A076631</u> <u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG;12.5MG</u>	<u>A076631</u> <u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG;12.5MG</u>	<u>A076631</u> <u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG;25MG</u>	<u>A076631</u> <u>004</u>	Feb 11, 2004

**LOTENSIN HCT**

<u>AB</u> +	US PHARMS HOLDINGS	<u>10MG;12.5MG</u>	<u>N020033</u> <u>002</u>	May 19, 1992
I				
<u>AB</u> +		<u>20MG;12.5MG</u>	<u>N020033</u> <u>004</u>	May 19, 1992
<u>AB</u> +!		<u>20MG;25MG</u>	<u>N020033</u> <u>003</u>	May 19, 1992

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-51 (of 436)

BENDAMUSTINE HYDROCHLORIDE

POWDER;IV (INFUSION)

**BENDAMUSTINE HYDROCHLORIDE**

<b>AP</b>	ACCORD HLTHCARE	<u>25MG/VIAL</u>	<b>A205574 001</b>	May 19, 2016
<b>AP</b>		<u>100MG/VIAL</u>	<b>A205574 002</b>	May 19, 2016
<b>AP</b>	BRECKENRIDGE PHARM	<u>25MG/VIAL</u>	<b>A205447 001</b>	Dec 29, 2016
<b>AP</b>		<u>100MG/VIAL</u>	<b>A205447 002</b>	Dec 29, 2016
<b>AP</b>	GLENMARK PHARMS LTD	<u>25MG/VIAL</u>	<b>A204771 001</b>	Mar 24, 2016
<b>AP</b>		<u>100MG/VIAL</u>	<b>A204771 002</b>	Mar 24, 2016
<b>AP</b>	HOSPIRA INC	<u>25MG/VIAL</u>	<b>A204086 001</b>	May 20, 2016
<b>AP</b>		<u>100MG/VIAL</u>	<b>A204086 002</b>	May 20, 2016
<b>AP</b>	INNOPHARMA LICENSING	<u>25MG/VIAL</u>	<b>A205476 001</b>	Mar 24, 2016
<b>AP</b>		<u>100MG/VIAL</u>	<b>A205476 002</b>	Mar 24, 2016

**TREANDA**

<b>AP</b> +!	CEPHALON	<u>25MG/VIAL</u>	<b>N022249 002</b>	May 01, 2009
<b>AP</b> +!		<u>100MG/VIAL</u>	<b>N022249 001</b>	Mar 20, 2008

SOLUTION;IV (INFUSION)

BENDEKA

+! EAGLE PHARMS 100MG/4ML (25MG/ML)

N208194 001 Dec 07, 2015

TREANDA

+! CEPHALON 45MG/0.5ML (90MG/ML)

N022249 003 Sep 13, 2013

+! 180MG/2ML (90MG/ML)

N022249 004 Sep 13, 2013

BENDROFLUMETHIAZIDE; NADOLOL

TABLET;ORAL

**CORZIDE**

<b>AB</b> +	KING PHARMS LLC	<u>5MG;40MG</u>	<b>N018647 001</b>	May 25, 1983
<b>AB</b> +!		<u>5MG;80MG</u>	<b>N018647 002</b>	May 25, 1983

**NADOLOL AND BENDROFLUMETHIAZIDE**

<b>AB</b>	IMPAK LABS	<u>5MG;40MG</u>	<b>A077833 001</b>	Mar 30, 2007
<b>AB</b>		<u>5MG;80MG</u>	<b>A077833 002</b>	Mar 30, 2007

BENOXINATE HYDROCHLORIDE; FLUORESCIN SODIUM

SOLUTION/DROPS;OPHTHALMIC

ALTAFLUOR BENOX

+! ALTAIRE PHARMS INC 0.4%;0.25%

N208582 001 Dec 14, 2017

BENZNIDAZOLE

TABLET;ORAL

BENZNIDAZOLE

+ CHEMO RESEARCH SL 12.5MG  
+! 100MG

N209570 001 Aug 29, 2017

N209570 002 Aug 29, 2017

BENZONATATE

CAPSULE;ORAL

**BENZONATATE**

<b>AA</b>	APOTEX INC	<u>100MG</u>	<b>A091310 001</b>	Jan 16, 2015
<b>AA</b>		<u>200MG</u>	<b>A091310 002</b>	Jan 16, 2015
<b>AA</b>	BIONPHARMA INC	<u>100MG</u>	<b>A081297 001</b>	Jan 29, 1993
<b>AA</b>		<u>200MG</u>	<b>A081297 002</b>	Oct 30, 2007
<b>AA</b>	CSPC NBP PHARM CO	<u>200MG</u>	<b>A202765 001</b>	Jul 31, 2015
<b>AA</b>	MIKART	<u>100MG</u>	<b>A040851 001</b>	Nov 09, 2009
<b>AA</b>		<u>150MG</u>	<b>A040851 002</b>	Nov 09, 2009
<b>AA</b>		<u>200MG</u>	<b>A040851 003</b>	Nov 09, 2009
<b>AA</b>	ORIT LABS LLC	<u>100MG</u>	<b>A040682 001</b>	Jul 30, 2007
<b>AA</b>		<u>200MG</u>	<b>A040682 002</b>	Jul 30, 2007
<b>AA</b>	STRIDES PHARMA	<u>100MG</u>	<b>A091133 001</b>	Jul 30, 2015
<b>AA</b>		<u>200MG</u>	<b>A091133 002</b>	Jul 30, 2015
<b>AA</b>	SUN PHARM INDs INC	<u>100MG</u>	<b>A040587 001</b>	Mar 19, 2008
<b>AA</b>		<u>200MG</u>	<b>A040587 002</b>	Mar 19, 2008
<b>AA</b> !	THE PHARMA NETWORK	<u>100MG</u>	<b>A040627 001</b>	Mar 30, 2007
<b>AA</b> !		<u>200MG</u>	<b>A040749 001</b>	Jul 25, 2007
<b>AA</b> !	THEPHARMANETWORK LLC	<u>150MG</u>	<b>A201209 001</b>	Sep 24, 2014
<b>AA</b>	ZYDUS PHARMS USA	<u>100MG</u>	<b>A040597 001</b>	Jun 08, 2007
<b>AA</b>		<u>200MG</u>	<b>A040597 002</b>	Jun 08, 2007
<b>AA</b>	<b>TESSALON</b>	<u>100MG</u>	<b>N011210 001</b>	
<b>AA</b> +	PFIZER	<u>100MG</u>		

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-52 (of 436)

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

<u>ACANYA</u>	<u>+!</u>	DOW PHARM	<u>2.5%:EQ 1.2% BASE</u>	<u>N050819 001</u>	Oct 23, 2008	
<u>BENZACLIN</u>			<u>5%:EQ 1% BASE</u>	<u>N050756 001</u>	Dec 21, 2000	
<u>CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE</u>			<u>2.5%:EQ 1.2% BASE</u>	<u>A205128 001</u>	Jun 19, 2015	
		ACTAVIS LABS UT INC	<u>5%:EQ 1% BASE</u>	<u>A065443 001</u>	Aug 11, 2009	
		MYLAN PHARMS INC	<u>5%:EQ 1% BASE</u>	<u>A202440 001</u>	Sep 21, 2015	
		PERRIGO ISRAEL	<u>5%:EQ 1% BASE</u>	<u>A090979 001</u>	Jun 26, 2012	
			<u>5%:1.2%</u>	<u>A206218 001</u>	Dec 15, 2017	
		TARO	<u>5%:1.2%</u>	<u>A204087 001</u>	Jun 27, 2017	
		TOLMAR	<u>5%:EQ 1% BASE</u>	<u>A203688 001</u>	Aug 25, 2016	
			<u>5%:1.2%</u>			
<u>DUAC</u>						
	<u>+!</u>	STIEFEL	<u>5%:1.2%</u>	<u>N050741 001</u>	Aug 26, 2002	
		ONEXTON				
		+!	DOW PHARM	3.75%;EQ 1.2% BASE	N050819 002	Nov 24, 2014

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL;TOPICAL

<u>BENZAMYCIN</u>						
<u>AB</u>	<u>+!</u>	VALEANT INTL	<u>5%:3%</u>	<u>N050557 001</u>	Oct 26, 1984	
		<u>ERYTHROMYCIN AND BENZOYL PEROXIDE</u>				
<u>AB</u>		LYNE	<u>5%:3%</u>	<u>A065385 001</u>	Sep 18, 2015	
<u>AB</u>		TOLMAR	<u>5%:3%</u>	<u>A065112 001</u>	Mar 29, 2004	
		AKTIPAK				
		+!	CUTANEA	5%:3%	N050769 001	Nov 27, 2000

BENZPHETAMINE HYDROCHLORIDE

TABLET;ORAL

<u>BENZPHETAMINE HYDROCHLORIDE</u>					
<u>AA</u>		ANDA REPOSITORY	<u>50MG</u>	<u>A040578 001</u>	Apr 17, 2006
<u>AA</u>		EMCURE PHARMS LTD	<u>50MG</u>	<u>A202061 001</u>	Jan 27, 2012
<u>AA</u>		EPIC PHARMA LLC	<u>50MG</u>	<u>A090346 001</u>	Dec 15, 2015
<u>AA</u>	<u>!</u>	KVK TECH	<u>50MG</u>	<u>A090968 001</u>	Jul 20, 2010
<u>AA</u>		MIKART	<u>25MG</u>	<u>A090473 001</u>	Sep 15, 2010
<u>AA</u>			<u>50MG</u>	<u>A090473 002</u>	Sep 15, 2010
<u>AA</u>		SPECGX LLC	<u>50MG</u>	<u>A040773 001</u>	Apr 25, 2007
<u>AA</u>		TEDOR PHARM	<u>25MG</u>	<u>A040747 002</u>	Nov 20, 2015
<u>AA</u>			<u>50MG</u>	<u>A040747 001</u>	Mar 30, 2007

BENZTROPINE MESYLATE

INJECTABLE;INJECTION

<u>BENZTROPINE MESYLATE</u>					
<u>AP</u>		FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A090233 001</u>	Jul 28, 2009
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A090287 001</u>	Aug 31, 2009
<u>AP</u>		LUITPOLD	<u>1MG/ML</u>	<u>A091152 001</u>	Mar 29, 2010
<u>AP</u>		NAVINTA LLC	<u>1MG/ML</u>	<u>A091525 001</u>	Feb 05, 2013
		<u>COGENTIN</u>			
<u>AP</u>	<u>+!</u>	OAK PHARMS AKORN	<u>1MG/ML</u>	<u>N012015 001</u>	

TABLET;ORAL

<u>BENZTROPINE MESYLATE</u>					
<u>AA</u>		ASPEN GLOBAL INC	<u>0.5MG</u>	<u>A204713 001</u>	Apr 14, 2015
<u>AA</u>			<u>1MG</u>	<u>A204713 002</u>	Apr 14, 2015
<u>AA</u>			<u>2MG</u>	<u>A204713 003</u>	Apr 14, 2015
<u>AA</u>		EPIC PHARMA LLC	<u>0.5MG</u>	<u>A072264 001</u>	Feb 27, 1989
<u>AA</u>			<u>1MG</u>	<u>A072265 001</u>	Feb 27, 1989
<u>AA</u>			<u>2MG</u>	<u>A072266 001</u>	Feb 27, 1989
<u>AA</u>		INVAGEN PHARMS	<u>0.5MG</u>	<u>A090294 001</u>	Mar 29, 2010
<u>AA</u>			<u>1MG</u>	<u>A090294 002</u>	Mar 29, 2010
<u>AA</u>			<u>2MG</u>	<u>A090294 003</u>	Mar 29, 2010
<u>AA</u>		LEADING PHARMA LLC	<u>0.5MG</u>	<u>A090168 001</u>	Nov 28, 2012
<u>AA</u>			<u>1MG</u>	<u>A090168 002</u>	Nov 28, 2012
<u>AA</u>			<u>2MG</u>	<u>A090168 003</u>	Nov 28, 2012
<u>AA</u>		PLIVA	<u>0.5MG</u>	<u>A089058 001</u>	Aug 10, 1988
<u>AA</u>			<u>1MG</u>	<u>A089059 001</u>	Aug 10, 1988
<u>AA</u>			<u>2MG</u>	<u>A089060 001</u>	Aug 10, 1988
<u>AA</u>	<u>!</u>	USL PHARMA	<u>0.5MG</u>	<u>A040103 001</u>	Dec 12, 1996
<u>AA</u>	<u>!</u>		<u>1MG</u>	<u>A040103 002</u>	Dec 12, 1996
<u>AA</u>	<u>!</u>		<u>2MG</u>	<u>A040103 003</u>	Dec 12, 1996
<u>AA</u>		VINTAGE	<u>0.5MG</u>	<u>A040715 001</u>	Aug 27, 2007
<u>AA</u>			<u>1MG</u>	<u>A040715 002</u>	Aug 27, 2007
<u>AA</u>			<u>2MG</u>	<u>A040715 003</u>	Aug 27, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-53 (of 436)

BENZTROPINE MESYLATE

TABLET;ORAL

BENZTROPINE MESYLATE

OXFORD PHARMS

0.5MG

A040706 002 Feb 14, 2008

1MG

A040706 003 Feb 14, 2008

BENZYL ALCOHOL

LOTION;TOPICAL

ULESFIA

+! SHIONOGI INC

5%

N022129 001 Apr 09, 2009

BENZYL PENICILLOYL POLYLYSINE

INJECTABLE;INJECTION

PRE-PEN

+! ALLERQUEST

60UMOLAR

N050114 001

BEPOTASTINE BESILATE

SOLUTION/DROPS;OPHTHALMIC

BEPREVE

+! BAUSCH AND LOMB INC

1.5%

N022288 001 Sep 08, 2009

BERACTANT

SUSPENSION;INTRATRACHEAL

SURVANTA

+! ABBVIE

25MG/ML

N020032 001 Jul 01, 1991

BESIFLOXACIN HYDROCHLORIDE

SUSPENSION/DROPS;OPHTHALMIC

BESIVANCE

+! BAUSCH AND LOMB

EQ 0.6% BASE

N022308 001 May 28, 2009

BETAINE HYDROCHLORIDE

FOR SOLUTION;ORAL

CYSTADANE

+! ORPHAN EUROPE

1GM/SCOOPFUL

N020576 001 Oct 25, 1996

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE

AB LUITPOLD 3MG/ML;EQ 3MG BASE/ML

A090747 001 Jul 31, 2009

CELESTONE SOLUSPAN

AB +! MERCK SHARP DOHME 3MG/ML;EQ 3MG BASE/ML

N014602 001

BETAMETHASONE DIPROPIONATE

CREAM;TOPICAL

BETAMETHASONE DIPROPIONATE

AB ACTAVIS MID EQ 0.05% BASE

A070885 001 Feb 03, 1987

ATLANTIC

AB +! FOUGERA PHARMS EQ 0.05% BASE

N019137 001 Jun 26, 1984

AB TARO EQ 0.05% BASE

A073552 001 Apr 30, 1992

CREAM, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

AB FOUGERA PHARMS EQ 0.05% BASE

A076215 001 Dec 09, 2003

AB GLENMARK GENERICS EQ 0.05% BASE

A078930 001 Sep 23, 2008

AB PERRIGO NEW YORK EQ 0.05% BASE

A076592 001 Dec 09, 2003

AB TARO EQ 0.05% BASE

A076543 001 Dec 09, 2003

AB TOLMAR EQ 0.05% BASE

A076603 001 Jan 23, 2004

DIPROLENE AF

AB +! MERCK SHARP DOHME EQ 0.05% BASE

N019555 001 Apr 27, 1987

GEL, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

AB ! FOUGERA PHARMS EQ 0.05% BASE

A075276 001 May 13, 2003

AB TARO EQ 0.05% BASE

A076508 001 Dec 02, 2003

LOTION;TOPICAL

BETAMETHASONE DIPROPIONATE

AB ACTAVIS MID EQ 0.05% BASE

A070281 001 Jul 31, 1985

ATLANTIC

AB ! FOUGERA PHARMS INC EQ 0.05% BASE

A070275 001 Aug 12, 1985

AB G AND W LABS INC EQ 0.05% BASE

A071467 001 Aug 10, 1987

AB PERRIGO NEW YORK EQ 0.05% BASE

A072538 001 Jan 31, 1990

LOTION, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

AB FOUGERA PHARMS EQ 0.05% BASE

A077111 001 May 21, 2007

AB TARO EQ 0.05% BASE

A077477 001 May 21, 2007

DIPROLENE

AB +! MERCK SHARP DOHME EQ 0.05% BASE

N019716 001 Aug 01, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-54 (of 436)

BETAMETHASONE DIPROPIONATE

OINTMENT;TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.05% BASE</u>	<u>A071012 001</u>	Feb 03, 1987
<u>AB</u>	+! FOUGERA PHARMS INC	<u>EQ 0.05% BASE</u>	<u>N019141 001</u>	Sep 04, 1984
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A074271 001</u>	Sep 15, 1994

OINTMENT, AUGMENTED;TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.05% BASE</u>	<u>A074304 001</u>	Aug 31, 1995
<u>AB</u>	FOUGERA PHARMS	<u>EQ 0.05% BASE</u>	<u>A075373 001</u>	Jun 22, 1999
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>A076753 001</u>	Oct 12, 2004
<u>AB</u>	TELIGENT PHARMA INC	<u>EQ 0.05% BASE</u>	<u>A206118 001</u>	Nov 09, 2017

DIPROLENE

<u>AB</u>	+! MERCK SHARP DOHME SPRAY;TOPICAL	<u>EQ 0.05% BASE</u>	<u>N018741 001</u>	Jul 27, 1983
	SERNIVO +! PROMIUS PHARMA LLC	<u>EQ 0.05% BASE/SPRAY</u>		N208079 001 Feb 05, 2016

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE

AEROSOL, FOAM;TOPICAL

ENSTILAR

+! LEO PHARMA AS 0.064%;0.005%

N207589 001 Oct 16, 2015

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE

OINTMENT;TOPICAL

CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	PERRIGO ISRAEL	<u>0.064%;0.005%</u>	<u>A200174 001</u>	Dec 12, 2014
<u>AB</u>	TOLMAR	<u>0.064%;0.005%</u>	<u>A201615 001</u>	Jan 14, 2013

TACLONEX

<u>AB</u>	+! LEO PHARMA AS SUSPENSION;TOPICAL	<u>0.064%;0.005%</u>	<u>N021852 001</u>	Jan 09, 2006
	TACLONEX +! LEO PHARMA AS	<u>0.064%;0.005%</u>		N022185 001 May 09, 2008

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM;TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.05% BASE;1%</u>	<u>A076002 001</u>	Aug 02, 2002
<u>AB</u>	FOUGERA PHARMS	<u>EQ 0.05% BASE;1%</u>	<u>A075502 001</u>	Jun 05, 2001
<u>AB</u>	GLENMARK PHARMS	<u>EQ 0.05% BASE;1%</u>	<u>A202894 001</u>	Oct 30, 2015
<u>AB</u>	TARO	<u>EQ 0.05% BASE;1%</u>	<u>A075673 001</u>	May 29, 2001

LOTRISONE

<u>AB</u>	+! MERCK SHARP DOHME LOTION;TOPICAL	<u>EQ 0.05% BASE;1%</u>	<u>N018827 001</u>	Jul 10, 1984
-----------	-------------------------------------	-------------------------	--------------------	--------------

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	FOUGERA PHARMS	<u>EQ 0.05% BASE;1%</u>	<u>A076516 001</u>	Jun 16, 2005
<u>AB</u>	TARO	<u>EQ 0.05% BASE;1%</u>	<u>A076493 001</u>	Jul 28, 2004

LOTRISONE

<u>AB</u>	+! MERCK SHARP DOHME	<u>EQ 0.05% BASE;1%</u>	<u>N020010 001</u>	Dec 08, 2000
-----------	----------------------	-------------------------	--------------------	--------------

BETAMETHASONE VALERATE

AEROSOL, FOAM;TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	PERRIGO UK FINCO	<u>0.12%</u>	<u>A078337 001</u>	Nov 26, 2012
<u>AB</u>	RICONPHARMA LLC	<u>0.12%</u>	<u>A207144 001</u>	May 24, 2017
<u>AB</u>	TARO PHARM	<u>0.12%</u>	<u>A208204 001</u>	May 24, 2017

LUXIO

<u>AB</u>	+! MYLAN PHARMS INC	<u>0.12%</u>	<u>N020934 001</u>	Feb 28, 1999
-----------	---------------------	--------------	--------------------	--------------

CREAM;TOPICAL

BETA-VAL

<u>AB</u>	G AND W LABS INC	<u>EQ 0.1% BASE</u>	<u>N018642 001</u>	Mar 24, 1983
-----------	------------------	---------------------	--------------------	--------------

BETAMETHASONE VALERATE

<u>AB</u>	+! FOUGERA PHARMS INC	<u>EQ 0.1% BASE</u>	<u>N018861 001</u>	Aug 31, 1983
-----------	-----------------------	---------------------	--------------------	--------------

DERMABET

<u>AB</u>	TARO	<u>EQ 0.1% BASE</u>	<u>A072041 001</u>	Jan 06, 1988
-----------	------	---------------------	--------------------	--------------

VALNAC

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>EQ 0.1% BASE</u>	<u>A070050 001</u>	Oct 10, 1984
	LOTION;TOPICAL			

BETA-VAL

<u>AB</u>	G AND W LABS INC	<u>EQ 0.1% BASE</u>	<u>A070072 001</u>	Jun 27, 1985
-----------	------------------	---------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-55 (of 436)

BETAMETHASONE VALERATE

LOTION;TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	+!	FOUGERA PHARMS INC	<u>EQ 0.1% BASE</u>	<u>N018866 001</u>	Aug 31, 1983
<u>AB</u>		STI PHARMA LLC	<u>EQ 0.1% BASE</u>	<u>A070052 001</u>	Jul 31, 1985

OINTMENT;TOPICAL

BETA-VAL

<u>AB</u>		G AND W LABS INC	<u>EQ 0.1% BASE</u>	<u>A070069 001</u>	Dec 19, 1985
-----------	--	------------------	---------------------	--------------------	--------------

BETAMETHASONE VALERATE

<u>AB</u>		ACTAVIS MID ATLANTIC	<u>EQ 0.1% BASE</u>	<u>A070051 001</u>	Oct 10, 1984
<u>AB</u>	+!	FOUGERA PHARMS INC	<u>EQ 0.1% BASE</u>	<u>N018865 001</u>	Aug 31, 1983

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

<u>AT</u>		AKORN	<u>EQ 0.5% BASE</u>	<u>A075386 001</u>	Jun 30, 2000
<u>AT</u>		MEDIMETRIKS PHARMS	<u>EQ 0.5% BASE</u>	<u>A075630 001</u>	Apr 12, 2001
<u>AT</u>		WOCKHARDT	<u>EQ 0.5% BASE</u>	<u>A078694 001</u>	Nov 16, 2009

BETOPTIC

<u>AT</u>	+!	ALCON	<u>EQ 0.5% BASE</u>	<u>N019270 001</u>	Aug 30, 1985
		SUSPENSION/DROPS;OPHTHALMIC			
		BETOPTIC S			

+! NOVARTIS PHARMS CORP

TABLET;ORAL

BETAXOLOL HYDROCHLORIDE

<u>AB</u>		EPIC PHARMA	<u>10MG</u>	<u>A075541 001</u>	Oct 22, 1999
<u>AB</u>	!		<u>20MG</u>	<u>A075541 002</u>	Oct 22, 1999
<u>AB</u>		KVK TECH	<u>10MG</u>	<u>A078962 001</u>	Jun 27, 2008
<u>AB</u>			<u>20MG</u>	<u>A078962 002</u>	Jun 27, 2008

BETHANECHOL CHLORIDE

TABLET;ORAL

BETHANECHOL CHLORIDE

<u>AA</u>		AMNEAL PHARM	<u>5MG</u>	<u>A040855 001</u>	Nov 21, 2007
<u>AA</u>			<u>10MG</u>	<u>A040855 002</u>	Nov 21, 2007
<u>AA</u>			<u>25MG</u>	<u>A040855 003</u>	Nov 21, 2007
<u>AA</u>			<u>50MG</u>	<u>A040855 004</u>	Nov 21, 2007
<u>AA</u>		ECI PHARMS LLC	<u>5MG</u>	<u>A040725 001</u>	Oct 26, 2007

<u>AA</u>			<u>10MG</u>	<u>A040726 001</u>	Oct 26, 2007
<u>AA</u>			<u>25MG</u>	<u>A040727 001</u>	Oct 26, 2007
<u>AA</u>			<u>50MG</u>	<u>A040728 001</u>	Oct 26, 2007
<u>AA</u>		HERITAGE PHARMA	<u>5MG</u>	<u>A091256 001</u>	May 04, 2010
<u>AA</u>			<u>10MG</u>	<u>A091256 002</u>	May 04, 2010
<u>AA</u>			<u>25MG</u>	<u>A091256 003</u>	May 04, 2010
<u>AA</u>			<u>50MG</u>	<u>A091256 004</u>	May 04, 2010
<u>AA</u>		LANNETT HOLDINGS INC	<u>5MG</u>	<u>A040677 002</u>	Mar 27, 2008

<u>AA</u>			<u>10MG</u>	<u>A040677 003</u>	Mar 27, 2008
<u>AA</u>			<u>25MG</u>	<u>A040677 004</u>	Mar 27, 2008
<u>AA</u>			<u>50MG</u>	<u>A040677 001</u>	Mar 27, 2008
<u>AA</u>		UPSHER-SMITH LABS	<u>5MG</u>	<u>A040633 001</u>	Jun 01, 2005
<u>AA</u>			<u>10MG</u>	<u>A040634 001</u>	Jun 01, 2005
<u>AA</u>			<u>25MG</u>	<u>A040635 001</u>	Jun 01, 2005
<u>AA</u>			<u>50MG</u>	<u>A040636 001</u>	Jun 01, 2005
<u>AA</u>		WOCKHARDT	<u>5MG</u>	<u>A040532 001</u>	Sep 29, 2003

<u>AA</u>			<u>10MG</u>	<u>A040533 001</u>	Sep 29, 2003
<u>AA</u>			<u>25MG</u>	<u>A040534 001</u>	Sep 29, 2003
<u>AA</u>			<u>50MG</u>	<u>A040518 001</u>	Sep 29, 2003

DUVOID

<u>AA</u>		BI-COASTAL PHARMA	<u>10MG</u>	<u>A086262 001</u>	
<u>AA</u>			<u>25MG</u>	<u>A086263 001</u>	
<u>AA</u>			<u>50MG</u>	<u>A085882 003</u>	

URECHOLINE

<u>AA</u>	!	ODYSSEY PHARMS	<u>5MG</u>	<u>A089095 001</u>	Dec 19, 1985
<u>AA</u>	!		<u>10MG</u>	<u>A088440 001</u>	May 29, 1984
<u>AA</u>	!		<u>25MG</u>	<u>A088441 001</u>	May 29, 1984
<u>AA</u>	!		<u>50MG</u>	<u>A089096 001</u>	Dec 19, 1985

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-56 (of 436)

BETRIXABAN

CAPSULE; ORAL

BEVYXXA

+ PORTOLA PHARMS INC 40MG  
+! 80MG

N208383 001 Jun 23, 2017  
N208383 002 Jun 23, 2017

BEXAROTENE

CAPSULE; ORAL

**BEXAROTENE**

**AB** BIONPHARMA INC **75MG**  
**TARGRETIN**  
**AB** +! VALEANT LUXEMBOURG **75MG**  
GEL; TOPICAL  
TARGRETIN  
+! VALEANT LUXEMBOURG 1%

**A203174 001** Aug 12, 2014  
**N021055 001** Dec 29, 1999  
N021056 001 Jun 28, 2000

BICALUTAMIDE

TABLET; ORAL

**BICALUTAMIDE**

**AB** ACCORD HLTHCARE **50MG**  
**AB** APOTEX INC **50MG**  
**AB** FRESENIUS KABI **50MG**  
ONCOL  
**AB** MYLAN **50MG**  
**AB** SANDOZ **50MG**  
**AB** SANTOS BIOTECH **50MG**  
**AB** SUN PHARMA GLOBAL **50MG**  
**AB** TEVA **50MG**  
**AB** WATSON LABS TEVA **50MG**  
**AB** ZYDUS PHARMS USA **50MG**  
INC

**A078917 001** Jul 06, 2009  
**A200274 001** May 21, 2015  
**A079045 001** May 13, 2010  
**A079185 001** Jul 06, 2009  
**A078575 001** Jul 06, 2009  
**A091011 001** Jun 10, 2015  
**A079110 001** Jul 06, 2009  
**A076932 001** Jul 06, 2009  
**A078634 001** Aug 28, 2009  
**A079089 001** Jul 06, 2009

**CASODEX**

**AB** +! ASTRAZENECA PHARMS **50MG**

**N020498 001** Oct 04, 1995

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC

**BIMATOPROST**

**AT** APOTEX INC **0.03%**  
**AT** ! LUPIN LTD **0.03%**  
**AT** SANDOZ INC **0.03%**  
LUMIGAN  
+! ALLERGAN 0.01%

**A090449 001** Jul 20, 2015  
**A203991 001** Feb 20, 2015  
**A202565 001** May 05, 2015

SOLUTION/DROPS; TOPICAL

**BIMATOPROST**

**AT** APOTEX INC **0.03%**  
**AT** SANDOZ INC **0.03%**

**A201894 001** Dec 01, 2014  
**A202719 001** Apr 19, 2016

**LATISSE**

**AT** +! ALLERGAN **0.03%**

**N022369 001** Dec 24, 2008

BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION, TABLET, DELAYED RELEASE; ORAL

PEG-3350, SODIUM CHLORIDE, SODIUM BICARBONATE, POTASSIUM CHLORIDE AND BISACODYL

! NOVEL LABS INC 5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM A202217 001 Aug 20, 2014  
; N/A, 5.6GM

BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE

CAPSULE; ORAL

PYLERA

+! FOREST LABS LLC 140MG; 125MG; 125MG

N050786 001 Sep 28, 2006

BISOPROLOL FUMARATE

TABLET; ORAL

**BISOPROLOL FUMARATE**

**AB** AUROBINDO PHARMA **5MG**  
**AB** **10MG**  
**AB** MYLAN **5MG**  
**AB** ! **10MG**  
**AB** ORIT LABS LLC **5MG**  
**AB** **10MG**  
**AB** SANDOZ **5MG**  
**AB** **10MG**  
**AB** TEVA PHARMS **5MG**  
**AB** **10MG**  
**AB** UNICHEM PHARMS (USA) **5MG**  
**AB** **10MG**

**A077910 001** Dec 27, 2006  
**A077910 002** Dec 27, 2006  
**A075831 001** Dec 14, 2005  
**A075831 002** Dec 14, 2005  
**A204891 001** Jan 11, 2017  
**A204891 002** Jan 11, 2017  
**A075643 001** Nov 16, 2000  
**A075643 002** Nov 16, 2000  
**A075644 001** Jun 26, 2001  
**A075644 002** Jun 26, 2001  
**A078635 001** Aug 18, 2009  
**A078635 002** Aug 18, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-57 (of 436)

**BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE**

TABLET;ORAL

**BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE**

<b>AB</b>	MYLAN	<u>2.5MG;6.25MG</u>	<b>A075768 001</b>	Sep 25, 2000
<b>AB</b>		<u>5MG;6.25MG</u>	<b>A075768 002</b>	Sep 25, 2000
<b>AB</b>		<u>10MG;6.25MG</u>	<b>A075768 003</b>	Sep 25, 2000
<b>AB</b>	SANDOZ	<u>2.5MG;6.25MG</u>	<b>A075579 001</b>	Sep 25, 2000
<b>AB</b>		<u>5MG;6.25MG</u>	<b>A075579 002</b>	Sep 25, 2000
<b>AB</b>		<u>10MG;6.25MG</u>	<b>A075579 003</b>	Sep 25, 2000
<b>AB</b>	UNICHEM	<u>2.5MG;6.25MG</u>	<b>A079106 001</b>	Jul 28, 2010
<b>AB</b>		<u>5MG;6.25MG</u>	<b>A079106 002</b>	Jul 28, 2010
<b>AB</b>		<u>10MG;6.25MG</u>	<b>A079106 003</b>	Jul 28, 2010
	<b>ZIAC</b>			
<b>AB</b>	+ TEVA BRANDED PHARM	<u>2.5MG;6.25MG</u>	<b>N020186 003</b>	Mar 26, 1993
<b>AB</b>	+	<u>5MG;6.25MG</u>	<b>N020186 001</b>	Mar 26, 1993
<b>AB</b>	++!	<u>10MG;6.25MG</u>	<b>N020186 002</b>	Mar 26, 1993

**BIVALIRUDIN**

INJECTABLE;INTRAVENOUS

**ANGIOMAX**

<b>AP</b>	+! THE MEDICINES CO	<u>250MG/VIAL</u>	<b>N020873 001</b>	Dec 15, 2000
	<b>BIVALIRUDIN</b>			
<b>AP</b>	ACCORD HLTHCARE	<u>250MG/VIAL</u>	<b>A206551 001</b>	Nov 22, 2017
<b>AP</b>	APOTEX INC	<u>250MG/VIAL</u>	<b>A204876 001</b>	Jul 06, 2017
<b>AP</b>	DR REDDYS LABS LTD	<u>250MG/VIAL</u>	<b>A201577 001</b>	May 26, 2017
<b>AP</b>	FRESENIUS KABI USA	<u>250MG/VIAL</u>	<b>A090189 001</b>	Oct 28, 2016
<b>AP</b>	HOSPIRA INC	<u>250MG/VIAL</u>	<b>A090811 001</b>	Jul 14, 2015
<b>AP</b>		<u>250MG/VIAL</u>	<b>A090816 001</b>	Jul 14, 2015

SOLUTION;IV (INFUSION)

BIVALIRUDIN IN 0.9% SODIUM CHLORIDE

+!	CELERITY PHARMS	250MG/50ML (5MG/ML)	N208374 001	Dec 21, 2017
+!		500MG/100ML (5MG/ML)	N208374 002	Dec 21, 2017

**BLEOMYCIN SULFATE**

INJECTABLE;INJECTION

**BLEOMYCIN SULFATE**

<b>AP</b>	! FRESENIUS KABI USA	<u>EQ 15 UNITS BASE/VIAL</u>	<b>A065185 001</b>	Jan 28, 2008
<b>AP</b>	!	<u>EQ 30 UNITS BASE/VIAL</u>	<b>A065185 002</b>	Jan 28, 2008
<b>AP</b>	HOSPIRA	<u>EQ 15 UNITS BASE/VIAL</u>	<b>A065031 001</b>	Mar 10, 2000
<b>AP</b>		<u>EQ 30 UNITS BASE/VIAL</u>	<b>A065031 002</b>	Mar 10, 2000
<b>AP</b>	TEVA PHARMS USA	<u>EQ 15 UNITS BASE/VIAL</u>	<b>A065033 001</b>	Jun 27, 2000
<b>AP</b>		<u>EQ 30 UNITS BASE/VIAL</u>	<b>A065033 002</b>	Jun 27, 2000
<b>AP</b>	WEST-WARD PHARMS INT	<u>EQ 15 UNITS BASE/VIAL</u>	<b>A065042 002</b>	Oct 17, 2001
<b>AP</b>		<u>EQ 30 UNITS BASE/VIAL</u>	<b>A065042 001</b>	Oct 17, 2001

**BORTEZOMIB**

INJECTABLE;INTRAVENOUS, SUBCUTANEOUS

VELCADE

+!	MILLENNIUM PHARMS	3.5MG/VIAL		
----	-------------------	------------	--	--

POWDER;INTRAVENOUS

BORTEZOMIB

FRESENIUS KABI USA	3.5MG/VIAL			
--------------------	------------	--	--	--

**BOSENTAN**

TABLET;ORAL

TRACLEER

+ ACTELION PHARMS LTD	62.5MG			
+!	125MG			

TABLET, FOR SUSPENSION;ORAL

TRACLEER

+! ACTELION PHARMS	32MG			
--------------------	------	--	--	--

N021602 001 May 13, 2003

N205004 001 Nov 06, 2017

N209279 001 Sep 05, 2017

**BOSUTINIB MONOHYDRATE**

TABLET;ORAL

BOSULIF

+! PF PRISM CV	EQ 100MG BASE			
+	EQ 400MG BASE			
+	EQ 500MG BASE			

N203341 001 Sep 04, 2012

N203341 003 Oct 27, 2017

N203341 002 Sep 04, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-58 (of 436)

**BREXPIPRAZOLE**

TABLET;ORAL

REXULTI

+ OTSUKA PHARM CO LTD	0.25MG	N205422 001	Jul 10, 2015
+	0.5MG	N205422 002	Jul 10, 2015
+	1MG	N205422 003	Jul 10, 2015
++!	2MG	N205422 004	Jul 10, 2015
+	3MG	N205422 005	Jul 10, 2015
+	4MG	N205422 006	Jul 10, 2015

**BRIGATINIB**

TABLET;ORAL

ALUNBRIG

+ ARIAD	30MG	N208772 001	Apr 28, 2017
++!	90MG	N208772 002	Apr 28, 2017
+	180MG	N208772 003	Oct 02, 2017

**BRIMONIDINE TARTRATE**

GEL;TOPICAL

MIRVASO

+! GALDERMA LABS LP	EQ 0.33% BASE	N204708 001	Aug 23, 2013
SOLUTION/DROPS;OPHTHALMIC			

**ALPHAGAN P**

<b>AT</b>	+! ALLERGAN	<b>0.15%</b>	<b>N021262 001</b>	Mar 16, 2001
-----------	-------------	--------------	--------------------	--------------

**BRIMONIDINE TARTRATE**

<b>AT</b>	AKORN	<b>0.2%</b>	<b>A076439 001</b>	Mar 14, 2006
<b>AT</b>	! BAUSCH AND LOMB	<b>0.2%</b>	<b>A076260 001</b>	May 28, 2003
<b>AT</b>	INDOCO REMEDIES	<b>0.2%</b>	<b>A091691 001</b>	Nov 18, 2014
<b>AT</b>	SANDOZ INC	<b>0.2%</b>	<b>A076254 001</b>	Sep 16, 2003
<b>AT</b>		<b>0.2%</b>	<b>A078075 001</b>	Jan 30, 2008

**OOLIANA**

<b>AT</b>	SANDOZ INC	<b>0.15%</b>	<b>N021764 001</b>	May 22, 2006
	ALPHAGAN P			
+!	ALLERGAN	0.1%	N021770 001	Aug 19, 2005

**BRIMONIDINE TARTRATE; BRINZOLAMIDE**

SUSPENSION/DROPS;OPHTHALMIC

SIMBRINZA

+! NOVARTIS PHARMS CORP	0.2%;1%	N204251 001	Apr 19, 2013
-------------------------	---------	-------------	--------------

**BRIMONIDINE TARTRATE; TIMOLOL MALEATE**

SOLUTION/DROPS;OPHTHALMIC

COMBIGAN

+! ALLERGAN	0.2%;EQ 0.5% BASE	N021398 001	Oct 30, 2007
-------------	-------------------	-------------	--------------

**BRINZOLAMIDE**

SUSPENSION/DROPS;OPHTHALMIC

AZOPT

+! NOVARTIS PHARMS CORP	1%	N020816 001	Apr 01, 1998
-------------------------	----	-------------	--------------

**BRIVARACETAM**

SOLUTION;INTRAVENOUS

BRIVIACT

+! UCB INC	50MG/5ML (10MG/ML)	N205837 001	May 12, 2016
------------	--------------------	-------------	--------------

SOLUTION;ORAL

BRIVIACT

+! UCB INC	10MG/ML	N205838 001	May 12, 2016
------------	---------	-------------	--------------

TABLET;ORAL

BRIVIACT

+ UCB INC	10MG	N205836 001	May 12, 2016
+	25MG	N205836 002	May 12, 2016
+	50MG	N205836 003	May 12, 2016
+	75MG	N205836 004	May 12, 2016
++!	100MG	N205836 005	May 12, 2016

**BROMFENAC SODIUM**

SOLUTION/DROPS;OPHTHALMIC

**BROMFENAC SODIUM**

<b>AT1</b>	AMRING PHARMS	<b>EQ 0.09% ACID</b>	<b>A202030 001</b>	Jan 09, 2013
<b>AT1</b>	! APOTEX INC	<b>EQ 0.09% ACID</b>	<b>A202435 001</b>	Jun 19, 2014
<b>AT1</b>	PADDICK LLC	<b>EQ 0.09% ACID</b>	<b>A201941 001</b>	Feb 10, 2015
<b>AT2</b>	APOTEX INC	<b>EQ 0.09% ACID</b>	<b>A202620 001</b>	Jun 23, 2014
<b>AT2</b>	! HI-TECH PHARMACAL	<b>EQ 0.09% ACID</b>	<b>A203395 001</b>	Jan 22, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-59 (of 436)

BROMFENAC SODIUM

SOLUTION/DROPS;OPHTHALMIC

BROMSITE

+! SUN PHARMA GLOBAL EQ 0.075% ACID

N206911 001 Apr 08, 2016

PROLENSA

+! BAUSCH AND LOMB EQ 0.07% ACID

N203168 001 Apr 05, 2013

BROMOCRIPTINE MESYLATE

CAPSULE;ORAL

**BROMOCRIPTINE MESYLATE**

**AB** ! MYLAN **EQ 5MG BASE**

**A077226 001** Apr 04, 2005

**AB** ZYDUS PHARMS USA INC **EQ 5MG BASE**

**A078899 001** Jul 30, 2008

**PARLODEL**

**AB** + US PHARMS HOLDINGS I **EQ 5MG BASE**

**N017962 002** Mar 01, 1982

TABLET;ORAL

**BROMOCRIPTINE MESYLATE**

**AB** MYLAN **EQ 2.5MG BASE**

**A076962 001** Sep 24, 2004

**AB** ! PADDOCK LLC **EQ 2.5MG BASE**

**A077646 001** Oct 01, 2008

**AB** SANDOZ INC **EQ 2.5MG BASE**

**A074631 001** Jan 13, 1998

**PARLODEL**

**AB** + US PHARMS HOLDINGS I **EQ 2.5MG BASE**

**N017962 001**

CYCLOSET

+! VEROSCIENCE EQ 0.8MG BASE

N020866 001 May 05, 2009

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP;ORAL

**BROMFED-DM**

**AA** ! WOCKHARDT **2MG/5ML;10MG/5ML;30MG/5ML**

**A088811 001** Jun 07, 1985

**BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE**

**AA** ACELLA PHARMS LLC **2MG/5ML;10MG/5ML;30MG/5ML**

**A203375 001** Sep 20, 2016

**AA** PADDOCK LLC **2MG/5ML;10MG/5ML;30MG/5ML**

**A205292 001** Jul 15, 2014

**AA** TARO PHARM **2MG/5ML;10MG/5ML;30MG/5ML**

**A205112 001** Feb 27, 2017

**AA** VINTAGE PHARMS INC **2MG/5ML;10MG/5ML;30MG/5ML**

**A202940 001** Jul 21, 2014

BUDESONIDE

AEROSOL, FOAM;RECTAL

UCERIS

+! VALEANT PHARMS INTL 2MG/ACTUATION

N205613 001 Oct 07, 2014

CAPSULE;ORAL

**BUDESONIDE**

**AB** ALVOGEN MALTA **3MG**

**A206724 001** Nov 23, 2016

**AB** AMNEAL PHARMS **3MG**

**A206200 001** Jul 31, 2017

**AB** APPCO PHARMA LLC **3MG**

**A207367 001** Apr 07, 2017

**AB** BARR LABS DIV TEVA **3MG**

**A090379 001** Apr 02, 2014

**AB** MAYNE PHARMA **3MG**

**A206623 001** Apr 08, 2016

**AB** MYLAN **3MG**

**A090410 001** May 16, 2011

**AB** SCIECURE PHARMA INC **3MG**

**A209041 001** Sep 28, 2017

**AB** ZYDUS PHARMS USA INC **3MG**

**A206134 001** May 04, 2017

**ENTOCORT EC**

**AB** +! PERRIGO PHARMA INTL **3MG**

**N021324 001** Oct 02, 2001

POWDER, METERED;INHALATION

PULMICORT FLEXHALER

+! ASTRAZENECA 0.08MG/INH

N021949 001 Jul 12, 2006

+! 0.16MG/INH

N021949 002 Jul 12, 2006

SUSPENSION;INHALATION

**BUDESONIDE**

**AN** APOTEX INC **0.25MG/2ML**

**A078202 001** Mar 30, 2009

**AN** 0.5MG/2ML

**A078202 002** Mar 30, 2009

**AN** CIPLA LTD **0.25MG/2ML**

**A205710 001** Nov 16, 2017

**AN** 0.5MG/2ML

**A205710 002** Nov 16, 2017

**AN** 1MG/2ML

**A205710 003** Nov 16, 2017

**AN** IMPAX LABS INC **0.25MG/2ML**

**A078404 001** Jul 31, 2012

**AN** 0.5MG/2ML

**A078404 002** Jul 31, 2012

**AN** SANDOZ INC **0.25MG/2ML**

**A201966 003** Sep 27, 2013

**AN** 0.5MG/2ML

**A201966 002** Sep 27, 2013

**AN** 1MG/2ML

**A201966 001** Sep 27, 2013

**AN** TEVA PHARMS **0.25MG/2ML**

**A077519 001** Nov 18, 2008

**AN** 0.5MG/2ML

**A077519 002** Nov 18, 2008

**AN** TEVA PHARMS USA **1MG/2ML**

**A204548 001** Mar 08, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-60 (of 436)

BUDESONIDE

SUSPENSION; INHALATION

**PULMICORT RESPULES**

<u>AN</u>	+	ASTRAZENECA PHARMS	<u>0.25MG/2ML</u>
<u>AN</u>	+		<u>0.5MG/2ML</u>
<u>AN</u>	+!		<u>1MG/2ML</u>

TABLET, EXTENDED RELEASE; ORAL  
 UCERIS

+! VALEANT PHARMS INTL 9MG

N020929 001 Aug 08, 2000  
N020929 002 Aug 08, 2000  
N020929 003 Aug 08, 2000

N203634 001 Jan 14, 2013

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

AEROSOL, METERED; INHALATION

SYMBICORT

+! ASTRAZENECA 0.08MG/INH; 0.0045MG/INH  
 +! 0.16MG/INH; 0.0045MG/INH

N021929 001 Jul 21, 2006  
N021929 002 Jul 21, 2006

BUMETANIDE

INJECTABLE; INJECTION

**BUMETANIDE**

<u>AP</u>	!	ATHENEX INC	<u>0.25MG/ML</u>
<u>AP</u>		HOSPIRA	<u>0.25MG/ML</u>
<u>AP</u>		WEST-WARD PHARMS INT	<u>0.25MG/ML</u>

TABLET; ORAL

**BUMETANIDE**

<u>AB</u>	AMNEAL PHARMS CO	<u>0.5MG</u>
<u>AB</u>		<u>1MG</u>
<u>AB</u>		<u>2MG</u>
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>0.5MG</u>
<u>AB</u>		<u>1MG</u>
<u>AB</u>		<u>2MG</u>
<u>AB</u>		<u>2MG</u>
		<b>BUMEX</b>
<u>AB</u>	VALIDUS PHARMS	<u>0.5MG</u>
<u>AB</u>	+	<u>1MG</u>
<u>AB</u>	+	<u>2MG</u>

A074441 001 Jan 27, 1995  
A074332 001 Oct 31, 1994  
A079196 001 Apr 30, 2008

A209724 001 Oct 18, 2017  
A209724 002 Oct 18, 2017  
A209724 003 Oct 18, 2017  
A074225 001 Apr 24, 1995  
A074225 002 Apr 24, 1995  
A074225 003 Apr 24, 1995  
A074700 001 Nov 21, 1996  
A074700 002 Nov 21, 1996  
A074700 003 Nov 21, 1996

N018225 002 Feb 28, 1983  
N018225 001 Feb 28, 1983  
N018225 003 Jun 14, 1985

BUPIVACAINE

INJECTABLE, LIPOSOMAL; INJECTION

EXPAREL

+! PACIRA PHARMS INC 133MG/10ML (13.3MG/ML)  
 +! 266MG/20ML (13.3MG/ML)

N022496 001 Oct 28, 2011  
 N022496 002 Oct 28, 2011

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

**BUPIVACAINE HYDROCHLORIDE**

<u>AP</u>	AUROBINDO PHARMA LTD	<u>0.25%</u>
<u>AP</u>		<u>0.5%</u>
<u>AP</u>	HOSPIRA	<u>0.25%</u>
<u>AP</u>		<u>0.25%</u>
<u>AP</u>		<u>0.25%</u>
<u>AP</u>		<u>0.5%</u>
<u>AP</u>		<u>0.75%</u>
<u>AP</u>		<u>0.75%</u>
<u>AP</u>	SAGENT AGILA	<u>0.25%</u>
<u>AP</u>		<u>0.5%</u>
		<b>BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE</b>
<u>AP</u>	AUROBINDO PHARMA LTD	<u>0.25%</u>
<u>AP</u>		<u>0.5%</u>
<u>AP</u>		<u>0.75%</u>
<u>AP</u>	SAGENT AGILA	<u>0.25%</u>
<u>AP</u>		<u>0.5%</u>
<u>AP</u>		<u>0.75%</u>
		<b>MARCAINE HYDROCHLORIDE</b>
<u>AP</u>	+! HOSPIRA	<u>0.25%</u>
<u>AP</u>	+	<u>0.5%</u>

A207183 001 May 13, 2016  
A207183 002 May 13, 2016  
A070583 001 Feb 17, 1987  
A070586 001 Mar 03, 1987  
A070590 001 Feb 17, 1987  
N018053 002  
A070584 001 Feb 17, 1986  
A070597 001 Mar 03, 1987  
A070609 001 Mar 03, 1987  
N018053 001  
A070585 001 Mar 03, 1987  
N018053 003  
A091503 001 Oct 18, 2011  
A091503 002 Oct 18, 2011

A203895 001 Nov 05, 2013  
A203895 002 Nov 05, 2013  
A203895 003 Nov 05, 2013  
A091487 002 Oct 18, 2011  
A091487 001 Oct 18, 2011  
A091487 003 Oct 18, 2011

N016964 001  
N016964 006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-61 (of 436)

## BUPIVACAINE HYDROCHLORIDE

## INJECTABLE; INJECTION

**MARCAINE HYDROCHLORIDE PRESERVATIVE FREE**

<u>AP</u>	+!	HOSPIRA	<u>0.25%</u>	<u>N016964</u>	<u>012</u>
<u>AP</u>	+!		<u>0.5%</u>	<u>N016964</u>	<u>005</u>
<u>AP</u>	+!		<u>0.75%</u>	<u>N016964</u>	<u>009</u>
<b><u>SENSORCAINE</u></b>					
<u>AP</u>		FRESENIUS KABI USA	<u>0.25%</u>	<u>A070552</u>	<u>001</u> May 21, 1986
<u>AP</u>			<u>0.25%</u>	<u>N018304</u>	<u>001</u>
<u>AP</u>			<u>0.5%</u>	<u>A070553</u>	<u>001</u> May 21, 1986
<u>AP</u>			<u>0.5%</u>	<u>N018304</u>	<u>002</u>
<u>AP</u>			<u>0.75%</u>	<u>A070554</u>	<u>001</u> May 21, 1986
<u>AP</u>			<u>0.75%</u>	<u>N018304</u>	<u>003</u>

## INJECTABLE; SPINAL

## BUPIVACAINE HYDROCHLORIDE

<u>AP</u>	BAXTER HLTHCARE CORP	<u>0 .75%</u>	<u>A207266</u> <u>001</u>	Jul 25, 2016
<u>AP</u>	HOSPIRA	<u>0 .75%</u>	<u>A071810</u> <u>001</u>	Dec 11, 1987
	<u>MARCaine</u>			
<u>AP</u>	+! HOSPIRA	<u>0 .75%</u>	<u>N018692</u> <u>001</u>	May 04, 1984
	<u>SENSORCAINE</u>			
<u>AP</u>	PRESENTIUS KART USA	<u>0 .75%</u>	<u>A071202</u> <u>001</u>	Apr 15, 1987

#### BUPIVACAINE HYDROCHLORIDE: EPINEPHRINE

#### INJECTABLE: INJECTION

#### **BUPIVACAINe HYDROCHLORIDE AND EPINEPHRINE**

AP ! HOSPIRA 0.5%;0.005MG/ML 0.5%;0.005MG/ML A071168 001 Jun 16, 1988  
AP ! 0.25%;0.005MG/ML 0.25%;0.005MG/ML A071170 001 Jun 16, 1988  
A071165 001 Jun 16, 1988  
A071167 001 Jun 16, 1988

**BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE**

## INJECTABLE; INJECTION

## BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

AP SEPTODONT 0.5%;0.0091MG/ML A077250 001 Sep 27, 2006  
BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE

AP +! HOSPIRA 0.5%;0.0091MG/ML N022046 001 Jul 13, 1983  
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE

+! HOSPIRA 0.25% ; 0.0091M

<u>AP</u>	+!	<u>0.5%:0.0091MG/ML</u>	<u>N016964</u>	<u>008</u>	
<u>MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE</u>					
<u>AP</u>	+!	<u>HOSPIRA</u>	<u>0.25%:0.0091MG/ML</u>	<u>N016964</u>	<u>013</u>
<u>AP</u>	+!		<u>0.5%:0.0091MG/ML</u>	<u>N016964</u>	<u>007</u>
<u>AP</u>	+!		<u>0.75%:0.0091MG/ML</u>	<u>N016964</u>	<u>010</u>
<u>SENSORCAINE</u>					
<u>AP</u>	FRESENIUS KABI USA	<u>0.25%:0.0091MG/ML</u>	<u>A070966</u>	<u>001</u>	Oct 13, 1987
<u>AP</u>		<u>0.25%:0.0091MG/ML</u>	<u>A070967</u>	<u>001</u>	Oct 13, 1987
<u>AP</u>		<u>0.5%:0.0091MG/ML</u>	<u>A070968</u>	<u>001</u>	Oct 13, 1987
<u>AP</u>		<u>0.5%:0.0091MG/ML</u>	<u>N018304</u>	<u>004</u>	Sep 02, 1983
<u>AP</u>		<u>0.75%:0.0091MG/ML</u>	<u>N018304</u>	<u>005</u>	Sep 02, 1983

BURBENOBRTNE

Film Extended Release Transdermal

BUTTRANS

+ PURDUE PHARMA LP 5MCG/HR N021306 001 Jun 30, 2010  
+ 7.5MCG/HR N021306 005 Jun 30, 2014  
+ 10MCG/HR N021306 002 Jun 30, 2010  
+ 15MCG/HR N021306 004 Jul 25, 2013  
+! 20MCG/HR N021306 003 Jun 30, 2010

SOLUTION, EXTENDED RELEASE; SUBCUTANEOUS

SUBLOCODE

+ INDIVIOR INC 100MG/0.5ML (100MG/0.5ML) N209819 001 Nov 29, 2017  
+! 300MG/1.5ML (200MG/ML) N209819 002 Nov 29, 2017

## BUPRENORPHINE HYDROCHLORIDE

### FILM; BUCCAL

BELBUCA

+ BDSI EQ 0.075MG BASE N207932 001 Oct 23, 2015  
+ EQ 0.15MG BASE N207932 002 Oct 23, 2015  
+ EQ 0.3MG BASE N207932 003 Oct 23, 2015  
+ EQ 0.45MG BASE N207932 004 Oct 23, 2015  
+ EQ 0.6MG BASE N207932 005 Oct 23, 2015  
+ EQ 0.75MG BASE N207932 006 Oct 23, 2015  
+ EQ 0.9MG BASE N207932 007 Oct 23, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-62 (of 436)

BUPRENORPHINE HYDROCHLORIDE

IMPLANT; IMPLANTATION

PROBUPHINE

+! BRAEBURN PHARMS INC EQ 80MG BASE/IMPLANT  
 INJECTABLE; INJECTION

N204442 001 May 26, 2016

BUPRENEK

AP +! INDIVIOR INC EQ 0.3MG BASE/ML

N018401 001

BUPRENORPHINE HYDROCHLORIDE

AP HOSPIRA EQ 0.3MG BASE/ML  
AP LUITPOLD EQ 0.3MG BASE/ML  
AP PAR STERILE EQ 0.3MG BASE/ML  
AP PRODUCTS  
AP WEST-WARD PHARMS EQ 0.3MG BASE/ML  
 INT

A074137 001 Jun 03, 1996  
A078331 001 Mar 27, 2007  
A206586 001 Jul 28, 2015  
A076931 001 Mar 02, 2005

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE

AB ACTAVIS ELIZABETH EQ 2MG BASE  
AB EQ 8MG BASE  
AB BARR EQ 2MG BASE  
AB EQ 8MG BASE  
AB ETHYPHARM EQ 2MG BASE  
AB EQ 8MG BASE  
AB MYLAN PHARMS INC EQ 2MG BASE  
AB EQ 8MG BASE  
AB RHODES PHARMS EQ 2MG BASE  
AB EQ 8MG BASE  
AB SANDOZ INC EQ 2MG BASE  
AB EQ 8MG BASE  
AB SUN PHARM IND'S LTD EQ 2MG BASE  
AB EQ 8MG BASE  
AB WEST-WARD PHARMS EQ 2MG BASE  
 INT  
AB ! EQ 8MG BASE

A090819 001 Feb 19, 2015  
A090819 002 Feb 19, 2015  
A090360 001 May 07, 2010  
A090360 002 May 07, 2010  
A090622 001 Sep 24, 2010  
A090622 002 Sep 24, 2010  
A201066 001 Mar 06, 2015  
A201066 002 Mar 06, 2015  
A207276 001 Mar 27, 2017  
A207276 002 Mar 27, 2017  
A090279 001 Jun 10, 2015  
A090279 002 Jun 10, 2015  
A201760 001 Jan 29, 2016  
A201760 002 Jan 29, 2016  
A078633 001 Oct 08, 2009  
A078633 002 Oct 08, 2009

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

FILM; Buccal

BUNAVAIL

+ BDSI EQ 2.1MG BASE; EQ 0.3MG BASE  
 + EQ 4.2MG BASE; EQ 0.7MG BASE  
 +! EQ 6.3MG BASE; EQ 1MG BASE

N205637 001 Jun 06, 2014  
 N205637 002 Jun 06, 2014  
 N205637 003 Jun 06, 2014

FILM; Buccal, SUBLINGUAL

SUBOXONE

+ INDIVIOR INC EQ 2MG BASE; EQ 0.5MG BASE  
 + EQ 4MG BASE; EQ 1MG BASE  
 + EQ 8MG BASE; EQ 2MG BASE  
 +! EQ 12MG BASE; EQ 3MG BASE

N022410 001 Aug 30, 2010  
 N022410 003 Aug 10, 2012  
 N022410 002 Aug 30, 2010  
 N022410 004 Aug 10, 2012

TABLET; SUBLINGUAL

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

AB ACTAVIS ELIZABETH EQ 2MG BASE; EQ 0.5MG BASE  
AB ! EQ 8MG BASE; EQ 2MG BASE  
AB AMNEAL PHARMS EQ 8MG BASE; EQ 2MG BASE  
AB ETHYPHARM USA CORP EQ 2MG BASE; EQ 0.5MG BASE  
AB EQ 8MG BASE; EQ 2MG BASE  
AB KREMERS URBAN EQ 2MG BASE; EQ 0.5MG BASE  
 PHARMS  
AB EQ 8MG BASE; EQ 2MG BASE  
AB SPECGX LLC EQ 2MG BASE; EQ 0.5MG BASE  
AB EQ 8MG BASE; EQ 2MG BASE  
AB SUN PHARM IND'S LTD EQ 8MG BASE; EQ 2MG BASE  
AB TEVA PHARMS USA EQ 2MG BASE; EQ 0.5MG BASE  
AB EQ 8MG BASE; EQ 2MG BASE  
AB WEST-WARD PHARMS EQ 2MG BASE; EQ 0.5MG BASE  
 INT  
AB EQ 8MG BASE; EQ 2MG BASE

A091422 001 Feb 22, 2013  
A091422 002 Feb 22, 2013  
A203136 002 Feb 22, 2013  
A204431 001 Oct 16, 2015  
A204431 002 Oct 16, 2015  
A205022 001 Sep 19, 2016  
A205022 002 Sep 19, 2016  
A207000 001 Dec 13, 2017  
A207000 002 Dec 13, 2017  
A201633 002 Aug 05, 2016  
A091149 001 Sep 08, 2014  
A091149 002 Sep 08, 2014  
A203326 001 Jun 27, 2014  
A203326 002 Jun 27, 2014

BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE

AB AMNEAL PHARMS EQ 2MG BASE; EQ 0.5MG BASE  
AB SUN PHARM IND'S LTD EQ 2MG BASE; EQ 0.5MG BASE  
 ZUBSOLV  
 + OREXO US INC EQ 0.7MG BASE; EQ 0.18MG BASE  
 + EQ 1.4MG BASE; EQ 0.36MG BASE  
 + EQ 2.9MG BASE; EQ 0.71MG BASE  
 + EQ 5.7MG BASE; EQ 1.4MG BASE  
 + EQ 8.6MG BASE; EQ 2.1MG BASE  
 +! EQ 11.4MG BASE; EQ 2.9MG BASE

N204242 006 Oct 04, 2016  
 N204242 001 Jul 03, 2013  
 N204242 005 Jun 04, 2015  
 N204242 002 Jul 03, 2013  
 N204242 003 Dec 11, 2014  
 N204242 004 Dec 11, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-63 (of 436)

**BUPROPION HYDROBROMIDE**

TABLET, EXTENDED RELEASE; ORAL

APLENZIN

+ VALEANT PHARMS NORTH	174MG	N022108 001	Apr 23, 2008
+ !	348MG	N022108 002	Apr 23, 2008
	522MG	N022108 003	Apr 23, 2008

**BUPROPION HYDROCHLORIDE**

TABLET; ORAL

**BUPROPION HYDROCHLORIDE**

<b>AB</b>	APOTEX INC	<b><u>75MG</u></b>	<b>A076143 001</b>	Jan 17, 2006
<b>AB</b> !		<b><u>100MG</u></b>	<b>A076143 002</b>	Jan 17, 2006
<b>AB</b>	HERITAGE PHARMA	<b><u>75MG</u></b>	<b>A206975 001</b>	Aug 19, 2016
<b>AB</b>		<b><u>100MG</u></b>	<b>A206975 002</b>	Aug 19, 2016
<b>AB</b>	INVAGEN PHARMS	<b><u>75MG</u></b>	<b>A207389 001</b>	Sep 18, 2017
<b>AB</b>		<b><u>100MG</u></b>	<b>A207389 002</b>	Sep 18, 2017
<b>AB</b>	MYLAN	<b><u>75MG</u></b>	<b>A075491 001</b>	Apr 17, 2000
<b>AB</b>		<b><u>100MG</u></b>	<b>A075491 002</b>	Apr 17, 2000
<b>AB</b>	SANDOZ	<b><u>75MG</u></b>	<b>A075584 001</b>	Feb 07, 2000
<b>AB</b>		<b><u>100MG</u></b>	<b>A075584 002</b>	Feb 07, 2000

TABLET, EXTENDED RELEASE; ORAL

**BUPROPION HYDROCHLORIDE**

<b>AB1</b>	ACTAVIS LABS FL INC	<b><u>100MG</u></b>	<b>A079095 001</b>	Mar 24, 2009
<b>AB1</b>		<b><u>150MG</u></b>	<b>A079095 002</b>	Mar 24, 2009
<b>AB1</b>		<b><u>200MG</u></b>	<b>A079095 003</b>	Mar 24, 2009
<b>AB1</b>	ANCHEN PHARMS	<b><u>100MG</u></b>	<b>A091459 001</b>	Jun 09, 2011
<b>AB1</b>		<b><u>150MG</u></b>	<b>A091459 002</b>	Jun 09, 2011
<b>AB1</b>		<b><u>200MG</u></b>	<b>A091459 003</b>	Jun 09, 2011
<b>AB1</b>	IMPAX LABS	<b><u>100MG</u></b>	<b>A075913 001</b>	Jan 28, 2004
<b>AB1</b>		<b><u>150MG</u></b>	<b>A075913 002</b>	Mar 22, 2004
<b>AB1</b>		<b><u>200MG</u></b>	<b>A076711 001</b>	Dec 03, 2004
<b>AB1</b>	INVAGEN PHARMS	<b><u>100MG</u></b>	<b>A206674 001</b>	Feb 09, 2016
<b>AB1</b>		<b><u>150MG</u></b>	<b>A206674 002</b>	Feb 09, 2016
<b>AB1</b>		<b><u>200MG</u></b>	<b>A206674 003</b>	Feb 09, 2016
<b>AB1</b>	JUBILANT GENERICS	<b><u>100MG</u></b>	<b>A202774 001</b>	Oct 11, 2013
<b>AB1</b>		<b><u>150MG</u></b>	<b>A202774 002</b>	Oct 11, 2013
<b>AB1</b>		<b><u>200MG</u></b>	<b>A202774 003</b>	Oct 11, 2013
<b>AB1</b>	MYLAN	<b><u>100MG</u></b>	<b>A090325 001</b>	Apr 08, 2010
<b>AB1</b>		<b><u>150MG</u></b>	<b>A090325 002</b>	Apr 08, 2010
<b>AB1</b>		<b><u>200MG</u></b>	<b>A090325 003</b>	Apr 08, 2010
<b>AB1</b>	PRINSTON INC	<b><u>100MG</u></b>	<b>A202304 001</b>	May 26, 2015
<b>AB1</b>		<b><u>150MG</u></b>	<b>A202304 002</b>	May 26, 2015
<b>AB1</b>		<b><u>200MG</u></b>	<b>A202304 003</b>	May 26, 2015
<b>AB1</b>	SANDOZ	<b><u>100MG</u></b>	<b>A075932 001</b>	Nov 25, 2003
<b>AB1</b>		<b><u>150MG</u></b>	<b>A075932 002</b>	Mar 22, 2004
<b>AB1</b>		<b><u>200MG</u></b>	<b>A075932 003</b>	Jun 22, 2005
<b>AB1</b>	SCIEGEN PHARMS INC	<b><u>100MG</u></b>	<b>A205794 001</b>	Mar 01, 2016
<b>AB1</b>		<b><u>150MG</u></b>	<b>A205794 002</b>	Mar 01, 2016
<b>AB1</b>		<b><u>200MG</u></b>	<b>A205794 003</b>	Mar 01, 2016
<b>AB1</b>	SUN PHARMA GLOBAL	<b><u>100MG</u></b>	<b>A078866 001</b>	Apr 06, 2010
<b>AB1</b>		<b><u>150MG</u></b>	<b>A078866 002</b>	Apr 06, 2010
<b>AB1</b>		<b><u>200MG</u></b>	<b>A078866 003</b>	Apr 06, 2010
<b>AB1</b>	TORRENT PHARMS LTD	<b><u>100MG</u></b>	<b>A203969 001</b>	Oct 31, 2014
<b>AB1</b>		<b><u>150MG</u></b>	<b>A203969 002</b>	Oct 31, 2014
<b>AB1</b>		<b><u>200MG</u></b>	<b>A203969 003</b>	Oct 31, 2014
<b>AB1</b>	WATSON LABS INC	<b><u>100MG</u></b>	<b>A077455 001</b>	Jul 19, 2010
<b>AB1</b>		<b><u>150MG</u></b>	<b>A077455 002</b>	Mar 12, 2008
<b>AB1</b>		<b><u>200MG</u></b>	<b>A077455 003</b>	Jul 19, 2010

**WELLBUTRIN SR**

<b>AB1</b> + GLAXOSMITHKLINE	<b><u>100MG</u></b>	<b>N020358 002</b>	Oct 04, 1996
<b>AB1</b> +	<b><u>150MG</u></b>	<b>N020358 003</b>	Oct 04, 1996
<b>AB1</b> +!	<b><u>200MG</u></b>	<b>N020358 004</b>	Jun 14, 2002

**BUPROPION HYDROCHLORIDE**

<b>AB2</b>	ACTAVIS LABS FL INC	<b><u>150MG</u></b>	<b>A079094 001</b>	Mar 24, 2009
<b>AB2</b>	ANCHEN PHARMS	<b><u>150MG</u></b>	<b>A091520 001</b>	Jun 09, 2011
<b>AB2</b>	IMPAX LABS	<b><u>150MG</u></b>	<b>A075914 001</b>	May 27, 2004
<b>AB2</b>	JUBILANT GENERICS	<b><u>150MG</u></b>	<b>A202775 001</b>	Oct 11, 2013
<b>AB2</b>	MYLAN	<b><u>150MG</u></b>	<b>A090941 001</b>	May 03, 2010
<b>AB2</b>	SANDOZ INC	<b><u>150MG</u></b>	<b>A077475 001</b>	Mar 12, 2008
<b>AB2</b>	TECH ORGANIZED	<b><u>150MG</u></b>	<b>A206122 001</b>	Aug 17, 2016

**ZYBAN**

<b>AB2</b> +! GLAXOSMITHKLINE	<b><u>150MG</u></b>	<b>N020711 003</b>	May 14, 1997
-------------------------------	---------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-64 (of 436)

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

<u>AB3</u>	ACTAVIS LABS FL INC	<u>150MG</u>	<u>A077715</u> <u>001</u>	Nov 26, 2008
<u>AB3</u>	ANBISON LAB CO LTD	<u>150MG</u>	<u>A207224</u> <u>001</u>	Jun 30, 2017
<u>AB3</u>		<u>300MG</u>	<u>A207224</u> <u>002</u>	Jun 30, 2017
<u>AB3</u>	ANCHEN PHARMS	<u>150MG</u>	<u>A077284</u> <u>001</u>	Dec 14, 2006
<u>AB3</u>		<u>300MG</u>	<u>A077284</u> <u>002</u>	Dec 14, 2006
<u>AB3</u>	IMPAX LABS	<u>150MG</u>	<u>A077415</u> <u>001</u>	Nov 26, 2008
<u>AB3</u>	INVAGEN PHARMS	<u>150MG</u>	<u>A206556</u> <u>001</u>	Aug 26, 2016
<u>AB3</u>		<u>300MG</u>	<u>A206556</u> <u>002</u>	Aug 26, 2016
<u>AB3</u>	JUBILANT GENERICS	<u>150MG</u>	<u>A207459</u> <u>001</u>	Jun 30, 2017
<u>AB3</u>		<u>300MG</u>	<u>A207459</u> <u>002</u>	Jun 30, 2017
<u>AB3</u>	LUPIN LTD	<u>150MG</u>	<u>A090693</u> <u>001</u>	Apr 06, 2017
<u>AB3</u>		<u>300MG</u>	<u>A090693</u> <u>002</u>	Apr 06, 2017
<u>AB3</u>	MYLAN	<u>150MG</u>	<u>A090942</u> <u>001</u>	Jul 14, 2010
<u>AB3</u>		<u>300MG</u>	<u>A090942</u> <u>002</u>	Jul 14, 2010
<u>AB3</u>	SCIEGEN PHARMS INC	<u>150MG</u>	<u>A207479</u> <u>001</u>	Apr 12, 2017
<u>AB3</u>		<u>300MG</u>	<u>A207479</u> <u>002</u>	Apr 12, 2017
<u>AB3</u>	SINOOTHERAPEUTICS INC	<u>150MG</u>	<u>A208652</u> <u>001</u>	Aug 21, 2017
<u>AB3</u>		<u>300MG</u>	<u>A208652</u> <u>002</u>	Aug 21, 2017
<u>AB3</u>	SUN PHARMA GLOBAL	<u>150MG</u>	<u>A200695</u> <u>001</u>	Dec 18, 2014
<u>AB3</u>	TWI PHARMS INC	<u>150MG</u>	<u>A210081</u> <u>001</u>	Nov 03, 2017
<u>AB3</u>		<u>300MG</u>	<u>A210081</u> <u>002</u>	Nov 03, 2017
<u>AB3</u>	WATSON LABS INC	<u>150MG</u>	<u>A077285</u> <u>001</u>	Nov 26, 2008
<u>AB3</u>		<u>300MG</u>	<u>A077285</u> <u>002</u>	Aug 15, 2008
<u>AB3</u>	WOCKHARDT LTD	<u>150MG</u>	<u>A202189</u> <u>001</u>	Nov 21, 2012
<u>AB3</u>	ZYDUS PHARMS USA INC	<u>300MG</u>	<u>A201567</u> <u>001</u>	Jan 17, 2014

WELLBUTRIN XL

<u>AB3</u> + VALEANT INTL	<u>150MG</u>	<u>N021515</u> <u>001</u>	Aug 28, 2003
<u>AB3</u> +! FORFIVO XL	<u>300MG</u>	<u>N021515</u> <u>002</u>	Aug 28, 2003
+! ALVOGEN	450MG		
		N022497 001	Nov 10, 2011

BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CONTRAVE

+! OREXIGEN 90MG; 8MG

N200063 001 Sep 10, 2014

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPIRONE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A202557</u> <u>001</u>	Dec 30, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202557</u> <u>002</u>	Dec 30, 2014
<u>AB</u>		<u>10MG</u>	<u>A202557</u> <u>003</u>	Dec 30, 2014
<u>AB</u>		<u>15MG</u>	<u>A202557</u> <u>004</u>	Dec 30, 2014
<u>AB</u>		<u>30MG</u>	<u>A202557</u> <u>005</u>	Dec 30, 2014
<u>AB</u>	AMNEAL PHARMS CO	<u>5MG</u>	<u>A208829</u> <u>001</u>	May 24, 2017
<u>AB</u>		<u>7.5MG</u>	<u>A208829</u> <u>002</u>	May 24, 2017
<u>AB</u>		<u>10MG</u>	<u>A208829</u> <u>003</u>	May 24, 2017
<u>AB</u>		<u>15MG</u>	<u>A208829</u> <u>004</u>	May 24, 2017
<u>AB</u>		<u>30MG</u>	<u>A208829</u> <u>005</u>	May 24, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A078246</u> <u>001</u>	Feb 27, 2009
<u>AB</u>		<u>10MG</u>	<u>A078246</u> <u>002</u>	Feb 27, 2009
<u>AB</u>		<u>15MG</u>	<u>A078246</u> <u>003</u>	Feb 27, 2009
<u>AB</u>		<u>30MG</u>	<u>A078246</u> <u>004</u>	Feb 27, 2009
<u>AB</u>	HERITAGE PHARMA	<u>5MG</u>	<u>A204582</u> <u>001</u>	Sep 18, 2015
<u>AB</u>		<u>10MG</u>	<u>A204582</u> <u>002</u>	Sep 18, 2015
<u>AB</u>		<u>15MG</u>	<u>A204582</u> <u>003</u>	Sep 18, 2015
<u>AB</u>		<u>30MG</u>	<u>A204582</u> <u>004</u>	Sep 18, 2015
<u>AB</u>	IMPAX LABS INC	<u>5MG</u>	<u>A074253</u> <u>001</u>	Mar 28, 2001
<u>AB</u>		<u>10MG</u>	<u>A074253</u> <u>002</u>	Mar 28, 2001
<u>AB</u>		<u>15MG</u>	<u>A074253</u> <u>003</u>	Mar 13, 2002
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076008</u> <u>003</u>	Mar 01, 2002
<u>AB</u>		<u>7.5MG</u>	<u>A075467</u> <u>002</u>	Mar 28, 2001
<u>AB</u>		<u>7.5MG</u>	<u>A076008</u> <u>002</u>	Jul 08, 2013
<u>AB</u>		<u>10MG</u>	<u>A076008</u> <u>004</u>	Mar 01, 2002
<u>AB</u>		<u>15MG</u>	<u>A076008</u> <u>005</u>	Mar 28, 2001
<u>AB</u>		<u>30MG</u>	<u>A076008</u> <u>001</u>	Jun 28, 2001
<u>AB</u>	ORION CORP ORION	<u>5MG</u>	<u>A202087</u> <u>001</u>	Dec 16, 2015
<u>AB</u>		<u>10MG</u>	<u>A202087</u> <u>002</u>	Dec 16, 2015
<u>AB</u>		<u>15MG</u>	<u>A202087</u> <u>003</u>	Dec 16, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-65 (of 436)

BUSPIRONE HYDROCHLORIDE

TABLET;ORAL

**BUSPIRONE HYDROCHLORIDE**

<b>AB</b>		<u>30MG</u>	<b>A202087 004</b>	Dec 16, 2015
<b>AB</b>	OXFORD PHARMS	<u>30MG</u>	<b>A078302 001</b>	Dec 17, 2007
<b>AB</b>	STRIDES PHARMA	<u>5MG</u>	<b>A202330 001</b>	Aug 25, 2014
<b>AB</b>		<u>7.5MG</u>	<b>A202330 005</b>	Feb 17, 2017
<b>AB</b>		<u>10MG</u>	<b>A202330 002</b>	Aug 25, 2014
<b>AB</b>		<u>15MG</u>	<b>A202330 003</b>	Aug 25, 2014
<b>AB</b>		<u>30MG</u>	<b>A202330 004</b>	Aug 25, 2014
<b>AB</b>	TEVA	<u>5MG</u>	<b>A075022 001</b>	Feb 28, 2002
<b>AB</b>		<u>10MG</u>	<b>A075022 002</b>	Feb 28, 2002
<b>AB</b>	!	<u>15MG</u>	<b>A075022 003</b>	Feb 28, 2002
<b>AB</b>		<u>30MG</u>	<b>A075022 004</b>	Mar 25, 2004
<b>AB</b>	ZYDUS PHARMS USA INC	<u>5MG</u>	<b>A078888 001</b>	Feb 07, 2014
<b>AB</b>		<u>10MG</u>	<b>A078888 002</b>	Feb 07, 2014
<b>AB</b>		<u>15MG</u>	<b>A078888 003</b>	Feb 07, 2014
<b>AB</b>		<u>30MG</u>	<b>A078888 004</b>	Feb 07, 2014

BUSULFAN

INJECTABLE; INJECTION

**BUSULFAN**

<b>AP</b>	ACTAVIS LLC	<u>6MG/ML</u>	<b>A205139 001</b>	Dec 08, 2017
<b>AP</b>	AMNEAL PHARMS CO	<u>6MG/ML</u>	<b>A209580 001</b>	Dec 18, 2017
<b>AP</b>	LUITPOLD PHARMS INC	<u>6MG/ML</u>	<b>A202259 001</b>	Dec 22, 2015
<b>AP</b>	PHARMASCIENCE INC	<u>6MG/ML</u>	<b>A207050 001</b>	Mar 24, 2017
<b>AP</b>	SANDOZ INC	<u>6MG/ML</u>	<b>A205106 001</b>	Jul 07, 2017
	<b>BUSULFEX</b>			
<b>AP</b>	+! OTSUKA PHARM	<u>6MG/ML</u>	<b>N020954 001</b>	Feb 04, 1999
	<b>MYLERAN</b>			
<b>AP</b>	ASPEN GLOBAL INC	<u>6MG/ML</u>	<b>A208536 001</b>	Nov 20, 2017

TABLET;ORAL

MYLERAN

+! ASPEN GLOBAL 2MG

N009386 001

BUTABARBITAL SODIUM

TABLET;ORAL

BUTISOL SODIUM

+! MYLAN SPECIALITY LP 30MG

N000793 004

BUTENAFINE HYDROCHLORIDE

CREAM;TOPICAL

MENTAX

+! MYLAN 1%

N020524 001 Oct 18, 1996

BUTOCONAZOLE NITRATE

CREAM;VAGINAL

GYNAZOLE-1

+! PERRIGO ISRAEL 2%

A200923 001 May 18, 2012

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

**BUTORPHANOL TARTRATE**

<b>AP</b>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<b>A078400 001</b>	May 01, 2009
<b>AP</b>		<u>2MG/ML</u>	<b>A078400 002</b>	May 01, 2009
<b>AP</b>	WEST-WARD PHARMS INT	<u>2MG/ML</u>	<b>A075046 001</b>	Aug 12, 1998

**BUTORPHANOL TARTRATE PRESERVATIVE FREE**

<b>AP</b>	HOSPIRA	<u>1MG/ML</u>	<b>A074626 001</b>	Jan 23, 1997
<b>AP</b>		<u>2MG/ML</u>	<b>A074626 002</b>	Jan 23, 1997
<b>AP</b>	WEST-WARD PHARMS INT	<u>1MG/ML</u>	<b>A075045 001</b>	Aug 12, 1998
<b>AP</b>		<u>2MG/ML</u>	<b>A075045 002</b>	Aug 12, 1998

SPRAY, METERED;NASAL

**BUTORPHANOL TARTRATE**

<b>AB</b>	APOTEX INC	<u>1MG/SPRAY</u>	<b>A075499 001</b>	Dec 04, 2002
<b>AB</b>	! MYLAN	<u>1MG/SPRAY</u>	<b>A075759 001</b>	Aug 08, 2001
<b>AB</b>	WEST-WARD PHARMS INT	<u>1MG/SPRAY</u>	<b>A075824 001</b>	Mar 12, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-66 (of 436)

**CABAZITAXEL**

SOLUTION;IV (INFUSION)  
 JEVDTANA KIT  
 +! SANOFI AVENTIS US 60MG/1.5ML (40MG/ML)

N201023 001 Jun 17, 2010

**CABERGOLINE**

TABLET;ORAL

**CABERGOLINE**

<b>AB</b>	ACTAVIS LABS FL INC	<b>0.5MG</b>	<b>A078035 001</b>	Apr 21, 2008
<b>AB</b>	APOTEX CORP	<b>0.5MG</b>	<b>A201503 001</b>	Mar 08, 2013
<b>AB</b>	IVAX SUB TEVA PHARMS	<b>0.5MG</b>	<b>A077750 001</b>	Mar 07, 2007
<b>AB</b>	MYLAN PHARMS INC	<b>0.5MG</b>	<b>A202947 001</b>	Dec 02, 2013
<b>AB</b>	+! PAR PHARM	<b>0.5MG</b>	<b>A076310 001</b>	Dec 29, 2005

**CABOZANTINIB S-MALATE**

CAPSULE;ORAL

COMETRIQ

+! EXELIXIS EQ 20MG BASE  
 +! EQ 80MG BASE

N203756 001 Nov 29, 2012  
 N203756 002 Nov 29, 2012

TABLET;ORAL

CABOMETYX

+ EXELIXIS INC EQ 20MG BASE  
 + EQ 40MG BASE  
 +! EQ 60MG BASE

N208692 001 Apr 25, 2016  
 N208692 002 Apr 25, 2016  
 N208692 003 Apr 25, 2016

**CAFFEINE CITRATE**

SOLUTION;INTRAVENOUS

**CAFECIT**

<b>AP</b>	+! WEST-WARD PHARMS INT	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>N020793 001</b>	Sep 21, 1999
-----------	-------------------------	---	--------------------	--------------

**CAFFEINE CITRATE**

<b>AP</b>	AUROBINDO PHARMA LTD	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A205013 001</b>	Sep 22, 2015
<b>AP</b>	EXELA PHARMA SCIENCE	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A077233 001</b>	Sep 21, 2006
<b>AP</b>	FRESENIUS KABI USA	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A077997 001</b>	Jul 20, 2007
<b>AP</b>	LUITPOLD	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A077906 001</b>	May 15, 2007
<b>AP</b>	MICRO LABS	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A207400 001</b>	Dec 14, 2017
<b>AP</b>	SAGENT PHARMS	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A090827 001</b>	Aug 29, 2012
<b>AP</b>	SUN PHARMA GLOBAL	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A090077 001</b>	Sep 30, 2009

SOLUTION;ORAL

**CAFECIT**

<b>AA</b>	+! WEST-WARD PHARMS INT	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>N020793 002</b>	Apr 12, 2000
-----------	-------------------------	---	--------------------	--------------

**CAFFEINE CITRATE**

<b>AA</b>	EXELA PHARMA SCS LLC	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A077304 001</b>	Sep 21, 2006
<b>AA</b>	FRESENIUS KABI USA	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A078002 001</b>	Jan 31, 2008
<b>AA</b>	LUITPOLD	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A090064 001</b>	Nov 20, 2009
<b>AA</b>	SAGENT PHARMS	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A091102 001</b>	Aug 29, 2012
<b>AA</b>	SUN PHARMA GLOBAL	<b>EQ 30MG BASE/3ML (EQ 10MG BASE/ML)</b>	<b>A090357 001</b>	Sep 30, 2009

**CAFFEINE; ERGOTAMINE TARTRATE**

SUPPOSITORY;RECTAL

MIGERGOT

! HORIZON PHARMA 100MG;2MG

A086557 001 Oct 04, 1983

TABLET;ORAL

**CAFERGOT**

<b>AA</b>	+! SANDOZ	<b>100MG;1MG</b>	<b>A084294 001</b>	
-----------	-----------	------------------	--------------------	--

**ERGOTAMINE TARTRATE AND CAFFEINE**

<b>AA</b>	HIKMA INTL PHARMS	<b>100MG;1MG</b>	<b>A040510 001</b>	Sep 17, 2004
<b>AA</b>	MIKART	<b>100MG;1MG</b>	<b>A040590 001</b>	Sep 16, 2005

**CALCIFEDIOL**

CAPSULE, EXTENDED RELEASE;ORAL

RAYALDEE

+! OPKO IRELAND GLOBAL 0.03MG

N208010 001 Jun 17, 2016

**CALCIPOTRIENE**

AEROSOL, FOAM;TOPICAL

SORILUX

+! MAYNE PHARMA 0.005%

N022563 001 Oct 06, 2010

CREAM;TOPICAL

**CALCIPOTRIENE**

<b>AB</b>	GLENMARK PHARMS	<b>0.005%</b>	<b>A205772 001</b>	Jun 09, 2015
<b>AB</b>	TOLMAR	<b>0.005%</b>	<b>A200935 001</b>	May 30, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-67 (of 436)

CALCIPOTRIENE

CREAM;TOPICAL

DOVONEX

<u>AB</u>	+!	LEO PHARMA AS	<u>0.005%</u>	<u>N020554 001</u>	Jul 22, 1996
OINTMENT;TOPICAL					
CALCIPOTRIENE					
! GLENMARK PHARMS INC	0.005%			A090633 001	Mar 24, 2010

SOLUTION;TOPICAL

CALCIPOTRIENE

<u>AT</u>	FOUGERA PHARMS	<u>0.005%</u>	<u>A078305 001</u>	May 06, 2008
<u>AT</u>	G AND W LABS INC	<u>0.005%</u>	<u>A078468 001</u>	Mar 24, 2011
<u>AT</u>	HI TECH PHARMA	<u>0.005%</u>	<u>A077579 001</u>	Nov 19, 2009
<u>AT</u>	NOVEL LABS INC	<u>0.005%</u>	<u>A207163 001</u>	Dec 26, 2017
<u>AT</u>	! TOLMAR	<u>0.005%</u>	<u>A077029 001</u>	Nov 20, 2009

CALCITONIN SALMON

INJECTABLE; INJECTION

MIACALCIN

! MYLAN IRELAND LTD	200 IU/ML	N017808 002	Mar 29, 1991
SPRAY, METERED;NASAL			

CALCITONIN-SALMON

<u>AB</u>	! APOTEX INC	<u>200 IU/SPRAY</u>	<u>A076396 001</u>	Nov 17, 2008
<u>AB</u>	PAR PHARM	<u>200 IU/SPRAY</u>	<u>A076979 001</u>	Jun 08, 2009

CALCITRIOL

CAPSULE;ORAL

CALCITRIOL

<u>AB</u>	AMNEAL PHARMS	<u>0.25MCG</u>	<u>A203289 002</u>	Jun 14, 2017
<u>AB</u>		<u>0.5MCG</u>	<u>A203289 001</u>	Jun 14, 2017
<u>AB</u>	BIONPHARMA INC	<u>0.25MCG</u>	<u>A091174 001</u>	May 24, 2013
<u>AB</u>		<u>0.5MCG</u>	<u>A091174 002</u>	May 24, 2013
<u>AB</u>	STRIDES PHARMA	<u>0.25MCG</u>	<u>A091356 001</u>	Dec 12, 2014
<u>AB</u>		<u>0.5MCG</u>	<u>A091356 002</u>	Dec 12, 2014
<u>AB</u>	TEVA	<u>0.25MCG</u>	<u>A075765 001</u>	Oct 12, 2001
<u>AB</u>		<u>0.5MCG</u>	<u>A075765 002</u>	Oct 12, 2001
<u>AB</u>	WEST-WARD PHARMS	<u>0.25MCG</u>	<u>A076917 001</u>	Mar 27, 2006
	INT			

ROCALTROL

<u>AB</u>	+ VALIDUS PHARMS	<u>0.25MCG</u>	<u>N018044 001</u>	
<u>AB</u>	+!	<u>0.5MCG</u>	<u>N018044 002</u>	

INJECTABLE; INJECTION

CALCITRIOL

!	AKORN	0.001MG/ML	A078066 001	Jan 29, 2008
OINTMENT;TOPICAL				
VECTICAL				
+! GALDERMA LABS LP	3MCG/GM		N022087 001	Jan 23, 2009

SOLUTION;ORAL

CALCITRIOL

<u>AA</u>	WEST-WARD PHARMS	<u>1MCG/ML</u>	<u>A076242 001</u>	Jul 18, 2003
<u>AA</u>	INT			

ROCALTROL

<u>AA</u>	+! VALIDUS PHARMS	<u>1MCG/ML</u>	<u>N021068 001</u>	Nov 20, 1998
-----------	-------------------	----------------	--------------------	--------------

CALCIUM ACETATE

CAPSULE;ORAL

CALCIUM ACETATE

<u>AB</u>	AMNEAL PHARMS	<u>667MG</u>	<u>A201658 001</u>	Oct 06, 2014
<u>AB</u>	CHARTWELL RX	<u>667MG</u>	<u>A091312 001</u>	Jun 01, 2012
<u>AB</u>	ECI PHARMS LLC	<u>667MG</u>	<u>A203298 001</u>	Jul 26, 2016
<u>AB</u>	HERITAGE PHARMS INC	<u>667MG</u>	<u>A202315 001</u>	Jun 29, 2015
<u>AB</u>	INVAGEN PHARMS	<u>667MG</u>	<u>A203135 001</u>	Feb 07, 2013
<u>AB</u>	LUPIN LTD	<u>667MG</u>	<u>A202127 001</u>	Jul 09, 2015
<u>AB</u>	NOSTRUM LABS INC	<u>667MG</u>	<u>A203179 001</u>	Oct 26, 2015
<u>AB</u>	WEST-WARD PHARMS	<u>667MG</u>	<u>A077728 001</u>	Feb 26, 2008
	INT			

PHOSLO GELCAPS

<u>AB</u>	+! FRESENIUS MEDCL	<u>667MG</u>	<u>N021160 003</u>	Apr 02, 2001
OINTMENT;ORAL				
PHOSLYRA				

+! FRESENIUS MEDCL

TABLET;ORAL

CALCIUM ACETATE

<u>AB</u>	HERITAGE PHARMS INC	<u>667MG</u>	<u>A202885 001</u>	Jan 22, 2015
<u>AB</u>	INVAGEN PHARMS	<u>667MG</u>	<u>A202420 001</u>	Feb 05, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-68 (of 436)

CALCIUM ACETATE

TABLET;ORAL

CALCIUM ACETATE

**AB** ! PADDOCK LLC **667MG** **A091561 001** Apr 13, 2011

ELIPHOS

**AB** CYPRESS PHARM **667MG** **A078502 001** Nov 25, 2008

CALCIUM CHLORIDE

INJECTABLE;INJECTION

CALCIUM CHLORIDE 10%

**AP** LUITPOLD PHARMS INC **100MG/ML** **A209088 001** Jul 27, 2017

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

**AP** +! HOSPIRA **100MG/ML** **N021117 001** Jan 28, 2000

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION;IRRIGATION

BSS PLUS

**AT** +! ALCON **0.154MG/ML;0.92MG/ML;0.184MG/ML;0.2MG/M  
L;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/M  
L** **N018469 001**

ENDOSOL EXTRA

**AT** +! AKORN **0.154MG/ML;0.92MG/ML;0.184MG/ML;0.2MG/M  
L;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/M  
L** **N020079 001** Nov 27, 1991

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

PRISMASOL B22GK 2/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.0  
CORP 5GM/1000ML;0.157GM/1000ML;2.21GM/1000ML  
;7.07GM/1000ML (5000ML)

PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.0  
CORP 5GM/1000ML;0.314GM/1000ML;2.21GM/1000ML  
;7.07GM/1000ML (5000ML)

PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;  
CORP 3.05GM/1000ML;0.314GM/1000ML;2.21GM/100  
0ML;7.07GM/1000ML (5000ML)

PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;  
CORP 3.05GM/1000ML;N/A/1000ML;3.09GM/1000ML;  
6.46GM/1000ML (5000ML)

PRISMASOL BGK 2/0 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML;20GM/1000ML;5.4GM/1000ML;2.0  
CORP 3GM/1000ML;0.157GM/1000ML;3.09GM/1000ML  
;6.46GM/1000ML (5000ML)

PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 5.15GM/1000ML;20GM/1000ML;5.4GM/1000ML;  
CORP 2.03GM/1000ML;0.157GM/1000ML;3.09GM/100  
0ML;6.46GM/1000ML (5000ML)

PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML;20GM/1000ML;5.4GM/1000ML;2.4  
CORP 4GM/1000ML;0.314GM/1000ML;3.09GM/1000ML  
;6.46GM/1000ML (5000ML)

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;  
CORP 3.05GM/1000ML;0.314GM/1000ML;3.09GM/1000ML  
;6.46GM/1000ML (5000ML)

PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE N/A/1000ML;N/A/1000ML;5.4GM/1000ML;2.44  
CORP GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46  
GM/1000ML (5000ML)

PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 5.15GM/1000ML;N/A/1000ML;5.4GM/1000ML;2  
0.03GM/1000ML;N/A/1000ML;3.09GM/1000ML;6  
.46GM/1000ML (5000ML)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION;INTRAPERITONEAL

DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

**AT** FRESENIUS MEDCL **25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5  
67MG/100ML;392MG/100ML** **N018883 001** Nov 30, 1984

DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

**AT** FRESENIUS MEDCL **25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5  
38MG/100ML;448MG/100ML** **N018883 004** Nov 30, 1984

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-69 (of 436)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE  
 SOLUTION; INTRAPERITONEAL

<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER</b> <u>18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N020171 001</b>	Aug 19, 1992
<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5      67MG/100ML;392MG/100ML</u>	<b>N018883 002</b>	Nov 30, 1984
<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N018883 005</b>	Nov 30, 1984
<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER</b> <u>18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N020171 002</b>	Aug 19, 1992
<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;      567MG/100ML;392MG/100ML</u>	<b>N018883 003</b>	Nov 30, 1984
<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;      538MG/100ML;448MG/100ML</u>	<b>N018883 006</b>	Nov 30, 1984
<b>AT</b>	FRESENIUS MEDCL	<b>DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER</b> <u>18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;      538MG/100ML;448MG/100ML</u>	<b>N020171 003</b>	Aug 19, 1992
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5      67MG/100ML;392MG/100ML</u>	<b>N017512 001</b>	
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5      67MG/100ML;392MG/100ML</u>	<b>N017512 003</b>	
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;      567MG/100ML;392MG/100ML</u>	<b>N017512 002</b>	
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</b> <u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N020183 001</b>	Dec 04, 1992
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5      67MG/100ML;392MG/100ML</u>	<b>N017512 007</b>	Jul 09, 1984
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5      67MG/100ML;392MG/100ML</u>	<b>N017512 008</b>	Jul 09, 1984
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;      567MG/100ML;392MG/100ML</u>	<b>N017512 009</b>	Jul 09, 1984
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</b> <u>18.3MG/100ML;1.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N017512 004</b>	
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N020163 001</b>	Dec 04, 1992
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;      538MG/100ML;448MG/100ML</u>	<b>N017512 005</b>	
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;      538MG/100ML;448MG/100ML</u>	<b>N020163 002</b>	Dec 04, 1992
<b>AT</b>	BAXTER HLTHCARE	<b>DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;      538MG/100ML;448MG/100ML</u>	<b>N020163 003</b>	Dec 04, 1992
		<b>DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</b> <u>18.3MG/100ML;2.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N020183 002</b>	Dec 04, 1992
		<b>DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</b> <u>18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N020183 003</b>	Dec 04, 1992
		<b>DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</b> <u>18.3MG/100ML;4.25GM/100ML;5.08MG/100ML;      538MG/100ML;448MG/100ML</u>	<b>N020183 004</b>	Dec 04, 1992
		<b>DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5      67MG/100ML;392MG/100ML</u>	<b>N017512 010</b>	Nov 18, 1985
		<b>DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</b> <u>25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5      38MG/100ML;448MG/100ML</u>	<b>N017512 011</b>	Nov 18, 1985

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-70 (of 436)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INTRATHECAL

ELLIOTTS B SOLUTION

+! LUKARE MEDICAL LLC 0.2MG/ML;0.8MG/ML;0.3MG/ML;0.3MG/ML;1.9 MG/ML;7.3MG/ML;0.2MG/ML N020577 001 Sep 27, 1996

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER

B BRAUN 33MG/100ML;5GM/100ML;30MG/100ML;860MG/1 00ML N020000 001 Apr 17, 1992

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP + ICU MEDICAL INC 20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML N017608 001

DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

AP B BRAUN 20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML N019634 003 Feb 24, 1988

LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML N016679 001

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;254MG/100ML;600MG/ 100ML;310MG/100ML N019367 006 Apr 05, 1985

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML N019367 004 Apr 05, 1985

AP 20MG/100ML;5GM/100ML;328MG/100ML;600MG/ 100ML;310MG/100ML N019367 005 Apr 05, 1985

AP ICU MEDICAL INC 20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML N019685 002 Oct 17, 1988

AP 20MG/100ML;5GM/100ML;328MG/100ML;600MG/ 100ML;310MG/100ML N019685 008 Oct 17, 1988

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;254MG/100ML;600MG/ 100ML;310MG/100ML N019367 007 Apr 05, 1985

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20MG/100ML;5GM/100ML;328MG/100ML;600MG/ 100ML;310MG/100ML N019367 008 Apr 05, 1985

AP ICU MEDICAL INC 20MG/100ML;5GM/100ML;328MG/100ML;600MG/ 100ML;310MG/100ML N019685 004 Oct 17, 1988

DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER

B BRAUN 10MG/100ML;2.5GM/100ML;15MG/100ML;300MG /100ML;160MG/100ML N019634 001 Feb 24, 1988

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML;5GM/100ML;105MG/100ML;600MG/ 100ML;310MG/100ML N019367 002 Apr 05, 1985

20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML N019367 003 Apr 05, 1985

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE 20MG/100ML;5GM/100ML;105MG/100ML;600MG/ 100ML;310MG/100ML N019367 001 Apr 05, 1985

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

+! HOSPIRA 16.5MG/ML;25.4MG/ML;74.6MG/ML;121MG/ML; 16.1MG/ML N018895 001 Jul 20, 1984

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

SOLUTION; IRRIGATION

BALANCED SALT

AT AKORN 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6 .4MG/ML;1.7MG/ML A075503 001 Sep 27, 2006

AT B BRAUN 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6 .4MG/ML;1.7MG/ML A091387 001 Feb 03, 2010

BSS

AT +! ALCON 0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6 .4MG/ML;1.7MG/ML N020742 001 Dec 10, 1997



38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-72 (of 436)

CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

INVOKAMET XR

+ JANSSEN PHARMS	50MG;500MG
+	50MG;1GM
+	150MG;500MG
+!	150MG;1GM

N205879 001	Sep 20, 2016
N205879 002	Sep 20, 2016
N205879 003	Sep 20, 2016
N205879 004	Sep 20, 2016

CANDESARTAN CILEXETIL

TABLET; ORAL

**ATACAND**

<b>AB</b> + ASTRAZENECA	<b>4MG</b>
<b>AB</b> +	<b>8MG</b>
<b>AB</b> +	<b>16MG</b>
<b>AB</b> +!	<b>32MG</b>

<b>N020838 001</b>	Jun 04, 1998
<b>N020838 002</b>	Jun 04, 1998
<b>N020838 003</b>	Jun 04, 1998
<b>N020838 004</b>	Jun 04, 1998

**CANDESARTAN CILEXETIL**

<b>AB</b> ALEMBIC PHARMS LTD	<b>32MG</b>
<b>AB</b> APOTEX INC	<b>4MG</b>
<b>AB</b>	<b>8MG</b>
<b>AB</b>	<b>16MG</b>
<b>AB</b>	<b>32MG</b>
<b>AB</b> MACLEODS PHARMS LTD	<b>4MG</b>
<b>AB</b>	<b>8MG</b>
<b>AB</b>	<b>16MG</b>
<b>AB</b>	<b>32MG</b>
<b>AB</b> SANDOZ	<b>4MG</b>
<b>AB</b>	<b>8MG</b>
<b>AB</b>	<b>16MG</b>
<b>AB</b>	<b>32MG</b>
<b>AB</b> ZYDUS PHARMS USA INC	<b>4MG</b>
<b>AB</b>	<b>8MG</b>
<b>AB</b>	<b>16MG</b>
<b>AB</b>	<b>32MG</b>

<b>A209119 001</b>	Jun 20, 2017
<b>A202079 001</b>	Jan 10, 2014
<b>A202079 002</b>	Jan 10, 2014
<b>A202079 003</b>	Jan 10, 2014
<b>A202079 004</b>	Jan 10, 2014
<b>A203813 001</b>	Dec 05, 2016
<b>A203813 002</b>	Dec 05, 2016
<b>A203813 003</b>	Dec 05, 2016
<b>A203813 004</b>	Dec 05, 2016
<b>A078702 001</b>	May 03, 2013
<b>A078702 002</b>	May 03, 2013
<b>A078702 003</b>	May 03, 2013
<b>A078702 004</b>	May 03, 2013
<b>A091390 001</b>	Aug 23, 2017
<b>A091390 002</b>	Aug 23, 2017
<b>A091390 003</b>	Aug 23, 2017
<b>A091390 004</b>	Aug 23, 2017

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

**ATACAND HCT**

<b>AB</b> + ASTRAZENECA	<b>16MG;12.5MG</b>
<b>AB</b> +	<b>32MG;12.5MG</b>
<b>AB</b> +!	<b>32MG;25MG</b>

<b>N021093 001</b>	Sep 05, 2000
<b>N021093 002</b>	Sep 05, 2000
<b>N021093 003</b>	May 16, 2008

**CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE**

<b>AB</b> APOTEX INC	<b>16MG;12.5MG</b>
<b>AB</b>	<b>32MG;12.5MG</b>
<b>AB</b>	<b>32MG;25MG</b>
<b>AB</b> DR REDDYS LABS LTD	<b>16MG;12.5MG</b>
<b>AB</b>	<b>32MG;12.5MG</b>
<b>AB</b>	<b>32MG;25MG</b>
<b>AB</b> MACLEODS PHARMS LTD	<b>16MG;12.5MG</b>
<b>AB</b>	<b>32MG;12.5MG</b>
<b>AB</b>	<b>32MG;25MG</b>
<b>AB</b> MYLAN PHARMS INC	<b>16MG;12.5MG</b>
<b>AB</b>	<b>32MG;12.5MG</b>
<b>AB</b>	<b>32MG;25MG</b>
<b>AB</b> ZYDUS PHARMS USA INC	<b>16MG;12.5MG</b>
<b>AB</b>	<b>32MG;12.5MG</b>
<b>AB</b>	<b>32MG;25MG</b>

<b>A202884 001</b>	Dec 04, 2012
<b>A202884 002</b>	Dec 04, 2012
<b>A202884 003</b>	Jun 03, 2013
<b>A202965 001</b>	Jun 03, 2013
<b>A202965 002</b>	Jun 03, 2013
<b>A202965 003</b>	Jun 03, 2013
<b>A204100 001</b>	Feb 27, 2015
<b>A204100 002</b>	Feb 27, 2015
<b>A204100 003</b>	Feb 27, 2015
<b>A090704 001</b>	Dec 04, 2012
<b>A090704 002</b>	Dec 04, 2012
<b>A090704 003</b>	Dec 04, 2012
<b>A203466 001</b>	Nov 27, 2017
<b>A203466 002</b>	Nov 27, 2017
<b>A203466 003</b>	Nov 27, 2017

CANGRELOR

POWDER; IV (INFUSION)

KENGREAL

+! CHIESI USA INC	50MG/VIAL
-------------------	-----------

N204958 001 Jun 22, 2015

CAPECITABINE

TABLET; ORAL

**CAPECITABINE**

<b>AB</b> ACCORD HLTHCARE	<b>150MG</b>
<b>AB</b>	<b>500MG</b>
<b>AB</b> ALKEM LABS LTD	<b>150MG</b>
<b>AB</b>	<b>500MG</b>
<b>AB</b> AMNEAL PHARMS	<b>150MG</b>
<b>AB</b>	<b>500MG</b>
<b>AB</b> MYLAN PHARMS INC	<b>150MG</b>
<b>AB</b>	<b>500MG</b>

<b>A202593 001</b>	Apr 23, 2015
<b>A202593 002</b>	Apr 23, 2015
<b>A207652 001</b>	Nov 24, 2017
<b>A207652 002</b>	Nov 24, 2017
<b>A204741 001</b>	Feb 28, 2017
<b>A204741 002</b>	Feb 28, 2017
<b>A090943 001</b>	Aug 08, 2014
<b>A090943 002</b>	Aug 08, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PREScription DRUG PRODUCT LIST**

3-73 (of 436)

## CAPECITABINE

**TABLET; ORAL**

## **CAPECITABINE**

<u><b>AB</b></u>	SHILPA MEDICARE LTD	<u><b>150MG</b></u>	<u><b>A207456</b></u>	<u><b>001</b></u>	Dec 12, 2016
<u><b>AB</b></u>		<u><b>500MG</b></u>	<u><b>A207456</b></u>	<u><b>002</b></u>	Dec 12, 2016
<u><b>AB</b></u>	TEVA PHARMS USA	<u><b>150MG</b></u>	<u><b>A091649</b></u>	<u><b>001</b></u>	Sep 16, 2013
<u><b>AB</b></u>		<u><b>500MG</b></u>	<u><b>A091649</b></u>	<u><b>002</b></u>	Sep 16, 2013
<u><b>AB</b></u>	WEST-WARD PHARMS INT	<u><b>150MG</b></u>	<u><b>A200483</b></u>	<u><b>001</b></u>	Jul 14, 2016
<u><b>AB</b></u>		<u><b>500MG</b></u>	<u><b>A200483</b></u>	<u><b>002</b></u>	Jul 14, 2016
<u><b>XELODA</b></u>					
<u><b>AB</b></u>	+ HOFFMANN LA ROCHE	<u><b>150MG</b></u>	<u><b>N020896</b></u>	<u><b>001</b></u>	Apr 30, 1998
<u><b>AB</b></u>	+!	<u><b>500MG</b></u>	<u><b>N020896</b></u>	<u><b>002</b></u>	Apr 30, 1998

## CAPREOMYCIN SULFATE

## INJECTABLE; INJECTION

## **CAPASTAT SULFATE**

<u>AP</u>	+!	AKORN	<u>EQ 1GM BASE/VIAL</u>	<u>N050095 001</u>
<u>CAPREOMYCIN SULFATE</u>				
<u>AP</u>		MYLAN LABS LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A202634 001</u> NOV 27, 2017

#### CAPSAICIN

## PATCH; TOPICAL

QUOTENZA

+ ! ACORDA

8%

N050095 001

#### CAPTOPRIL

**TABLET; ORAL**

## CAPTOPRIL

<u>AB</u>	HIKMA INTL PHARMS	<u>12 .5MG</u>	<u>A074505</u> <u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074505</u> <u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074505</u> <u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>A074505</u> <u>004</u>	Feb 13, 1996
<u>AB</u>	MYLAN PHARMS INC	<u>12 .5MG</u>	<u>A074434</u> <u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074434</u> <u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074434</u> <u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>A074434</u> <u>004</u>	Feb 13, 1996
<u>AB</u>	PRINSTON INC	<u>12 .5MG</u>	<u>A074477</u> <u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074477</u> <u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074477</u> <u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>A074477</u> <u>004</u>	Feb 13, 1996
<u>AB</u>	TEVA	<u>12 .5MG</u>	<u>A074322</u> <u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>A074322</u> <u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>A074322</u> <u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>A074322</u> <u>004</u>	Feb 13, 1996
<u>AB</u>	WATSON LABS	<u>12 .5MG</u>	<u>A074386</u> <u>001</u>	May 23, 1996
<u>AB</u>		<u>25MG</u>	<u>A074386</u> <u>002</u>	May 23, 1996
<u>AB</u>		<u>50MG</u>	<u>A074386</u> <u>003</u>	May 23, 1996
<u>AB</u>		<u>100MG</u>	<u>A074386</u> <u>004</u>	May 23, 1996
<u>AB</u>	WOCKHARDT LTD	<u>12 .5MG</u>	<u>A074532</u> <u>001</u>	Mar 28, 1997
<u>AB</u>		<u>25MG</u>	<u>A074532</u> <u>002</u>	Mar 28, 1997
<u>AB</u>		<u>50MG</u>	<u>A074532</u> <u>003</u>	Mar 28, 1997
<u>AB</u>		<u>100MG</u>	<u>A074532</u> <u>004</u>	Mar 28, 1997

## CAPTOPRIL; HYDROCHLOROTHIAZIDE

**TABLET; ORAL**

## **CAPTOPRIL AND HYDROCHLOROTHIAZIDE**

<u>AB</u>	G AND W LABS INC	<u>25MG;15MG</u>	<u>A074827_001</u>	Dec 29, 1997
<u>AB</u>		<u>25MG;25MG</u>	<u>A074827_002</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;15MG</u>	<u>A074827_004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>A074827_003</u>	Dec 29, 1997
<u>AB</u>	MYLAN	<u>25MG;15MG</u>	<u>A074896_001</u>	Dec 29, 1997
<u>AB</u>	!	<u>25MG;25MG</u>	<u>A074896_002</u>	Dec 29, 1997
<u>AB</u>	!	<u>50MG;15MG</u>	<u>A074896_004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>A074896_003</u>	Dec 29, 1997

CARBACHOT

#### SOLUTION: INTRAOCULAR

MTOSTAT

1

0.01%

N016968 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-74 (of 436)

**CARBAMAZEPINE**

CAPSULE, EXTENDED RELEASE;ORAL

**CARBAMAZEPINE**

<b>AB</b>	APOTEX INC	<u>100MG</u>	<b>A078986 001</b>	Nov 25, 2011
<b>AB</b>		<u>200MG</u>	<b>A078986 002</b>	Nov 25, 2011
<b>AB</b>		<u>300MG</u>	<b>A078986 003</b>	Nov 25, 2011
<b>AB</b>	MYLAN IRELAND LTD	<u>100MG</u>	<b>A076697 001</b>	May 20, 2011
<b>AB</b>		<u>200MG</u>	<b>A076697 002</b>	May 20, 2011
<b>AB</b>		<u>300MG</u>	<b>A076697 003</b>	May 20, 2011
<b>AB</b>	TARO	<u>100MG</u>	<b>A201106 001</b>	Jun 21, 2013
<b>AB</b>		<u>200MG</u>	<b>A201106 002</b>	Jun 21, 2013
<b>AB</b>		<u>300MG</u>	<b>A201106 003</b>	Jun 21, 2013
<b>AB</b>	TEVA PHARMS	<u>100MG</u>	<b>A078592 001</b>	Sep 20, 2012
<b>AB</b>		<u>200MG</u>	<b>A078592 002</b>	Sep 20, 2012
<b>AB</b>		<u>300MG</u>	<b>A078592 003</b>	Sep 20, 2012

**CARBATROL**

<b>AB</b>	+	SHIRE	<u>100MG</u>	<b>N020712 003</b>	Sep 30, 1997
<b>AB</b>	+		<u>200MG</u>	<b>N020712 001</b>	Sep 30, 1997
<b>AB</b>	+!		<u>300MG</u>	<b>N020712 002</b>	Sep 30, 1997
	EQUETRO				
	+	VALIDUS PHARMS	100MG	N021710 001	Dec 10, 2004
	+		200MG	N021710 002	Dec 10, 2004
	+!		300MG	N021710 003	Dec 10, 2004

SOLUTION;IV (INFUSION)

CARNEXIV

+! LUNDBECK PHARMS LLC 200MG/20ML (10MG/ML)

N206030 001 Oct 07, 2016

SUSPENSION;ORAL

**CARBAMAZEPINE**

<b>AB</b>	WOCHARDT BIO AG	<u>100MG/5ML</u>	<b>A075714 001</b>	Jun 05, 2002
<b>AB</b>	<b>TEGRETOL</b>			
<b>AB</b>	+!	NOVARTIS	<u>100MG/5ML</u>	<b>N018927 001</b> Dec 18, 1987

**TERIL**

<b>AB</b>	TARO PHARM	<u>100MG/5ML</u>	<b>A076729 001</b>	Sep 20, 2004
-----------	------------	------------------	--------------------	--------------

TABLET;ORAL

**CARBAMAZEPINE**

<b>AB</b>	APOTEX INC	<u>200MG</u>	<b>A075948 001</b>	Feb 27, 2002
<b>AB</b>	TARO	<u>200MG</u>	<b>A074649 001</b>	Oct 03, 1996
<b>AB</b>	TORRENT PHARMS	<u>200MG</u>	<b>A077272 002</b>	Dec 07, 2005

**EPITOL**

<b>AB</b>	TEVA	<u>200MG</u>	<b>A070541 001</b>	Sep 17, 1986
-----------	------	--------------	--------------------	--------------

**TEGRETOL**

<b>AB</b>	+!	NOVARTIS	<u>200MG</u>	<b>N016608 001</b>
	CARBAMAZEPINE			
	TORRENT PHARMS	100MG	A077272 001	Dec 07, 2005
		300MG	A077272 003	Dec 07, 2005
		400MG	A077272 004	Dec 07, 2005

TABLET, CHEWABLE;ORAL

**CARBAMAZEPINE**

<b>AB</b>	TARO PHARM INDS	<u>100MG</u>	<b>A075687 001</b>	Oct 24, 2000
<b>AB</b>	TORRENT PHARMS	<u>100MG</u>	<b>A075712 001</b>	Jul 05, 2001

**EPITOL**

<b>AB</b>	TEVA	<u>100MG</u>	<b>A073524 001</b>	Jul 29, 1992
<b>AB</b>	<b>TEGRETOL</b>			
<b>AB</b>	+!	NOVARTIS	<u>100MG</u>	<b>N018281 001</b>
	CARBAMAZEPINE			

!

TARO PHARM INDS

200MG

A075687 002 Jul 29, 2002

TABLET, EXTENDED RELEASE;ORAL

**CARBAMAZEPINE**

<b>AB</b>	TARO	<u>100MG</u>	<b>A078115 001</b>	Mar 31, 2009
<b>AB</b>		<u>200MG</u>	<b>A078115 002</b>	Mar 31, 2009
<b>AB</b>		<u>400MG</u>	<b>A078115 003</b>	Mar 31, 2009

**TEGRETOL-XR**

<b>AB</b>	+	NOVARTIS	<u>100MG</u>	<b>N020234 001</b>	Mar 25, 1996
<b>AB</b>	+		<u>200MG</u>	<b>N020234 002</b>	Mar 25, 1996
<b>AB</b>	+!		<u>400MG</u>	<b>N020234 003</b>	Mar 25, 1996

**CARBIDOPA**

TABLET;ORAL

**CARBIDOPA**

<b>AB</b>	ALVOGEN MALTA	<u>25MG</u>	<b>A204291 001</b>	Jan 08, 2016
<b>AB</b>	AMERIGEN PHARMS LTD	<u>25MG</u>	<b>A203261 001</b>	Mar 10, 2014
<b>AB</b>	EDENBRIDGE PHARMS	<u>25MG</u>	<b>A205304 001</b>	Feb 17, 2016
<b>AB</b>	NOVEL LABS INC	<u>25MG</u>	<b>A204763 001</b>	Oct 20, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-75 (of 436)

CARBIDOPA

TABLET;ORAL

IODOSYN

AB +! ATON 25MG N017830 001

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET;ORAL

CARBIDOPA, LEVODOPA AND ENTACAPONE

<u>AB</u>	SUN PHARMA GLOBAL	<u>25MG;200MG;100MG</u>	<u>A079085 001</u>	May 10, 2012
<u>AB</u>		<u>37.5MG;200MG;150MG</u>	<u>A079085 002</u>	May 10, 2012
<u>AB</u>	WOCKHARDT LTD	<u>12.5MG;200MG;50MG</u>	<u>A090786 001</u>	Nov 20, 2012
<u>AB</u>		<u>18.75MG;200MG;75MG</u>	<u>A090833 001</u>	Nov 20, 2012
<u>AB</u>		<u>25MG;200MG;100MG</u>	<u>A090833 002</u>	Nov 20, 2012
<u>AB</u>		<u>31.25MG;200MG;125MG</u>	<u>A090833 003</u>	Nov 20, 2012
<u>AB</u>		<u>37.5MG;200MG;150MG</u>	<u>A090833 004</u>	Nov 20, 2012
<u>AB</u>		<u>50MG;200MG;200MG</u>	<u>A090833 005</u>	Nov 20, 2012
		<u>STALEVO 100</u>		
<u>AB</u>	+ ORION PHARMA	<u>25MG;200MG;100MG</u>	<u>N021485 002</u>	Jun 11, 2003
		<u>STALEVO 125</u>		
<u>AB</u>	+ ORION PHARMA	<u>31.25MG;200MG;125MG</u>	<u>N021485 006</u>	Aug 29, 2008
		<u>STALEVO 150</u>		
<u>AB</u>	+ ORION PHARMA	<u>37.5MG;200MG;150MG</u>	<u>N021485 003</u>	Jun 11, 2003
		<u>STALEVO 200</u>		
<u>AB</u>	+! ORION PHARMA	<u>50MG;200MG;200MG</u>	<u>N021485 004</u>	Aug 02, 2007
		<u>STALEVO 50</u>		
<u>AB</u>	+! ORION PHARMA	<u>12.5MG;200MG;50MG</u>	<u>N021485 001</u>	Jun 11, 2003
		<u>STALEVO 75</u>		
<u>AB</u>	+ ORION PHARMA	<u>18.75MG;200MG;75MG</u>	<u>N021485 005</u>	Aug 29, 2008

CARBIDOPA; LEVODOPA

CAPSULE, EXTENDED RELEASE;ORAL

RYTARY

+	IMPAK LABS INC	<u>23.75MG;95MG</u>	<u>N203312 001</u>	Jan 07, 2015
+		<u>36.25MG;145MG</u>	<u>N203312 002</u>	Jan 07, 2015
+		<u>48.75MG;195MG</u>	<u>N203312 003</u>	Jan 07, 2015
+		<u>61.25MG;245MG</u>	<u>N203312 004</u>	Jan 07, 2015

SUSPENSION;ENTERAL

DUOPA

+	ABBVIE INC	<u>4.63MG/ML;20MG/ML</u>	<u>N203952 001</u>	Jan 09, 2015
---	------------	--------------------------	--------------------	--------------

TABLET;ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG;100MG</u>	<u>A074260 001</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;100MG</u>	<u>A074260 002</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;250MG</u>	<u>A074260 003</u>	Sep 03, 1993
<u>AB</u>	APOTEX INC	<u>10MG;100MG</u>	<u>A077120 001</u>	Jun 02, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A077120 002</u>	Jun 02, 2008
<u>AB</u>		<u>25MG;250MG</u>	<u>A077120 003</u>	Jun 02, 2008
<u>AB</u>	MAYNE PHARMA	<u>10MG;100MG</u>	<u>A073618 001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;100MG</u>	<u>A073589 001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;250MG</u>	<u>A073607 001</u>	Aug 28, 1992
<u>AB</u>	MYLAN	<u>10MG;100MG</u>	<u>A090324 001</u>	Sep 28, 2009
<u>AB</u>		<u>25MG;100MG</u>	<u>A090324 002</u>	Sep 28, 2009
<u>AB</u>		<u>25MG;250MG</u>	<u>A090324 003</u>	Sep 28, 2009
<u>AB</u>	SUN PHARM INDs	<u>10MG;100MG</u>	<u>A078536 001</u>	Oct 28, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A078536 002</u>	Oct 28, 2008
<u>AB</u>		<u>25MG;250MG</u>	<u>A078536 003</u>	Oct 28, 2008

SINEMET

<u>AB</u>	+ MERCK SHARP DOHME	<u>10MG;100MG</u>	<u>N017555 001</u>	
<u>AB</u>	+	<u>25MG;100MG</u>	<u>N017555 003</u>	
<u>AB</u>	+!	<u>25MG;250MG</u>	<u>N017555 002</u>	

TABLET, EXTENDED RELEASE;ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	ACCORD HLTHCARE	<u>25MG;100MG</u>	<u>A202323 001</u>	Feb 08, 2013
<u>AB</u>		<u>50MG;200MG</u>	<u>A202323 002</u>	Feb 08, 2013
<u>AB</u>	APOTEX	<u>25MG;100MG</u>	<u>A076212 001</u>	Jun 16, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>A076212 002</u>	Jun 16, 2004
<u>AB</u>	IMPAK LABS	<u>25MG;100MG</u>	<u>A076521 001</u>	May 14, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>A076521 002</u>	May 14, 2004
<u>AB</u>	MYLAN	<u>25MG;100MG</u>	<u>A075091 002</u>	Apr 21, 2000
<u>AB</u>		<u>50MG;200MG</u>	<u>A075091 001</u>	Sep 30, 1999
<u>AB</u>	SUN PHARM INDs	<u>25MG;100MG</u>	<u>A077828 001</u>	Aug 23, 2007
<u>AB</u>		<u>50MG;200MG</u>	<u>A077828 002</u>	Aug 23, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-76 (of 436)

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE;ORAL

SINemet CR

<u>AB</u>	+	MERCK SHARP DOHME	<u>25MG;100MG</u>	<u>N019856 002</u>	Dec 24, 1992
<u>AB</u>	!		<u>50MG;200MG</u>	<u>N019856 001</u>	May 30, 1991

TABLET, ORALLY DISINTEGRATING;ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	MYLAN	<u>10MG;100MG</u>	<u>A078893 001</u>	Sep 18, 2008
<u>AB</u>		<u>25MG;100MG</u>	<u>A078893 002</u>	Sep 18, 2008
<u>AB</u>	!	<u>25MG;250MG</u>	<u>A078893 003</u>	Sep 18, 2008
<u>AB</u>	SUN PHARMA GLOBAL	<u>10MG;100MG</u>	<u>A078690 001</u>	Jul 31, 2009
<u>AB</u>		<u>25MG;100MG</u>	<u>A078690 002</u>	Jul 31, 2009
<u>AB</u>		<u>25MG;250MG</u>	<u>A078690 003</u>	Jul 31, 2009

CARBINOXAMINE MALEATE

SOLUTION;ORAL

CARBINOXAMINE MALEATE

<u>AA</u>	CYPRESS PHARM	<u>4MG/5ML</u>	<u>A090418 001</u>	May 04, 2010
<u>AA</u>	!	<u>4MG/5ML</u>	<u>A040458 001</u>	Apr 25, 2003
<u>AA</u>	VINTAGE PHARMS	<u>4MG/5ML</u>	<u>A040814 001</u>	Feb 26, 2008

SUSPENSION, EXTENDED RELEASE;ORAL

KARBINAL ER

+! TRIS PHARMA INC

4MG/5ML

N022556 001 Mar 28, 2013

TABLET;ORAL

CARBINOXAMINE MALEATE

<u>AA</u>	CYPRESS PHARM	<u>4MG</u>	<u>A090417 001</u>	Aug 23, 2010
<u>AA</u>	INVAGEN PHARMS	<u>4MG</u>	<u>A090435 001</u>	Apr 15, 2010
<u>AA</u>	!	<u>4MG</u>	<u>A040442 001</u>	Mar 19, 2003
<u>AA</u>	MISSION PHARMACAL CO	<u>4MG</u>	<u>A090756 001</u>	May 27, 2011
<u>AA</u>	VINTAGE PHARMS	<u>4MG</u>	<u>A040639 002</u>	May 30, 2008
	MIKART INC	6MG	A207484 001	May 31, 2016

CARBOPLATIN

INJECTABLE;IV (INFUSION)

CARBOPLATIN

<u>AP</u>	ACCORD HLTHCARE	<u>50MG/5ML (10MG/ML)</u>	<u>A206775 001</u>	Feb 09, 2017
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A206775 002</u>	Feb 09, 2017
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A206775 003</u>	Feb 09, 2017
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A206775 004</u>	Feb 09, 2017
<u>AP</u>	AKORN	<u>50MG/5ML (10MG/ML)</u>	<u>A090475 001</u>	Jul 29, 2009
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A090475 002</u>	Jul 29, 2009
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A090475 003</u>	Jul 29, 2009
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A091268 002</u>	Jul 28, 2010
<u>AP</u>	CIPLA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A077861 001</u>	Jan 18, 2007
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077861 002</u>	Jan 18, 2007
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077861 003</u>	Jan 18, 2007
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077861 004</u>	Jan 18, 2007
<u>AP</u>	FRESENIUS KABI ONCOL	<u>50MG/5ML (10MG/ML)</u>	<u>A077432 001</u>	Sep 29, 2006
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077432 002</u>	Sep 29, 2006
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077432 003</u>	Sep 29, 2006
<u>AP</u>	FRESENIUS KABI USA	<u>450MG/45ML (10MG/ML)</u>	<u>A077247 003</u>	Oct 21, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077266 003</u>	Feb 15, 2006
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077266 004</u>	Feb 15, 2006
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A207324 001</u>	Feb 15, 2017
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A207324 002</u>	Feb 15, 2017
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A207324 003</u>	Feb 15, 2017
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A207324 004</u>	Feb 15, 2017
<u>AP</u>	HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A076517 001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A076517 002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A076517 003</u>	Oct 14, 2004
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077059 001</u>	Nov 23, 2004
<u>AP</u>	INGENUS PHARMS LLC	<u>50MG/5ML (10MG/ML)</u>	<u>A208487 001</u>	Apr 26, 2017
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A208487 002</u>	Apr 26, 2017
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A208487 003</u>	Apr 26, 2017
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A208487 004</u>	Apr 26, 2017
<u>AP</u>	MYLAN INSTITUTIONAL	<u>50MG/5ML (10MG/ML)</u>	<u>A077998 001</u>	Apr 24, 2007
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077998 002</u>	Apr 24, 2007
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077998 003</u>	Apr 24, 2007
<u>AP</u>	MYLAN LABS LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A091063 001</u>	Nov 09, 2011
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A091063 002</u>	Nov 09, 2011
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A091063 003</u>	Nov 09, 2011
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A091063 004</u>	Nov 09, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-77 (of 436)

CARBOPLATIN

INJECTABLE; IV (INFUSION)

CARBOPLATIN

<u>AP</u>	NANJING KING-FRIEND	<u>50MG/5ML (10MG/ML)</u>	<u>A077096 001</u>	Jun 14, 2005
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077096 002</u>	Jun 14, 2005
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077096 003</u>	Jun 14, 2005
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077096 004</u>	Jun 03, 2013
<u>AP</u>	! PHARMACHEMIE BV	<u>50MG/5ML (10MG/ML)</u>	<u>A077269 001</u>	Oct 14, 2004
<u>AP</u>	!	<u>150MG/15ML (10MG/ML)</u>	<u>A077269 002</u>	Oct 14, 2004
<u>AP</u>	!	<u>450MG/45ML (10MG/ML)</u>	<u>A077269 003</u>	Oct 14, 2004
<u>AP</u>	!	<u>600MG/60ML (10MG/ML)</u>	<u>A077269 004</u>	Dec 28, 2007
<u>AP</u>	PLIVA LACHEMA	<u>50MG/5ML (10MG/ML)</u>	<u>A078631 001</u>	Dec 02, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A078631 002</u>	Dec 02, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078631 003</u>	Dec 02, 2008
<u>AP</u>	SANDOZ INC	<u>50MG/5ML (10MG/ML)</u>	<u>A078631 004</u>	Dec 02, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A078280 001</u>	May 08, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A078280 002</u>	May 08, 2008
<u>AP</u>	SANJA PHARMS CO	<u>50MG/5ML (10MG/ML)</u>	<u>A078280 003</u>	May 08, 2008
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A205487 001</u>	Mar 28, 2016
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A205487 002</u>	Mar 28, 2016
<u>AP</u>	SUN PHARMA GLOBAL	<u>50MG/5ML (10MG/ML)</u>	<u>A205487 003</u>	Mar 28, 2016
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077926 001</u>	Sep 19, 2008
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077926 002</u>	Sep 19, 2008
<u>AP</u>	! TEVA PHARMS USA	<u>50MG/5ML (10MG/ML)</u>	<u>A077926 003</u>	Sep 19, 2008
<u>AP</u>	!	<u>150MG/15ML (10MG/ML)</u>	<u>A077139 001</u>	Sep 21, 2005
<u>AP</u>	!	<u>450MG/45ML (10MG/ML)</u>	<u>A077139 002</u>	Sep 21, 2005
<u>AP</u>	!	<u>600MG/60ML (10MG/ML)</u>	<u>A077139 003</u>	Sep 21, 2005
<u>AP</u>	WEST-WARD PHARMS INT	<u>50MG/5ML (10MG/ML)</u>	<u>A077139 004</u>	Sep 21, 2005
<u>AP</u>		<u>150MG/15ML (10MG/ML)</u>	<u>A077244 001</u>	Oct 15, 2004
<u>AP</u>		<u>450MG/45ML (10MG/ML)</u>	<u>A077244 002</u>	Oct 15, 2004
<u>AP</u>		<u>600MG/60ML (10MG/ML)</u>	<u>A077244 003</u>	Oct 15, 2004
	! MYLAN LABS LTD	1GM/100ML (10MG/ML)	<u>A077244 004</u>	Jan 20, 2006
			A091478 001	Nov 23, 2011

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION  
 HEMABATE

+!	PHARMACIA AND UPJOHN	EQ 0.25MG BASE/ML	N017989 001
----	----------------------	-------------------	-------------

CARFILZOMIB

POWDER; INTRAVENOUS  
 KYPROLIS

+	ONYX THERAP	30MG/VIAL	N202714 002	Jun 03, 2016
+		60MG/VIAL	N202714 001	Jul 20, 2012

CARGLUMIC ACID

TABLET; ORAL  
 CARBAGLU

+!	ORPHAN EUROPE	200MG	N022562 001	Mar 18, 2010
----	---------------	-------	-------------	--------------

CARIPRAZINE HYDROCHLORIDE

CAPSULE; ORAL  
 VRAYLAR

+	FOREST RES INST INC	EQ 1.5MG BASE	N204370 001	Sep 17, 2015
+		EQ 3MG BASE	N204370 002	Sep 17, 2015
+		EQ 4.5MG BASE	N204370 003	Sep 17, 2015
+		EQ 6MG BASE	N204370 004	Sep 17, 2015

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

<u>AA</u>	ACCELRX LABS	<u>350MG</u>	<u>A040576 001</u>	Jun 07, 2005
<u>AA</u>	AUROBINDO PHARMA	<u>350MG</u>	<u>A040792 001</u>	Aug 06, 2009
<u>AA</u>	HIKMA INTL PHARMS	<u>350MG</u>	<u>A040124 001</u>	Jan 24, 1996
<u>AA</u>	INGENUS PHARMS NJ	<u>350MG</u>	<u>A040823 001</u>	Oct 22, 2008
<u>AA</u>	NATCO PHARMA LTD	<u>350MG</u>	<u>A090988 001</u>	Oct 28, 2014
<u>AA</u>	ORIENT PHARMA CO LTD	<u>350MG</u>	<u>A205085 001</u>	Oct 28, 2014
<u>AA</u>	SCIEGEN PHARMS INC	<u>350MG</u>	<u>A203374 001</u>	Jan 27, 2014
<u>AA</u>	STRIDES PHARMA	<u>350MG</u>	<u>A205513 002</u>	Nov 12, 2015
<u>AA</u>	SUN PHARM INDUSTRIES	<u>350MG</u>	<u>A089346 001</u>	Oct 17, 1991
<u>AA</u>	VINTAGE PHARMS	<u>350MG</u>	<u>A040245 001</u>	Sep 08, 1997
<u>AA</u>	WATSON LABS	<u>350MG</u>	<u>A087499 001</u>	Apr 20, 1982

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-78 (of 436)

CARISOPRODOL

TABLET;ORAL

CARISOPRODOL

<u>AA</u>	WILSHIRE PHARMS INC	<u>350MG</u>	<u>A205126 002</u>	Jul 08, 2015
		<u>SOMA</u>		
<u>AA</u>	+ MYLAN SPECIALITY LP	<u>350MG</u>	<u>N011792 001</u>	
		<u>CARISOPRODOL</u>		
<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>	<u>A040792 002</u>	Nov 08, 2016
<u>AB</u>	NOSTRUM LABS INC	<u>250MG</u>	<u>A207237 001</u>	May 11, 2017
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A205513 001</u>	Nov 12, 2015
<u>AB</u>	WILSHIRE PHARMS INC	<u>250MG</u>	<u>A205126 001</u>	Jul 08, 2015
		<u>SOMA</u>		
<u>AB</u>	+! MYLAN SPECIALITY LP	<u>250MG</u>	<u>N011792 004</u>	Sep 13, 2007

CARMUSTINE

IMPLANT; INTRACRANIAL  
 GLIADEL

+! ARBOR PHARMS LLC 7.7MG

INJECTABLE; INJECTION

BICNU

+! EMCURE PHARMS LTD 100MG/VIAL

N020637 001 Sep 23, 1996

N017422 001

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>A075546 001</u>	Jan 20, 2000
<u>AT</u>	! SANDOZ INC	<u>1%</u>	<u>A075476 001</u>	Jan 03, 2000

CARVEDILOL

TABLET;ORAL

CARVEDILOL

<u>AB</u>	APOTEK INC	<u>3.125MG</u>	<u>A078165 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078165 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078165 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078165 004</u>	Sep 05, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>3.125MG</u>	<u>A078332 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078332 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078332 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078332 004</u>	Sep 05, 2007
<u>AB</u>	BEXIMCO USA	<u>3.125MG</u>	<u>A078384 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078384 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078384 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078384 004</u>	Sep 05, 2007
<u>AB</u>	CIPLA LTD	<u>3.125MG</u>	<u>A077474 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077474 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077474 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077474 004</u>	Sep 05, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>3.125MG</u>	<u>A076649 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076649 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076649 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A076649 004</u>	Sep 05, 2007
<u>AB</u>	GLENMARK GENERICS	<u>3.125MG</u>	<u>A078251 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078251 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078251 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078251 004</u>	Sep 05, 2007
<u>AB</u>	LUPIN	<u>3.125MG</u>	<u>A078217 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078217 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078217 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078217 004</u>	Sep 05, 2007
<u>AB</u>	MYLAN	<u>3.125MG</u>	<u>A077316 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077316 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077316 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077316 004</u>	Sep 05, 2007
<u>AB</u>	SANDOZ	<u>3.125MG</u>	<u>A078227 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A078227 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A078227 003</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A078227 004</u>	Sep 05, 2007
<u>AB</u>	SUN PHARM IND'S INC	<u>3.125MG</u>	<u>A077346 004</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A077346 001</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A077346 002</u>	Sep 05, 2007
<u>AB</u>		<u>25MG</u>	<u>A077346 003</u>	Sep 05, 2007
<u>AB</u>	SUN PHARM IND'S LTD	<u>3.125MG</u>	<u>A076989 001</u>	Sep 05, 2007
<u>AB</u>		<u>6.25MG</u>	<u>A076989 002</u>	Sep 05, 2007
<u>AB</u>		<u>12.5MG</u>	<u>A076989 003</u>	Sep 05, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-79 (of 436)

CARVEDILOL

TABLET;ORAL

CARVEDILOL

<u>AB</u>		<u>25MG</u>	<u>A076989 004</u>	Sep 05, 2007	
<u>AB</u>	TARO	<u>3.125MG</u>	<u>A077780 001</u>	Sep 05, 2007	
<u>AB</u>		<u>6.25MG</u>	<u>A077780 002</u>	Sep 05, 2007	
<u>AB</u>		<u>12.5MG</u>	<u>A077780 003</u>	Sep 05, 2007	
<u>AB</u>		<u>25MG</u>	<u>A077780 004</u>	Sep 05, 2007	
<u>AB</u>	TEVA	<u>3.125MG</u>	<u>A076373 001</u>	Sep 05, 2007	
<u>AB</u>		<u>6.25MG</u>	<u>A076373 002</u>	Sep 05, 2007	
<u>AB</u>		<u>12.5MG</u>	<u>A076373 003</u>	Sep 05, 2007	
<u>AB</u>		<u>25MG</u>	<u>A076373 004</u>	Sep 05, 2007	
<u>AB</u>	ZYDUS PHARMS USA INC	<u>3.125MG</u>	<u>A077614 004</u>	Sep 05, 2007	
<u>AB</u>		<u>6.25MG</u>	<u>A077614 001</u>	Sep 05, 2007	
<u>AB</u>		<u>12.5MG</u>	<u>A077614 002</u>	Sep 05, 2007	
<u>AB</u>		<u>25MG</u>	<u>A077614 003</u>	Sep 05, 2007	
<u>COREG</u>					
<u>AB</u>	+	SMITHKLINE BEECHAM	<u>3.125MG</u>	<u>N020297 004</u>	May 29, 1997
<u>AB</u>	+		<u>6.25MG</u>	<u>N020297 003</u>	Sep 14, 1995
<u>AB</u>	+!		<u>12.5MG</u>	<u>N020297 002</u>	Sep 14, 1995
<u>AB</u>	+		<u>25MG</u>	<u>N020297 001</u>	Sep 14, 1995

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE;ORAL

CARVEDILOL PHOSPHATE

<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A090132 001</u>	Oct 25, 2017	
<u>AB</u>		<u>20MG</u>	<u>A090132 002</u>	Oct 25, 2017	
<u>AB</u>		<u>40MG</u>	<u>A090132 003</u>	Oct 25, 2017	
<u>AB</u>		<u>80MG</u>	<u>A090132 004</u>	Oct 25, 2017	
<u>COREG CR</u>					
<u>AB</u>	+	SMITHKLINE BEECHAM	<u>10MG</u>	<u>N022012 001</u>	Oct 20, 2006
<u>AB</u>	+		<u>20MG</u>	<u>N022012 002</u>	Oct 20, 2006
<u>AB</u>	+!		<u>40MG</u>	<u>N022012 003</u>	Oct 20, 2006
<u>AB</u>	+		<u>80MG</u>	<u>N022012 004</u>	Oct 20, 2006

CASPOFUNGIN ACETATE

POWDER;IV (INFUSION)

CANCIDAS

<u>AP</u>	++ MERCK	<u>50MG/VIAL</u>	<u>N021227 001</u>	Jan 26, 2001
<u>AP</u>	++	<u>70MG/VIAL</u>	<u>N021227 002</u>	Jan 26, 2001
<u>CASPOFUNGIN ACETATE</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>N206110 001</u>	Dec 30, 2016
<u>AP</u>		<u>70MG/VIAL</u>	<u>N206110 002</u>	Dec 30, 2016
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/VIAL</u>	<u>A207092 001</u>	Sep 29, 2017
<u>AP</u>		<u>70MG/VIAL</u>	<u>A207092 002</u>	Sep 29, 2017
<u>AP</u>	MYLAN LABS LTD	<u>50MG/VIAL</u>	<u>A207650 001</u>	Sep 29, 2017
<u>AP</u>		<u>70MG/VIAL</u>	<u>A207650 002</u>	Sep 29, 2017

CEFACLOR

CAPSULE;ORAL

CEFACLOR

<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>A065350 001</u>	Apr 03, 2007
<u>AB</u>	!	<u>EQ 500MG BASE</u>	<u>A065350 002</u>	Apr 03, 2007
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 250MG BASE</u>	<u>A065146 001</u>	Jan 22, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065146 002</u>	Jan 22, 2004
FOR SUSPENSION;ORAL				
	YUNG SHIN PHARM	EQ 125MG BASE/5ML	A065412 001	Feb 17, 2012
		EQ 187MG BASE/5ML	A065412 002	Feb 17, 2012
		EQ 250MG BASE/5ML	A065412 003	Feb 17, 2012
	!	EQ 375MG BASE/5ML	A065412 004	Feb 17, 2012
TABLET, EXTENDED RELEASE;ORAL				
	CEFACLOR			
	TEVA	EQ 375MG BASE	A065058 001	Sep 04, 2002
	!	EQ 500MG BASE	A065058 002	Sep 04, 2002

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE;ORAL

CEFADROXIL

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A065352 001</u>	Jan 25, 2007
<u>AB</u>	CSPC OUYI PHARM CO	<u>EQ 500MG BASE</u>	<u>A205072 001</u>	Jul 28, 2017
<u>AB</u>	HIKMA	<u>EQ 500MG BASE</u>	<u>A065311 001</u>	Feb 07, 2006
<u>AB</u>	LUPIN	<u>EQ 500MG BASE</u>	<u>A065392 001</u>	May 29, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-80 (of 436)

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

<u>AB</u>	ORCHID HLTHCARE	<u>EQ 500MG BASE</u>	<u>A065309 001</u>	Sep 18, 2006
<u>AB</u> !	TEVA PHARMS	<u>EQ 500MG BASE</u>	<u>A065282 001</u>	Jan 20, 2006

FOR SUSPENSION; ORAL

CEFADROXIL

<u>AB</u>	AUROBINDO	<u>EQ 250MG BASE/5ML</u>	<u>A065349 001</u>	Apr 25, 2013
<u>AB</u>	HIKMA PHARMS	<u>EQ 250MG BASE/5ML</u>	<u>A065349 002</u>	Apr 25, 2013
<u>AB</u>	LUPIN	<u>EQ 250MG BASE/5ML</u>	<u>A091036 001</u>	Nov 28, 2012
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE/5ML</u>	<u>A091036 002</u>	Nov 28, 2012
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE/5ML</u>	<u>A065396 001</u>	Feb 21, 2008
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE/5ML</u>	<u>A065396 002</u>	Feb 21, 2008
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 500MG BASE/5ML</u>	<u>A065307 002</u>	Oct 16, 2006
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 500MG BASE/5ML</u>	<u>A065307 003</u>	Oct 16, 2006

TABLET; ORAL

CEFADROXIL

<u>AB</u>	HIKMA	<u>EQ 1GM BASE</u>	<u>A065260 001</u>	Mar 30, 2006
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 1GM BASE</u>	<u>A065301 001</u>	Sep 18, 2006
!	TEVA PHARMS	EQ 1GM BASE	A062774 001	Apr 08, 1987

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A065303 001</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065303 002</u>	Oct 22, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065306 001</u>	Oct 22, 2008
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 500MG BASE/VIAL</u>	<u>A065047 001</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065047 002</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065143 001</u>	Oct 18, 2004
<u>AP</u>	! HOSPIRA INC	<u>EQ 500MG BASE/VIAL</u>	<u>A065226 001</u>	Apr 21, 2005
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A065226 002</u>	Apr 21, 2005
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A065244 001</u>	Aug 12, 2005
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A201654 001</u>	Feb 03, 2016
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A065247 001</u>	Aug 12, 2005
<u>AP</u>	QILU PHARM CO LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A203661 001</u>	Dec 28, 2015
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A062831 001</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A062831 002</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065345 001</u>	May 09, 2007
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A062831 003</u>	Sep 25, 1992

KEFZOL

<u>AP</u>	ACS DOBFAR	<u>EQ 500MG BASE/VIAL</u>	<u>A061773 002</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A061773 003</u>	
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A061773 004</u>	
	ANCEF IN PLASTIC CONTAINER			
!	BAXTER HLTHCARE	EQ 10MG BASE/ML	A063002 001	Mar 28, 1991
!		EQ 20MG BASE/ML	A063002 002	Mar 28, 1991
	CEFAZOLIN AND DEXTROSE			
+	! B BRAUN	EQ 1GM BASE/VIAL	N050779 002	Jul 27, 2000
+		EQ 2GM BASE/VIAL	N050779 003	Jan 13, 2012
	CEFAZOLIN SODIUM			
!	ACS DOBFAR	EQ 20GM BASE/VIAL	A065306 002	Aug 18, 2014
!	SAMSON MEDCL	EQ 100GM BASE/VIAL	A065141 001	Nov 29, 2006
!		EQ 300GM BASE/VIAL	A065141 002	Nov 29, 2006

SOLUTION; INTRAVENOUS

CEFAZOLIN IN PLASTIC CONTAINER

BAXTER HLTHCARE EQ 2GM BASE/100ML (EQ 20MG BASE/ML)

N207131 001 Aug 07, 2015

CEFDINIR

CAPSULE; ORAL

CEFDINIR

<u>AB</u>	AUROBINDO PHARMA	<u>300MG</u>	<u>A065434 001</u>	Jan 07, 2008
<u>AB</u>	LUPIN	<u>300MG</u>	<u>A065264 001</u>	May 19, 2006
<u>AB</u>	ORCHID HLTHCARE	<u>300MG</u>	<u>A065418 001</u>	Jul 18, 2007
<u>AB</u>	! SANDOZ	<u>300MG</u>	<u>A065330 001</u>	Apr 06, 2007
<u>AB</u>	TEVA PHARMS	<u>300MG</u>	<u>A065368 001</u>	May 09, 2007

FOR SUSPENSION; ORAL

CEFDINIR

<u>AB</u>	AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065473 001</u>	Dec 14, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065473 002</u>	Dec 14, 2007
<u>AB</u>	LUPIN	<u>125MG/5ML</u>	<u>A065259 001</u>	May 31, 2006
<u>AB</u>		<u>250MG/5ML</u>	<u>A065259 002</u>	May 07, 2007
<u>AB</u>	ORCHID HLTHCARE	<u>125MG/5ML</u>	<u>A065429 001</u>	Jul 18, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-81 (of 436)

**CEFDINIR**

FOR SUSPENSION;ORAL

**CEFDINIR**

<b><u>AB</u></b>		<b><u>250MG/5ML</u></b>	
<b><u>AB</u></b>	SANDOZ	<b><u>125MG/5ML</u></b>	
<b><u>AB</u></b>	!	<b><u>250MG/5ML</u></b>	
<b><u>AB</u></b>	TEVA PHARMS	<b><u>125MG/5ML</u></b>	
<b><u>AB</u></b>		<b><u>250MG/5ML</u></b>	

<b><u>A065429</u></b>	<b><u>002</u></b>	Jul 18, 2007
<b><u>A065337</u></b>	<b><u>001</u></b>	Apr 06, 2007
<b><u>A065337</u></b>	<b><u>002</u></b>	Apr 06, 2007
<b><u>A065332</u></b>	<b><u>001</u></b>	May 04, 2007
<b><u>A065332</u></b>	<b><u>002</u></b>	May 04, 2007

**CEFEPIME HYDROCHLORIDE**

INJECTABLE; INJECTION

**CEFEPIME HYDROCHLORIDE**

<b><u>AP</u></b>	ACS DOBFAR	<b><u>EQ 1GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 2GM BASE/VIAL</u></b>	
<b><u>AP</u></b>	HOSPIRA INC	<b><u>EQ 500MG BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 1GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 1GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 2GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 2GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 500MG BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 1GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 2GM BASE/VIAL</u></b>	
<b><u>AP</u></b>	SAGENT PHARMS	<b><u>EQ 1GM BASE/VIAL</u></b>	
<b><u>AP</u></b>		<b><u>EQ 2GM BASE/VIAL</u></b>	

<b><u>A065441</u></b>	<b><u>001</u></b>	Mar 20, 2008
<b><u>A065441</u></b>	<b><u>002</u></b>	Mar 20, 2008
<b><u>A065369</u></b>	<b><u>001</u></b>	Jun 18, 2007
<b><u>A065369</u></b>	<b><u>002</u></b>	Jun 18, 2007
<b><u>A202268</u></b>	<b><u>001</u></b>	Jul 30, 2012
<b><u>A065369</u></b>	<b><u>003</u></b>	Jun 18, 2007
<b><u>A202268</u></b>	<b><u>002</u></b>	Jul 30, 2012
<b><u>A203704</u></b>	<b><u>001</u></b>	Feb 01, 2016
<b><u>A203704</u></b>	<b><u>002</u></b>	Feb 01, 2016
<b><u>A203704</u></b>	<b><u>003</u></b>	Feb 01, 2016
<b><u>A091048</u></b>	<b><u>001</u></b>	Jan 04, 2017
<b><u>A091048</u></b>	<b><u>002</u></b>	Jan 04, 2017

**MAXIPIME**

<b><u>AP</u></b>	+! HOSPIRA INC	<b><u>EQ 500MG BASE/VIAL</u></b>	
<b><u>AP</u></b>	+!	<b><u>EQ 1GM BASE/VIAL</u></b>	
<b><u>AP</u></b>	+!	<b><u>EQ 2GM BASE/VIAL</u></b>	

<b><u>N050679</u></b>	<b><u>001</u></b>	Jan 18, 1996
<b><u>N050679</u></b>	<b><u>002</u></b>	Jan 18, 1996
<b><u>N050679</u></b>	<b><u>003</u></b>	Jan 18, 1996

CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER	
B BRAUN	EQ 1GM BASE/VIAL
	EQ 2GM BASE/VIAL

N050821	001	May 06, 2010
N050821	002	May 06, 2010

CEFEPIME IN PLASTIC CONTAINER

+! BAXTER HLTHCARE	EQ 1GM BASE/50ML (EQ 20MG BASE/ML)
+!	EQ 2GM BASE/100ML (EQ 20MG BASE/ML)

N050817	001	Aug 05, 2008
N050817	002	Aug 05, 2008

**CEFIXIME**

CAPSULE;ORAL

SUPRAX	
+! LUPIN LTD	400MG

N203195	001	Jun 01, 2012
---------	-----	--------------

**CEFIXIME**

<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>100MG/5ML</u></b>	
<b><u>AB</u></b>		<b><u>200MG/5ML</u></b>	
<b><u>AB</u></b>	BELCHER PHARMS LLC	<b><u>100MG/5ML</u></b>	
<b><u>AB</u></b>		<b><u>200MG/5ML</u></b>	
<b><u>AB</u></b>		<b><u>500MG/5ML</u></b>	
<b><u>AB</u></b>	SANDOZ INC	<b><u>100MG/5ML</u></b>	
<b><u>AB</u></b>		<b><u>200MG/5ML</u></b>	

<b><u>A204835</u></b>	<b><u>001</u></b>	Apr 14, 2015
<b><u>A204835</u></b>	<b><u>002</u></b>	Apr 14, 2015
<b><u>A206938</u></b>	<b><u>001</u></b>	Feb 06, 2017
<b><u>A206938</u></b>	<b><u>002</u></b>	Feb 06, 2017
<b><u>A206939</u></b>	<b><u>001</u></b>	Feb 06, 2017
<b><u>A206144</u></b>	<b><u>001</u></b>	Nov 17, 2017
<b><u>A206144</u></b>	<b><u>002</u></b>	Nov 17, 2017

**SUPRAX**

<b><u>AB</u></b>	LUPIN LTD	<b><u>500MG/5ML</u></b>	
<b><u>AB</u></b>	LUPIN PHARMS	<b><u>100MG/5ML</u></b>	
<b><u>AB</u></b>		<b><u>200MG/5ML</u></b>	

<b><u>N202091</u></b>	<b><u>001</u></b>	Feb 20, 2013
<b><u>A065129</u></b>	<b><u>001</u></b>	Feb 23, 2004
<b><u>A065355</u></b>	<b><u>001</u></b>	Apr 10, 2007

TABLET;ORAL

SUPRAX	
! LUPIN PHARMS	400MG
TABLET, CHEWABLE;ORAL	
SUPRAX	
LUPIN LTD	100MG
	150MG
!	200MG

A065380	001	Oct 25, 2010
A065380	002	Oct 25, 2010
A065380	003	Oct 25, 2010

**CEFOTAXIME SODIUM**

INJECTABLE; INJECTION

<b><u>CEFOTAXIME</u></b>		
<b><u>AP</u></b>	HIKMA	<b><u>EQ 500MG BASE/VIAL</u></b>
<b><u>AP</u></b>		<b><u>EQ 1GM BASE/VIAL</u></b>
<b><u>AP</u></b>		<b><u>EQ 2GM BASE/VIAL</u></b>
<b><u>AP</u></b>		<b><u>EQ 10GM BASE/VIAL</u></b>
<b><u>AP</u></b>	WOCKHARDT	<b><u>EQ 1GM BASE/VIAL</u></b>

<b><u>A065072</u></b>	<b><u>001</u></b>	Nov 20, 2002
<b><u>A065072</u></b>	<b><u>002</u></b>	Nov 20, 2002
<b><u>A065072</u></b>	<b><u>003</u></b>	Nov 20, 2002
<b><u>A065071</u></b>	<b><u>001</u></b>	Nov 20, 2002
<b><u>A065197</u></b>	<b><u>001</u></b>	Aug 29, 2006

**CEFOTAXIME SODIUM**

<b><u>AP</u></b>	HOSPIRA INC	<b><u>EQ 500MG BASE/VIAL</u></b>
<b><u>AP</u></b>		<b><u>EQ 1GM BASE/VIAL</u></b>
<b><u>AP</u></b>		<b><u>EQ 1GM BASE/VIAL</u></b>

<b><u>A065290</u></b>	<b><u>001</u></b>	Aug 11, 2006
<b><u>A065290</u></b>	<b><u>002</u></b>	Aug 11, 2006
<b><u>A065293</u></b>	<b><u>001</u></b>	Aug 10, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-82 (of 436)

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME SODIUM

<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A203132 001</u>	Feb 19, 2016
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065290 003</u>	Aug 11, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065293 002</u>	Aug 10, 2006
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A203132 002</u>	Feb 19, 2016
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A065292 001</u>	Aug 10, 2006
<u>AP</u>	LUPIN	<u>EQ 500MG BASE/VIAL</u>	<u>A065124 001</u>	Sep 24, 2003
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065124 002</u>	Sep 24, 2003
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065124 003</u>	Sep 24, 2003
<u>AP</u>	WOCKHARDT	<u>EQ 500MG BASE/VIAL</u>	<u>A065197 002</u>	Jun 20, 2008
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A065197 003</u>	Jun 20, 2008

CIAFORAN

<u>AP</u>	+!	US PHARM HOLDINGS	<u>EQ 500MG BASE/VIAL</u>	<u>N050547 001</u>
<u>AP</u>	+!		<u>EQ 1GM BASE/VIAL</u>	<u>N050547 002</u>
<u>AP</u>	+!		<u>EQ 2GM BASE/VIAL</u>	<u>N050547 003</u>
<u>AP</u>	+!		<u>EQ 10GM BASE/VIAL</u>	<u>N050547 004</u> Dec 29, 1983

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

<u>AP</u>	+	TELIGENT	<u>EQ 1GM BASE/VIAL</u>	<u>N050588 001</u> Dec 27, 1985
<u>AP</u>	+		<u>EQ 2GM BASE/VIAL</u>	<u>N050588 002</u> Dec 27, 1985

CEFOTETAN

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 1GM BASE/VIAL</u>	<u>A065374 001</u> Aug 09, 2007
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065374 002</u> Aug 09, 2007
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A065375 001</u> Aug 09, 2007
<u>AP</u>		HIKMA FARMACEUTICA	<u>EQ 1GM BASE/VIAL</u>	<u>A091031 001</u> Oct 26, 2011
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A091031 002</u> Oct 26, 2011
<u>AP</u>		WEST-WARD PHARM CORP	<u>EQ 10GM BASE/VIAL</u>	<u>A091030 001</u> Oct 26, 2011

CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER

+!	B BRAUN	EQ 1GM BASE/VIAL	N065430 001	Aug 09, 2007
+!		EQ 2GM BASE/VIAL	N065430 002	Aug 09, 2007

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

<u>AP</u>	!	ACS DOBFAR	<u>EQ 1GM BASE/VIAL</u>	<u>A065414 001</u> Jun 12, 2009
<u>AP</u>	!		<u>EQ 2GM BASE/VIAL</u>	<u>A065414 002</u> Jun 12, 2009
<u>AP</u>	!		<u>EQ 10GM BASE/VIAL</u>	<u>A065415 001</u> May 19, 2010
<u>AP</u>		HIKMA FARMACEUTICA	<u>EQ 1GM BASE/VIAL</u>	<u>A065238 001</u> Mar 12, 2010
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065238 002</u> Mar 12, 2010
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065239 001</u> Mar 02, 2010
<u>AP</u>		HOSPIRA INC	<u>EQ 1GM BASE/VIAL</u>	<u>A065313 001</u> Jan 23, 2006
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065313 002</u> Jan 23, 2006
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065312 001</u> Feb 13, 2006
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 1GM BASE/VIAL</u>	<u>A065051 001</u> Sep 11, 2000
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>A065051 002</u> Sep 11, 2000
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>A065050 001</u> Sep 11, 2000

CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	+!	B BRAUN	<u>EQ 1GM BASE/VIAL</u>	<u>N065214 001</u> Mar 10, 2006
<u>AP</u>	+!		<u>EQ 2GM BASE/VIAL</u>	<u>N065214 002</u> Mar 10, 2006
		MEFOXIN IN PLASTIC CONTAINER		
	!	MYLAN INSTITUTIONAL	EQ 20MG BASE/ML	A063182 001 Jan 25, 1993
	!		EQ 40MG BASE/ML	A063182 002 Jan 25, 1993

POWDER; IV (INFUSION)

CEFOXITIN IN PLASTIC CONTAINER	SAMSON MEDCL	EQ 100GM BASE	A200938 001	Nov 16, 2015
--------------------------------	--------------	---------------	-------------	--------------

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETIL

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 50MG BASE/5ML</u>	<u>A065409 001</u>	Jun 08, 2007
<u>AB</u>		<u>EQ 100MG BASE/5ML</u>	<u>A065409 002</u>	Jun 08, 2007
<u>AB</u>	SANDOZ	<u>EQ 50MG BASE/5ML</u>	<u>A090031 001</u>	Jan 14, 2009
<u>AB</u>	!	<u>EQ 100MG BASE/5ML</u>	<u>A090031 002</u>	Jan 14, 2009

TABLET; ORAL

CEFPODOXIME PROXETIL

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 100MG BASE</u>	<u>A065370 001</u>	Jun 11, 2007
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A065370 002</u>	Jun 11, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-83 (of 436)

CEFPODOXIME PROXETIL

TABLET;ORAL

CEFPODOXIME PROXETIL

<u>AB</u>	ORCHID HLTHCARE	<u>EQ 100MG BASE</u>	<u>A065388 001</u>	Nov 14, 2007
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A065388 002</u>	Nov 14, 2007
<u>AB</u>	SANDOZ	<u>EQ 100MG BASE</u>	<u>A065462 001</u>	May 28, 2008
<u>AB</u>	!	<u>EQ 200MG BASE</u>	<u>A065462 002</u>	May 28, 2008

CEFPROZIL

FOR SUSPENSION;ORAL

CEFPROZIL

<u>AB</u>	APOTEX INC	<u>125MG/5ML</u>	<u>A065351 001</u>	Feb 29, 2012
<u>AB</u>		<u>250MG/5ML</u>	<u>A065351 002</u>	Feb 29, 2012
<u>AB</u>	AUROBINDO PHARMA	<u>125MG/5ML</u>	<u>A065381 001</u>	Jan 30, 2007
<u>AB</u>		<u>250MG/5ML</u>	<u>A065381 002</u>	Jan 30, 2007
<u>AB</u>	LUPIN	<u>125MG/5ML</u>	<u>A065261 001</u>	Dec 19, 2005
<u>AB</u>	!	<u>250MG/5ML</u>	<u>A065261 002</u>	Dec 19, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>125MG/5ML</u>	<u>A065284 002</u>	Dec 30, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065284 001</u>	Dec 30, 2005
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>A065257 001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065257 002</u>	Dec 08, 2005
<u>AB</u>	TEVA PHARMS	<u>125MG/5ML</u>	<u>A065236 001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>A065236 002</u>	Dec 08, 2005

TABLET;ORAL

CEFPROZIL

<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A065327 001</u>	Mar 26, 2008
<u>AB</u>		<u>500MG</u>	<u>A065327 002</u>	Mar 26, 2008
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A065340 001</u>	May 24, 2007
<u>AB</u>		<u>500MG</u>	<u>A065340 002</u>	May 24, 2007
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A065276 001</u>	Dec 08, 2005
<u>AB</u>	!	<u>500MG</u>	<u>A065276 002</u>	Dec 08, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>250MG</u>	<u>A065267 001</u>	Dec 19, 2005
<u>AB</u>		<u>500MG</u>	<u>A065267 002</u>	Dec 19, 2005
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065235 001</u>	Nov 14, 2005
<u>AB</u>		<u>500MG</u>	<u>A065235 002</u>	Nov 14, 2005
<u>AB</u>	TEVA	<u>250MG</u>	<u>A065208 001</u>	Dec 06, 2005
<u>AB</u>		<u>500MG</u>	<u>A065208 002</u>	Dec 06, 2005
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A065428 001</u>	Jun 14, 2007
<u>AB</u>		<u>500MG</u>	<u>A065428 002</u>	Jun 14, 2007

CEFTAROLINE FOSAMIL

POWDER;IV (INFUSION)

TEFLARO

+ CEREXA	400MG/VIAL
+ !	600MG/VIAL

N200327 001 Oct 29, 2010  
N200327 002 Oct 29, 2010

CEFTAZIDIME

INJECTABLE;INJECTION

CEFTAZIDIME

<u>AP</u>	ACS DOBFAR	<u>500MG/VIAL</u>	<u>A062640 001</u>	Nov 20, 1985
<u>AP</u>		<u>1GM/VIAL</u>	<u>A062640 002</u>	Nov 20, 1985
<u>AP</u>		<u>2GM/VIAL</u>	<u>A062640 003</u>	Nov 20, 1985
<u>AP</u>		<u>6GM/VIAL</u>	<u>A062640 004</u>	Feb 03, 1992
<u>AP</u>	WOCKHARDT	<u>1GM/VIAL</u>	<u>A065196 001</u>	Oct 15, 2008
<u>FORTAZ</u>				
<u>AP</u>	+! TELIGENT	<u>500MG/VIAL</u>	<u>N050578 001</u>	Jul 19, 1985
<u>AP</u>	+!	<u>1GM/VIAL</u>	<u>N050578 002</u>	Jul 19, 1985
<u>AP</u>	+!	<u>2GM/VIAL</u>	<u>N050578 003</u>	Jul 19, 1985
<u>AP</u>	+!	<u>6GM/VIAL</u>	<u>N050578 004</u>	Jul 19, 1985

TAZICEF

<u>AP</u>	HOSPIRA	<u>500MG/VIAL</u>	<u>A062662 001</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>A062662 002</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>A064032 001</u>	Oct 31, 1993
<u>AP</u>		<u>2GM/VIAL</u>	<u>A062662 003</u>	Mar 06, 1986
<u>AP</u>		<u>2GM/VIAL</u>	<u>A064032 002</u>	Oct 31, 1993
<u>AP</u>		<u>6GM/VIAL</u>	<u>A062662 004</u>	Mar 06, 1986

CEFTAZIDIME IN DEXTROSE CONTAINER

+ B BRAUN	EQ 1GM BASE
+ !	EQ 2GM BASE

N050823 001 Jun 13, 2011  
N050823 002 Jun 13, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-84 (of 436)

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

FORTAZ IN PLASTIC CONTAINER

+!	TELIGENT	EQ 20MG BASE/ML
+!		EQ 40MG BASE/ML

N050634 002	Apr 28, 1989
N050634 003	Apr 28, 1989

CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM

POWDER; IV (INFUSION)

ZERBAXA

+!	CUBIST PHARMS LLC	EQ 1GM BASE/VIAL; EQ 0.5GM BASE/VIAL
----	-------------------	--------------------------------------

N206829 001	Dec 19, 2014
-------------	--------------

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

CEFTRIAXONE

AP ACS DOBFAR

EQ 500MG BASE/VIAL

A065329 001 Jul 24, 2008

AP

EQ 1GM BASE/VIAL

A065329 002 Jul 24, 2008

AP

EQ 2GM BASE/VIAL

A065329 003 Jul 24, 2008

AP

EQ 10GM BASE/VIAL

A065328 001 Jul 24, 2008

AP BEDFORD

EQ 10GM BASE/VIAL

A065475 001 Aug 18, 2008

AP HOSPIRA INC

EQ 10GM BASE/VIAL

A065232 001 Aug 02, 2005

AP LUPIN

EQ 10GM BASE/VIAL

A065263 001 Sep 12, 2006

AP ! SANDOZ

EQ 10GM BASE/VIAL

A065168 001 May 17, 2005

AP ! SANDOZ INC

EQ 1GM BASE/VIAL

A065204 001 May 03, 2005

AP !

EQ 2GM BASE/VIAL

A065204 002 May 03, 2005

AP WOCKHARDT

EQ 1GM BASE/VIAL

A065180 001 May 12, 2006

CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER

AP +! B BRAUN

EQ 1GM BASE/VIAL

N050796 001 Apr 20, 2005

AP +!

EQ 2GM BASE/VIAL

N050796 002 Apr 20, 2005

CEFTRIAXONE SODIUM

AP HIKMA FARMACEUTICA

EQ 10GM BASE/VIAL

A090701 001 Oct 04, 2017

AP SAGENT PHARMS

EQ 10GM BASE/VIAL

A091117 001 Jan 20, 2017

CEFTRIAXONE

SAMSON MEDCL

EQ 100GM BASE/VIAL

A090057 001 Apr 25, 2014

CEFTRIAXONE IN PLASTIC CONTAINER

! BAXTER HLTHCARE

EQ 20MG BASE/ML

A065224 001 Aug 23, 2005

!

EQ 40MG BASE/ML

A065224 002 Aug 23, 2005

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFTRIAXONE

AP AKORN INC

EQ 250MG BASE/VIAL

A065305 001 Jan 11, 2008

AP

EQ 500MG BASE/VIAL

A065305 002 Jan 11, 2008

AP

EQ 1GM BASE/VIAL

A065305 003 Jan 11, 2008

AP

EQ 2GM BASE/VIAL

A065305 004 Jan 11, 2008

AP HIKMA FARMACEUTICA

EQ 250MG BASE/VIAL

A065342 001 Jan 10, 2008

AP

EQ 500MG BASE/VIAL

A065342 002 Jan 10, 2008

AP

EQ 1GM BASE/VIAL

A065342 003 Jan 10, 2008

AP

EQ 2GM BASE/VIAL

A065342 004 Jan 10, 2008

HOSPIRA INC

EQ 250MG BASE/VIAL

A065230 001 Aug 02, 2005

AP

EQ 500MG BASE/VIAL

A065230 002 Aug 02, 2005

AP

EQ 1GM BASE/VIAL

A065230 003 Aug 02, 2005

AP

EQ 2GM BASE/VIAL

A065230 004 Aug 02, 2005

LUPIN

EQ 250MG BASE/VIAL

A065125 001 Sep 30, 2003

AP

EQ 500MG BASE/VIAL

A065125 002 Sep 30, 2003

AP

EQ 1GM BASE/VIAL

A065125 003 Sep 30, 2003

AP

EQ 2GM BASE/VIAL

A065125 004 Sep 30, 2003

QILU PHARM CO LTD

EQ 250MG BASE/VIAL

A203702 001 Jun 29, 2016

AP

EQ 500MG BASE/VIAL

A203702 002 Jun 29, 2016

AP

EQ 1GM BASE/VIAL

A203702 003 Jun 29, 2016

AP

EQ 2GM BASE/VIAL

A203702 004 Jun 29, 2016

SANDOZ

EQ 250MG BASE/VIAL

A065169 001 May 09, 2005

AP !

EQ 500MG BASE/VIAL

A065169 002 May 09, 2005

AP !

EQ 1GM BASE/VIAL

A065169 003 May 09, 2005

AP !

EQ 2GM BASE/VIAL

A065169 004 May 09, 2005

WOCKHARDT

EQ 250MG BASE/VIAL

A065391 001 Apr 12, 2007

AP

EQ 500MG BASE/VIAL

A065391 002 Apr 12, 2007

AP

EQ 2GM BASE/VIAL

A065391 003 Apr 12, 2007

CEFUROXIME AXETIL

FOR SUSPENSION; ORAL

CEFTIN

+ GLAXOSMITHKLINE

EQ 125MG BASE/5ML

N050672 001 Jun 30, 1994

+

EQ 250MG BASE/5ML

N050672 002 Apr 29, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-85 (of 436)

CEFUROXIME AXETIL

TABLET; ORAL

CEFTIN

<u>AB</u> + GLAXOSMITHKLINE	<u>EQ 125MG BASE</u>	<u>N050605 001</u>	Dec 28, 1987
<u>AB</u> +	<u>EQ 250MG BASE</u>	<u>N050605 002</u>	Dec 28, 1987
<u>AB</u> +!	<u>EQ 500MG BASE</u>	<u>N050605 003</u>	Dec 28, 1987
<b>CEFUROXIME AXETIL</b>			
<u>AB</u> ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A065496 001</u>	Jun 07, 2010
<u>AB</u>	<u>EQ 500MG BASE</u>	<u>A065496 002</u>	Jun 07, 2010
<u>AB</u> ANI PHARMS INC	<u>EQ 250MG BASE</u>	<u>A065190 001</u>	Oct 18, 2004
<u>AB</u>	<u>EQ 500MG BASE</u>	<u>A065190 002</u>	Oct 18, 2004
<u>AB</u> APOTEX	<u>EQ 250MG BASE</u>	<u>A065069 001</u>	Oct 02, 2002
<u>AB</u>	<u>EQ 500MG BASE</u>	<u>A065069 002</u>	Oct 02, 2002
<u>AB</u> AUROBINDO PHARMA LTD	<u>EQ 125MG BASE</u>	<u>A065308 001</u>	Mar 29, 2006
	<u>EQ 250MG BASE</u>	<u>A065308 002</u>	Mar 29, 2006
	<u>EQ 500MG BASE</u>	<u>A065308 003</u>	Mar 29, 2006
<u>AB</u> LUPIN	<u>EQ 250MG BASE</u>	<u>A065135 001</u>	Jul 25, 2003
	<u>EQ 500MG BASE</u>	<u>A065135 002</u>	Jul 25, 2003
<u>AB</u> ORCHID HLTHCARE	<u>EQ 125MG BASE</u>	<u>A065359 001</u>	Feb 15, 2008
	<u>EQ 250MG BASE</u>	<u>A065359 002</u>	Feb 15, 2008
	<u>EQ 500MG BASE</u>	<u>A065359 003</u>	Feb 15, 2008
<u>AB</u> WOCKHARDT	<u>EQ 125MG BASE</u>	<u>A065166 001</u>	Jul 29, 2005
	<u>EQ 250MG BASE</u>	<u>A065166 002</u>	Jul 29, 2005
	<u>EQ 500MG BASE</u>	<u>A065166 003</u>	Jul 29, 2005

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u> +! B BRAUN	<u>EQ 750MG BASE/VIAL</u>	<u>N050780 001</u>	Feb 21, 2001
<u>AP</u> +!	<u>EQ 1.5GM BASE/VIAL</u>	<u>N050780 002</u>	Feb 21, 2001

CEFUROXIME SODIUM

<u>AP</u> ACS DOBFAR SPA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A064125 002</u>	May 30, 1997
<u>AP</u>	<u>EQ 7.5GM BASE/VIAL</u>	<u>A064124 001</u>	May 30, 1997
<u>AP</u> HIKMA FARMACEUTICA	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065048 002</u>	Jan 09, 2004
<u>AP</u>	<u>EQ 7.5GM BASE/VIAL</u>	<u>A065046 001</u>	Jan 09, 2004
<u>AP</u> HOSPIRA INC	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065483 002</u>	Oct 15, 2008
<u>AP</u>	<u>EQ 1.5GM BASE/VIAL</u>	<u>A065503 001</u>	Oct 15, 2008
<u>AP</u>	<u>EQ 7.5GM BASE/VIAL</u>	<u>A065484 001</u>	Oct 15, 2008

ZINACEF

<u>AP</u> +! TELIGENT	<u>EQ 1.5GM BASE/VIAL</u>	<u>N050558 003</u>	Oct 19, 1983
<u>AP</u> +!	<u>EQ 7.5GM BASE/VIAL</u>	<u>N050558 004</u>	Oct 23, 1986

ZINACEF IN PLASTIC CONTAINER

+! TELIGENT EQ 30MG BASE/ML

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

<u>AB</u> ACS DOBFAR SPA	<u>EQ 750MG BASE/VIAL</u>	<u>A064125 001</u>	May 30, 1997
<u>AB</u> HIKMA FARMACEUTICA	<u>EQ 750MG BASE/VIAL</u>	<u>A065048 001</u>	Jan 09, 2004
<b>ZINACEF</b>			
<u>AB</u> +! TELIGENT	<u>EQ 750MG BASE/VIAL</u>	<u>N050558 002</u>	Oct 19, 1983
<b>CEFUROXIME SODIUM</b>			
<u>AP</u> HOSPIRA INC	<u>EQ 750MG BASE/VIAL</u>	<u>A065483 001</u>	Oct 15, 2008

CELECOXIB

CAPSULE; ORAL

CELEBREX

<u>AB</u> + GD SEARLE	<u>50MG</u>	<u>N020998 004</u>	Dec 15, 2006
<u>AB</u> +	<u>100MG</u>	<u>N020998 001</u>	Dec 31, 1998
<u>AB</u> +	<u>200MG</u>	<u>N020998 002</u>	Dec 31, 1998
<u>AB</u> +!	<u>400MG</u>	<u>N020998 003</u>	Aug 29, 2002
<b>CELECOXIB</b>			
<u>AB</u> ALEMBIC PHARMS LTD	<u>50MG</u>	<u>A204519 001</u>	Aug 21, 2015
	<u>100MG</u>	<u>A204519 002</u>	Aug 21, 2015
	<u>200MG</u>	<u>A204519 003</u>	Aug 21, 2015
	<u>400MG</u>	<u>A204519 004</u>	Aug 21, 2015
<u>AB</u> APOTEX INC	<u>50MG</u>	<u>A204197 001</u>	Jun 02, 2015
	<u>100MG</u>	<u>A204197 002</u>	Jun 02, 2015
	<u>200MG</u>	<u>A204197 003</u>	Jun 02, 2015
<u>AB</u> AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A206827 001</u>	Feb 01, 2016
	<u>100MG</u>	<u>A206827 002</u>	Feb 01, 2016
	<u>200MG</u>	<u>A206827 003</u>	Feb 01, 2016
	<u>400MG</u>	<u>A206827 004</u>	Feb 01, 2016
<u>AB</u> CIPLA LTD	<u>50MG</u>	<u>A207446 001</u>	Sep 23, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-86 (of 436)

CELECOXIB

CAPSULE; ORAL

CELECOXIB

<u>AB</u>		<u>100MG</u>	<u>A207446 002</u>	Sep 23, 2015
<u>AB</u>		<u>200MG</u>	<u>A207446 003</u>	Sep 23, 2015
<u>AB</u>		<u>400MG</u>	<u>A207446 004</u>	Sep 23, 2015
<u>AB</u>	JUBILANT GENERICS	<u>50MG</u>	<u>A207061 001</u>	Apr 04, 2017
<u>AB</u>		<u>100MG</u>	<u>A207061 002</u>	Apr 04, 2017
<u>AB</u>		<u>200MG</u>	<u>A207061 003</u>	Apr 04, 2017
<u>AB</u>		<u>400MG</u>	<u>A207061 004</u>	Apr 04, 2017
<u>AB</u>	LUPIN LTD	<u>50MG</u>	<u>A202240 001</u>	Oct 29, 2014
<u>AB</u>		<u>100MG</u>	<u>A202240 002</u>	Jun 09, 2015
<u>AB</u>		<u>200MG</u>	<u>A202240 003</u>	Jun 09, 2015
<u>AB</u>		<u>400MG</u>	<u>A202240 004</u>	Jun 09, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>50MG</u>	<u>A204590 001</u>	Mar 16, 2016
<u>AB</u>		<u>100MG</u>	<u>A204590 002</u>	Mar 16, 2016
<u>AB</u>		<u>200MG</u>	<u>A204590 003</u>	Mar 16, 2016
<u>AB</u>		<u>400MG</u>	<u>A204590 004</u>	Mar 16, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A078857 001</u>	May 30, 2014
<u>AB</u>		<u>100MG</u>	<u>A078857 002</u>	Feb 11, 2015
<u>AB</u>		<u>200MG</u>	<u>A078857 003</u>	Feb 11, 2015
<u>AB</u>		<u>400MG</u>	<u>A078857 004</u>	Feb 11, 2015
<u>AB</u>	TEVA	<u>50MG</u>	<u>A076898 001</u>	May 30, 2014
<u>AB</u>		<u>100MG</u>	<u>A076898 002</u>	May 30, 2014
<u>AB</u>		<u>200MG</u>	<u>A076898 003</u>	May 30, 2014
<u>AB</u>		<u>400MG</u>	<u>A076898 004</u>	May 30, 2014
<u>AB</u>	TORRENT PHARMS LTD	<u>50MG</u>	<u>A207677 001</u>	Dec 23, 2015
<u>AB</u>		<u>100MG</u>	<u>A207677 002</u>	Dec 23, 2015
<u>AB</u>		<u>200MG</u>	<u>A207677 003</u>	Dec 23, 2015
<u>AB</u>		<u>400MG</u>	<u>A207677 004</u>	Dec 23, 2015
<u>AB</u>	WATSON LABS INC	<u>50MG</u>	<u>A200562 001</u>	Feb 11, 2015
<u>AB</u>		<u>100MG</u>	<u>A200562 002</u>	Feb 11, 2015
<u>AB</u>		<u>200MG</u>	<u>A200562 003</u>	Feb 11, 2015
<u>AB</u>		<u>400MG</u>	<u>A200562 004</u>	Feb 11, 2015

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<u>AB</u>	ALKEM LABS LTD	<u>EQ 250MG BASE</u>	<u>A090836 001</u>	Dec 20, 2010
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A090836 002</u>	Dec 20, 2010
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A090836 004</u>	Mar 29, 2013
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>A065253 001</u>	Nov 16, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065253 002</u>	Nov 16, 2005
<u>AB</u>	BELCHER PHARMS	<u>EQ 250MG BASE</u>	<u>A062713 001</u>	Jul 15, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062713 002</u>	Jul 15, 1988
<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>A065215 001</u>	Jan 24, 2006
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065215 002</u>	Jan 24, 2006
<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>A065229 001</u>	Nov 25, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065229 002</u>	Nov 25, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 250MG BASE</u>	<u>A065248 001</u>	Jun 28, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065248 002</u>	Jun 28, 2005
<u>AB</u>	SUN PHARM INDs (IN)	<u>EQ 250MG BASE</u>	<u>A062791 001</u>	Jun 11, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062791 002</u>	Jun 11, 1987
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A062702 001</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A062702 002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 250MG BASE</u>	<u>A065152 001</u>	Feb 24, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A065152 002</u>	Feb 24, 2005

KEFLEX

<u>AB</u>	+	PRAGMA PHARMS LLC	<u>EQ 250MG BASE</u>	<u>N050405 002</u>
<u>AB</u>	+		<u>EQ 500MG BASE</u>	<u>N050405 003</u>
<u>AB</u>	!+		<u>EQ 750MG BASE</u>	<u>N050405 005</u> May 12, 2006

CEPHALEXIN

ALKEM LABS LTD

EQ 333MG BASE

A090836 003 Mar 29, 2013

FOR SUSPENSION; ORAL

CEPHALEXIN

<u>AB</u>	LUPIN	<u>EQ 125MG BASE/5ML</u>	<u>A065234 001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065234 002</u>	Aug 17, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 125MG BASE/5ML</u>	<u>A065326 001</u>	Jul 10, 2006
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065326 002</u>	Jul 10, 2006
<u>AB</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>A062703 001</u>	Feb 13, 1987
<u>AB</u>	!	<u>EQ 250MG BASE/5ML</u>	<u>A062703 002</u>	Feb 13, 1987
<u>AB</u>	YUNG SHIN PHARM	<u>EQ 125MG BASE/5ML</u>	<u>A065336 001</u>	Jul 25, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-87 (of 436)

CEPHALEXIN

FOR SUSPENSION;ORAL

CEPHALEXIN

<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>A065336 002</u>	Jul 25, 2007
TABLET;ORAL				
CEPHALEXIN				
TEVA	EQ 250MG BASE		A063023 001	Jan 12, 1989

!

EQ 500MG BASE

A063024 001 Jan 12, 1989

CERITINIB

CAPSULE;ORAL

ZYKADIA

+! NOVARTIS PHARMS CORP

150MG

N205755 001 Apr 29, 2014

CETIRIZINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ZERVIASTE

+! EYEVANCE PHARMS

EQ 0.24% BASE

N208694 001 May 30, 2017

SYRUP;ORAL

CETIRIZINE HYDROCHLORIDE

<u>AA</u>	ALLIED PHARMA INC	<u>5MG/5ML</u>	<u>A090191 001</u>	Nov 12, 2009
<u>AA</u>	AMNEAL PHARMS	<u>5MG/5ML</u>	<u>A090766 001</u>	Oct 07, 2009
<u>AA</u>	BIO PHARM INC	<u>5MG/5ML</u>	<u>A078870 001</u>	Apr 27, 2009
<u>AA</u>	BRECKENRIDGE PHARM	<u>5MG/5ML</u>	<u>A078488 001</u>	Oct 06, 2008
<u>AA</u> !	PERRIGO R AND D	<u>5MG/5ML</u>	<u>A078398 001</u>	Jun 17, 2008
<u>AA</u>	SILARX	<u>5MG/5ML</u>	<u>A078876 001</u>	May 11, 2012
<u>AA</u>	TARO	<u>5MG/5ML</u>	<u>A076601 001</u>	Jun 20, 2008
<u>AA</u>	TEVA PHARMS	<u>5MG/5ML</u>	<u>A077279 001</u>	May 27, 2008
<u>AA</u>	VINTAGE	<u>5MG/5ML</u>	<u>A078496 001</u>	Sep 25, 2009

CETRORELIX

INJECTABLE; INJECTION

CETROTIDE

+! EMD SERONO INC

EQ 0.25MG BASE/ML

N021197 001 Aug 11, 2000

CEVIMELINE HYDROCHLORIDE

CAPSULE;ORAL

CEVIMELINE HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>30MG</u>	<u>A091260 001</u>	Aug 25, 2011
<u>AB</u>	NOVEL LABS INC	<u>30MG</u>	<u>A204746 001</u>	Dec 30, 2016
<u>AB</u>	RISING PHARMS INC	<u>30MG</u>	<u>A203775 001</u>	Jun 04, 2014
<u>AB</u>	WEST-WARD PHARMS INT	<u>30MG</u>	<u>A091591 001</u>	Jul 08, 2013

EVOXAC

<u>AB</u> +!	DAIICHI SANKYO INC	<u>30MG</u>	<u>N020989 002</u>	Jan 11, 2000
--------------	--------------------	-------------	--------------------	--------------

CHENODIOL

TABLET;ORAL

CHENODIOL

! NEXGEN PHARMA

250MG

A091019 001 Oct 22, 2009

CHLORAMBUCIL

TABLET;ORAL

LEUKERAN

+! ASPEN GLOBAL INC

2MG

N010669 002

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

! FRESENIUS KABI USA EQ 1GM BASE/VIAL

A062365 001 Aug 25, 1982

CHLORDIAZEPoxide HYDROCHLORIDE

CAPSULE;ORAL

CHLORDIAZEPoxide HYDROCHLORIDE

<u>AB</u>	BARR	<u>5MG</u>	<u>A084768 001</u>	
<u>AB</u>		<u>10MG</u>	<u>A083116 001</u>	
<u>AB</u>		<u>25MG</u>	<u>A084769 001</u>	

LIBRIUM

<u>AB</u>	VALEANT PHARM INTL	<u>5MG</u>	<u>A085461 001</u>	
<u>AB</u>		<u>10MG</u>	<u>A085472 001</u>	
<u>AB</u> !		<u>25MG</u>	<u>A085475 001</u>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-88 (of 436)

CHLORDIAZEPOXIDE HYDROCHLORIDE; CLIDINIUM BROMIDE

CAPSULE; ORAL  
 LIBRAX  
 +! VALEANT PHARMS 5MG; 2.5MG N012750 001

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

<u>AT</u>	HI TECH PHARMA	<u>0.12%</u>	<u>A074356 001</u>	May 07, 1996
<u>AT</u>	LYNE	<u>0.12%</u>	<u>A074291 001</u>	Dec 28, 1995
<u>AT</u>	TEVA	<u>0.12%</u>	<u>A074522 001</u>	Dec 15, 1995
<u>AT</u>	WOCKHARDT BIO AG	<u>0.12%</u>	<u>A075006 001</u>	Mar 03, 2004
<u>AT</u>	XTTRIUM	<u>0.12%</u>	<u>A077789 001</u>	Jun 18, 2009

PAROEX

<u>AT</u>	SUNSTAR AMERICAS	<u>0.12%</u>	<u>A076434 001</u>	Nov 29, 2005
-----------	------------------	--------------	--------------------	--------------

PERIDEX

<u>AT</u>	+! 3M	<u>0.12%</u>	<u>N019028 001</u>	Aug 13, 1986
-----------	-------	--------------	--------------------	--------------

PERIOGARD

<u>AT</u>	COLGATE PALMOLIVE CO	<u>0.12%</u>	<u>A073695 001</u>	Jan 14, 1994
<u>AT</u>	COLGATE-PALMOLIVE CO	<u>0.12%</u>	<u>A203212 001</u>	Jan 28, 2016

TABLET; DENTAL

PERIOCHIP

+!	DEXCEL PHARMA	2.5MG	N020774 001	May 15, 1998
----	---------------	-------	-------------	--------------

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>2%</u>	<u>A087447 001</u>	Apr 16, 1982
<u>AP</u>		<u>3%</u>	<u>A087446 001</u>	Apr 16, 1982
<u>AP</u>	WEST-WARD PHARMS INT	<u>2%</u>	<u>A040273 001</u>	Sep 09, 1998
<u>AP</u>		<u>3%</u>	<u>A040273 002</u>	Sep 09, 1998

NESACAIN

<u>AP</u>	+ FRESENIUS KABI USA	<u>2%</u>	<u>N009435 002</u>	
-----------	----------------------	-----------	--------------------	--

NESACAIN-MPF

<u>AP</u>	+! FRESENIUS KABI USA	<u>2%</u>	<u>N009435 006</u>	May 02, 1996
<u>AP</u>	+!	<u>3%</u>	<u>N009435 007</u>	May 02, 1996

NESACAIN

+!	FRESENIUS KABI USA	1%	N009435 001	
----	--------------------	----	-------------	--

SOLUTION; INTRATHECAL

CLOROTEKAL

+	B BRAUN MEDICAL INC	50MG/5ML (10MG/ML)	N208791 001	Sep 26, 2017
---	---------------------	--------------------	-------------	--------------

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE

<u>AA</u>	! HIKMA PHARMS	<u>EQ 150MG BASE</u>	<u>A083082 001</u>	
<u>AA</u>		<u>EQ 300MG BASE</u>	<u>A083082 002</u>	Sep 17, 1999
<u>AA</u>	IPCA LABS LTD	<u>EQ 150MG BASE</u>	<u>A090610 001</u>	Dec 03, 2009
<u>AA</u>		<u>EQ 300MG BASE</u>	<u>A090249 001</u>	Dec 03, 2009
<u>AA</u>	NATCO PHARMA LTD	<u>EQ 150MG BASE</u>	<u>A091621 001</u>	Jan 21, 2011
<u>AA</u>	!	<u>EQ 300MG BASE</u>	<u>A090612 001</u>	Jan 21, 2011

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+!	SALIX PHARMS	250MG/5ML	N011870 001	
----	--------------	-----------	-------------	--

TABLET; ORAL

CHLOROTHIAZIDE

!	MYLAN	250MG	A084217 002	
		500MG	A084217 001	

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

CHLOROTHIAZIDE SODIUM

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 500MG BASE/VIAL</u>	<u>A090896 001</u>	Oct 16, 2009
<u>AP</u>	LUITPOLD	<u>EQ 500MG BASE/VIAL</u>	<u>A202561 001</u>	Apr 22, 2013
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 500MG BASE/VIAL</u>	<u>A202493 001</u>	Jun 18, 2014
<u>AP</u>	SAGENT PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>A202462 001</u>	May 29, 2015
<u>AP</u>	SUN PHARMA GLOBAL	<u>EQ 500MG BASE/VIAL</u>	<u>A091546 001</u>	Jul 26, 2011

DIURIL

+!	OAK PHARMS AKORN	<u>EQ 500MG BASE/VIAL</u>	<u>N011145 005</u>	
----	------------------	---------------------------	--------------------	--

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-89 (of 436)

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

AA ACELLA PHARMS LLC 4MG/5ML;5MG/5ML A206891 001 Jun 09, 2017

VITUZ

AA +! CYPRESS PHARM 4MG/5ML;5MG/5ML N204307 001 Feb 20, 2013

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AA BIO-PHARM INC 4MG/5ML;5MG/5ML;60MG/5ML A206660 001 May 15, 2017

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AA PADDOCK LLC 4MG/5ML;5MG/5ML;60MG/5ML A204627 001 Apr 29, 2014

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

AA MAYNE PHARMA INC 4MG/5ML;5MG/5ML;60MG/5ML A205657 001 Aug 03, 2015

ZUTRIPRO

AA +! CYPRESS PHARM 4MG/5ML;5MG/5ML;60MG/5ML N022439 001 Jun 08, 2011

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

TUZISTRA XR

+! VERNALIS R AND D LTD EQ 2.8MG BASE/5ML;EQ 14.7MG BASE/5ML N207768 001 Apr 30, 2015

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

CAPSULE, EXTENDED RELEASE;ORAL

TUSSICAPS

ECR PHARMA EQ 4MG MALEATE;EQ 5MG BITARTRATE A077273 002 Sep 24, 2007  
! EQ 8MG MALEATE;EQ 10MG BITARTRATE A077273 001 Sep 24, 2007

SUSPENSION, EXTENDED RELEASE;ORAL

HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX

AB TRIS PHARMA INC EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML A091632 001 Oct 01, 2010

HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX

AB NEOS THERAP INC EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML A091671 001 Jun 29, 2012

TUSSIONEX PENN KINETIC

AB +! UCB INC EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML N019111 001 Dec 31, 1987

CHLORPROMAZINE HYDROCHLORIDE

INJECTABLE;INJECTION

CHLORPROMAZINE HYDROCHLORIDE

! WEST-WARD PHARMS 25MG/ML INT A083329 001

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

USL PHARMA 10MG A083386 001  
! 25MG A084112 001  
50MG A084113 001  
! 100MG A084114 001  
200MG A084115 001

CHLORPROPAMIDE

TABLET;ORAL

CHLORPROPAMIDE

AB ANI PHARMS INC 100MG A088921 001 Apr 12, 1985

AB 250MG A088922 001 Apr 12, 1985

AB MYLAN 100MG A088549 002 Jun 01, 1984

AB 250MG A088549 001 Jun 01, 1984

DIABINESE

AB + PFIZER 100MG N011641 003

AB +! 250MG N011641 006

GLUCAMIDE

AB ANI PHARMS INC 250MG A088641 001 Oct 11, 1984

CHLORTHALIDONE

TABLET;ORAL

CHLORTHALIDONE

AB MYLAN 25MG A086831 002

AB ! 50MG A086831 001

AB RICONPHARMA LLC 25MG A206904 001 Mar 30, 2017

AB 50MG A206904 002 Mar 30, 2017

AB SUN PHARM INDUSTRIES 25MG A089286 002 Jul 21, 1986

AB 50MG A089286 001 Jul 21, 1986

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-90 (of 436)

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLORPRES

MYLAN	15MG; 0.1MG	A071325 003	Feb 09, 1987
	15MG; 0.2MG	A071325 002	Feb 09, 1987
!	15MG; 0.3MG	A071325 001	Feb 09, 1987

CHLORZOXAZONE

TABLET; ORAL

**CHLORZOXAZONE**

<b>AA</b>	BARR	<b>500MG</b>	<b>A089895 001</b>	May 04, 1988
<b>AA</b>	! WATSON LABS	<b>500MG</b>	<b>A089859 001</b>	May 04, 1988
	MIKART	375MG	A040861 001	Jun 01, 2010
!		750MG	A040861 002	Jun 01, 2010
!	MIKART INC	250MG	A207483 001	Jun 24, 2016

CHOLESTYRAMINE

POWDER; ORAL

**CHOLESTYRAMINE**

<b>AB</b>	PAR PHARM	<b>EQ 4GM RESIN/PACKET</b>	<b>A077204 001</b>	Aug 26, 2005
<b>AB</b>		<b>EQ 4GM RESIN/SCOOPFUL</b>	<b>A077204 002</b>	Aug 26, 2005
<b>AB</b>	! SANDOZ	<b>EQ 4GM RESIN/PACKET</b>	<b>A074557 001</b>	Aug 15, 1996
<b>AB</b>		<b>EQ 4GM RESIN/SCOOPFUL</b>	<b>A074557 002</b>	Aug 15, 1996

**CHOLESTYRAMINE LIGHT**

<b>AB</b>	PAR PHARM	<b>EQ 4GM RESIN/PACKET</b>	<b>A077203 001</b>	Aug 26, 2005
<b>AB</b>		<b>EQ 4GM RESIN/SCOOPFUL</b>	<b>A077203 002</b>	Aug 26, 2005
<b>AB</b>	! SANDOZ	<b>EQ 4GM RESIN/PACKET</b>	<b>A074558 001</b>	Aug 15, 1996
<b>AB</b>		<b>EQ 4GM RESIN/SCOOPFUL</b>	<b>A074558 002</b>	Aug 15, 1996
<b>AB</b>	ZYDUS PHARMS USA INC	<b>EQ 4GM RESIN/SCOOPFUL</b>	<b>A202902 001</b>	Apr 25, 2017

**PREVALITE**

<b>AB</b>	UPSHER-SMITH LABS	<b>EQ 4GM RESIN/PACKET</b>	<b>A073263 001</b>	Feb 22, 1996
<b>AB</b>		<b>EQ 4GM RESIN/SCOOPFUL</b>	<b>A073263 002</b>	Oct 30, 1997

CHOLIC ACID

CAPSULE; ORAL

CHOLBAM

+	RTRX	50MG	N205750 001	Mar 17, 2015
+		250MG	N205750 002	Mar 17, 2015

CHOLINE C-11

INJECTABLE; INTRAVENOUS

**CHOLINE C-11**

<b>AP</b>	GLOBAL ISOTOPES LLC	<b>4-33.1mCi/ML</b>	<b>A206319 001</b>	Nov 13, 2015
<b>AP</b>	+! MCPRF	<b>4-33.1mCi/ML</b>	<b>N203155 001</b>	Sep 12, 2012
<b>AP</b>	UCSF RODIOPHARM	<b>4-33.1mCi/ML</b>	<b>A208444 001</b>	Nov 20, 2017
<b>AP</b>	WA UNIV SCH MED	<b>4-33.1mCi/ML</b>	<b>A208413 001</b>	Jan 10, 2017
	UNIV TX MD ANDERSON	4-100mCi/ML	A205690 001	Oct 29, 2015

CHOLINE FENOFLIBRATE

CAPSULE, DELAYED RELEASE; ORAL

**FENOFLIBRIC ACID**

<b>AB</b>	ACTAVIS ELIZABETH	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>A200920 001</b>	Oct 07, 2015
<b>AB</b>		<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>A200920 002</b>	Oct 07, 2015
<b>AB</b>	ALEMBIC PHARMS LTD	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>A208705 001</b>	May 12, 2017
<b>AB</b>		<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>A208705 002</b>	May 12, 2017
<b>AB</b>	ANCHEN PHARMS	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>A201573 002</b>	Jul 18, 2013
<b>AB</b>		<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>A201573 001</b>	Jul 18, 2013
<b>AB</b>	IMPAX LABS INC	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>A200264 001</b>	Sep 07, 2016
<b>AB</b>		<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>A200264 002</b>	Sep 07, 2016
<b>AB</b>	LUPIN LTD	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>A200750 001</b>	Dec 04, 2013
<b>AB</b>		<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>A200750 002</b>	Dec 04, 2013
<b>AB</b>	MYLAN PHARMS INC	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>A200913 001</b>	Mar 25, 2013
<b>AB</b>		<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>A200913 002</b>	Mar 25, 2013

**TRILIPIX**

<b>AB</b>	++ ABBVIE	<b>EQ 45MG FENOFLIBRIC ACID</b>	<b>N022224 001</b>	Dec 15, 2008
<b>AB</b>	++!	<b>EQ 135MG FENOFLIBRIC ACID</b>	<b>N022224 002</b>	Dec 15, 2008

CHORIOGONADOTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

OVIDREL

++!	EMD SERONO	EQ 0.25MG / 0.5ML	N021149 002	Oct 06, 2003
-----	------------	-------------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-91 (of 436)

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA

EQ 0.004MG CHROMIUM/ML

N018961 001 Jun 26, 1986

CICLESONIDE

AEROSOL, METERED; INHALATION

ALVESCO

+! ASTRAZENECA PHARMS 0.08MG/INH

+! 0.16MG/INH

N021658 002 Jan 10, 2008

N021658 003 Jan 10, 2008

AEROSOL, METERED; NASAL

ZETONNA

+! ASTRAZENECA PHARMS 0.037MG/INH

N202129 001 Jan 20, 2012

SPRAY, METERED; NASAL

OMNARIS

+! ASTRAZENECA PHARMS 0.05MG/INH

N022004 001 Oct 20, 2006

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

<b>AB</b>	FOUGERA PHARMS	<b>0.77%</b>	
<b>AB</b>	G AND W LABS INC	<b>0.77%</b>	
<b>AB</b>	GLENMARK PHARMS	<b>0.77%</b>	
<b>AB</b>	PERRIGO NEW YORK	<b>0.77%</b>	
<b>AB</b>	TARO	<b>0.77%</b>	

<b>A076435 001</b>	Dec 29, 2004
<b>A078463 001</b>	Dec 20, 2010
<b>A090273 001</b>	Nov 10, 2009
<b>A077364 001</b>	Mar 03, 2006
<b>A076790 001</b>	Apr 12, 2005

LOPROX

<b>AB</b>	+! MEDIMETRIKS PHARMS	<b>0.77%</b>	
	GEL; TOPICAL		

**N018748 001** Dec 30, 1982

CICLOPIROX

<b>AB</b>	+! CNTY LINE PHARMS	<b>0.77%</b>	
<b>AB</b>	FOUGERA PHARMS	<b>0.77%</b>	
<b>AB</b>	GLENMARK GENERICS	<b>0.77%</b>	
<b>AB</b>	PADDOCK LLC	<b>0.77%</b>	

<b>N020519 001</b>	Jul 21, 1997
<b>A077896 001</b>	Jun 10, 2008
<b>A091595 001</b>	Feb 29, 2012
<b>A078266 001</b>	Jan 07, 2009

SHAMPOO; TOPICAL

CICLOPIROX

<b>AT</b>	ACTAVIS MID ATLANTIC	<b>1%</b>	
<b>AT</b>	FOUGERA PHARMS	<b>1%</b>	
<b>AT</b>	PERRIGO CO	<b>1%</b>	
<b>AT</b>	TARO	<b>1%</b>	

<b>A090490 001</b>	Nov 24, 2009
<b>A090146 001</b>	May 25, 2010
<b>A078594 001</b>	Feb 16, 2010
<b>A090269 001</b>	Feb 23, 2011

LOPROX

<b>AT</b>	+! MEDICIS SOLUTION; TOPICAL	<b>1%</b>	
-----------	------------------------------	-----------	--

**N021159 001** Feb 28, 2003

CICLOPIROX

<b>AT</b>	ACTAVIS MID ATLANTIC	<b>8%</b>	
<b>AT</b>	AKORN	<b>8%</b>	
<b>AT</b>	APOTEX INC	<b>8%</b>	
<b>AT</b>	CIPLA LTD	<b>8%</b>	
<b>AT</b>	G AND W LABS	<b>8%</b>	
<b>AT</b>	HI TECH PHARMA	<b>8%</b>	
<b>AT</b>	PERRIGO NEW YORK	<b>8%</b>	
<b>AT</b>	TARO PHARM IND	<b>8%</b>	
<b>AT</b>	TOLMAR	<b>8%</b>	

<b>A078046 001</b>	Sep 18, 2007
<b>A078975 001</b>	Feb 17, 2010
<b>A078172 001</b>	Sep 18, 2007
<b>A078124 001</b>	Sep 18, 2007
<b>A078233 001</b>	Sep 18, 2007
<b>A078270 001</b>	Sep 18, 2007
<b>A077623 001</b>	Sep 18, 2007
<b>A078144 001</b>	Sep 18, 2007
<b>A077687 001</b>	Sep 18, 2007

PENLAC

<b>AT</b>	+! VALEANT BERMUDA SUSPENSION; TOPICAL	<b>8%</b>	
-----------	--	-----------	--

**N021022 001** Dec 17, 1999

CICLOPIROX

<b>AB</b>	FOUGERA PHARMS	<b>0.77%</b>	
<b>AB</b>	PERRIGO NEW YORK	<b>0.77%</b>	
<b>AB</b>	TARO	<b>0.77%</b>	

<b>A076422 001</b>	Aug 06, 2004
<b>A077676 001</b>	Dec 15, 2006
<b>A077092 001</b>	Aug 10, 2005

LOPROX

<b>AB</b>	+! MEDIMETRIKS PHARMS	<b>0.77%</b>	
-----------	-----------------------	--------------	--

**N019824 001** Dec 30, 1988

CIDOFIVIR

INJECTABLE; INJECTION

CIDOFOVIR

<b>AP</b>	EMCURE PHARMS LTD	<b>EQ 75MG BASE/ML</b>	
<b>AP</b>	! MYLAN INSTITUTIONAL	<b>EQ 75MG BASE/ML</b>	

<b>A202501 001</b>	Jul 26, 2012
<b>A201276 001</b>	Jun 27, 2012



38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-93 (of 436)

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPROFLOXACIN

CORP

<u>AP</u>	!		<u>400MG/40ML (10MG/ML)</u>
<u>AP</u>		HIKMA FARMACEUTICA	<u>200MG/20ML (10MG/ML)</u>
<u>AP</u>			<u>400MG/40ML (10MG/ML)</u>
<u>AP</u>		HOSPIRA	<u>200MG/20ML (10MG/ML)</u>
<u>AP</u>			<u>400MG/40ML (10MG/ML)</u>

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>		ACS DOBFAR INFO SA	<u>200MG/100ML</u>
<u>AP</u>			<u>400MG/200ML</u>
<u>AP</u>		BAXTER HLTHCARE CORP	<u>200MG/100ML</u>
<u>AP</u>			<u>400MG/200ML</u>
<u>AP</u>		HIKMA FARMACEUTICA	<u>400MG/200ML</u>
<u>AP</u>	!	HOSPIRA	<u>200MG/100ML</u>
<u>AP</u>	!		<u>400MG/200ML</u>

INJECTABLE, SUSPENSION;OTIC

OTIPRIO

+! OTONOMY INC

6% (60MG/ML)

N207986 001 Dec 10, 2015

CIPROFLOXACIN HYDROCHLORIDE

OINTMENT;OPHTHALMIC

CILOXAN

+! NOVARTIS PHARMS CORP

SOLUTION/DROPS;OPHTHALMIC

CILOXAN

<u>AT</u>	+!	NOVARTIS PHARMS CORP	<u>EQ 0.3% BASE</u>
-----------	----	----------------------	---------------------

CIPROFLOXACIN HYDROCHLORIDE

<u>AT</u>		AKORN INC	<u>EQ 0.3% BASE</u>
<u>AT</u>		FDC LTD	<u>EQ 0.3% BASE</u>
<u>AT</u>		RISING PHARMS INC	<u>EQ 0.3% BASE</u>
<u>AT</u>		TELIGENT	<u>EQ 0.3% BASE</u>
<u>AT</u>		WATSON LABS INC	<u>EQ 0.3% BASE</u>

SOLUTION/DROPS;OTIC

CETRAXAL

+! WRASER PHARMS

EQ 0.2% BASE

N021918 001 May 01, 2009

TABLET;ORAL

CIPRO

<u>AB</u>	+	BAYER HLTHCARE	<u>EQ 100MG BASE</u>
<u>AB</u>	+		<u>EQ 250MG BASE</u>
<u>AB</u>	+		<u>EQ 500MG BASE</u>
<u>AB</u>	+		<u>EQ 750MG BASE</u>

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>		APOTEX	<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>
<u>AB</u>			<u>EQ 750MG BASE</u>

<u>AB</u>		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>
<u>AB</u>			<u>EQ 750MG BASE</u>

<u>AB</u>		CARLSBAD	<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>
<u>AB</u>			<u>EQ 750MG BASE</u>

<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 100MG BASE</u>
<u>AB</u>			<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>

<u>AB</u>			<u>EQ 750MG BASE</u>
<u>AB</u>		HIKMA	<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>

<u>AB</u>			<u>EQ 750MG BASE</u>
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>

<u>AB</u>			<u>EQ 750MG BASE</u>
<u>AB</u>		MYLAN	<u>EQ 100MG BASE</u>
<u>AB</u>			<u>EQ 250MG BASE</u>

<u>AB</u>			<u>EQ 500MG BASE</u>
<u>AB</u>		SUN PHARM INDs LTD	<u>EQ 750MG BASE</u>
<u>AB</u>			<u>EQ 250MG BASE</u>

<u>AB</u>			<u>EQ 750MG BASE</u>
<u>AB</u>			<u>EQ 250MG BASE</u>
<u>AB</u>			<u>EQ 500MG BASE</u>

A078062 002 Apr 29, 2008  
A076717 001 Dec 22, 2009  
A076717 002 Dec 22, 2009  
A077245 001 Aug 28, 2006  
A077245 002 Aug 28, 2006

A078252 001 Mar 18, 2008  
A078252 002 Mar 18, 2008  
A078024 001 Mar 18, 2008  
A078024 002 Mar 18, 2008  
A078431 001 Nov 18, 2009  
A077753 001 Mar 18, 2008  
A077753 002 Mar 18, 2008

N020369 001 Mar 30, 1998  
N019992 001 Dec 31, 1990  
A076555 001 Dec 11, 2008  
A077568 001 Jun 30, 2008  
A077689 001 Dec 13, 2006  
A076754 001 Jun 09, 2004  
A076673 001 Jan 21, 2005

N019537 001 Apr 08, 1996  
N019537 002 Oct 22, 1987  
N019537 003 Oct 22, 1987  
N019537 004 Oct 22, 1987  
A076896 001 Nov 04, 2004  
A076896 002 Nov 04, 2004  
A076896 003 Nov 04, 2004  
A077859 001 Apr 26, 2007  
A077859 002 Apr 26, 2007  
A077859 003 Apr 26, 2007  
A076126 002 Jun 09, 2004  
A076126 003 Jun 09, 2004  
A076126 004 Jun 09, 2004  
A075593 002 Jun 09, 2004  
A075593 003 Jun 09, 2004  
A075593 004 Jun 09, 2004  
A075593 001 Jun 09, 2004  
A076558 002 Jun 09, 2004  
A076558 003 Jun 09, 2004  
A076558 004 Jun 09, 2004  
A076089 002 Jun 09, 2004  
A076089 003 Jun 09, 2004  
A076089 004 Jun 09, 2004  
A075817 001 Jun 25, 2007  
A075817 002 Jun 09, 2004  
A075817 003 Jun 09, 2004  
A075817 004 Jun 09, 2004  
A075747 001 Jun 09, 2004  
A075747 002 Jun 09, 2004  
A075747 003 Jun 09, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-94 (of 436)

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

<u>AB</u>	TARO PHARM	<u>EQ 100MG BASE</u>	<u>A076912 001</u>	Feb 18, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A076912 002</u>	Oct 06, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076912 003</u>	Oct 06, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076912 004</u>	Oct 06, 2004
<u>AB</u>	UNIQUE PHARM LABS	<u>EQ 250MG BASE</u>	<u>A076639 001</u>	Sep 10, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076639 002</u>	Sep 10, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076639 003</u>	Sep 10, 2004
<u>AB</u>	WATSON LABS	<u>EQ 100MG BASE</u>	<u>A076794 001</u>	Feb 10, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A076794 002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A076794 003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A076794 004</u>	Jun 09, 2004

CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE

SOLUTION/DROPS; OTIC

OTOVEL

+! LABORATORIOS SALVAT EQ 0.3% BASE; 0.025%

N208251 001 Apr 29, 2016

CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS; OTIC

CIPRO HC

+! NOVARTIS PHARMS EQ 0.2% BASE; 1%  
CORP

N020805 001 Feb 10, 1998

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPROFLOXACIN EXTENDED RELEASE

<u>AB</u>	ANCHEN PHARMS	<u>212.6MG;EQ 287.5MG BASE</u>	<u>A078166 002</u>	Nov 27, 2007
<u>AB</u>		<u>425.2MG;EQ 574.9MG BASE</u>	<u>A078166 001</u>	Nov 27, 2007
<u>AB</u>	DR REDDYS LABS LTD	<u>425.2MG;EQ 574.9MG BASE</u>	<u>A077701 001</u>	Mar 26, 2007
<u>AB</u>	! MYLAN PHARMS INC	<u>212.6MG;EQ 287.5MG BASE</u>	<u>A078183 001</u>	Mar 22, 2007
<u>AB</u>	!	<u>425.2MG;EQ 574.9MG BASE</u>	<u>A078183 002</u>	Mar 22, 2007

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS; OTIC

CIPRODEX

+! NOVARTIS PHARMS 0.3%; 0.1%  
CORP

N021537 001 Jul 18, 2003

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

CISATRACURIUM BESYLATE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A205873 001</u>	Jun 16, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203183 001</u>	Feb 26, 2015
<u>AP</u>	JIANGSU HENGRIUI MED	<u>EQ 2MG BASE/ML</u>	<u>A209334 001</u>	Aug 30, 2017
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200159 001</u>	Feb 03, 2012

CISATRACURIUM BESYLATE PRESERVATIVE FREE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A205872 001</u>	Jun 16, 2017
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A205872 002</u>	Jun 16, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A203182 001</u>	Feb 26, 2015
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A203182 002</u>	Feb 26, 2015
<u>AP</u>	JIANGSU HENGRIUI MED	<u>EQ 2MG BASE/ML</u>	<u>A204960 001</u>	Jan 27, 2017
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A204960 002</u>	Sep 19, 2017
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A200154 001</u>	Feb 03, 2012
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A200154 002</u>	Feb 03, 2012

NIMBEX

<u>AP</u>	+! ABBVIE	<u>EQ 2MG BASE/ML</u>	<u>N020551 001</u>	Dec 15, 1995
<u>AP</u>	<u>NIMBEX PRESERVATIVE FREE</u>	<u>EQ 2MG BASE/ML</u>	<u>N020551 003</u>	Dec 15, 1995
<u>AP</u>	+!	<u>EQ 10MG BASE/ML</u>	<u>N020551 002</u>	Dec 15, 1995

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

<u>AP</u>	ACCORD HLTHCARE	<u>1MG/ML</u>	<u>A206774 001</u>	Aug 18, 2015
<u>AP</u>	! FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A074735 001</u>	Jul 16, 1999
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A207323 001</u>	Mar 17, 2017
<u>AP</u>	+	<u>1MG/ML</u>	<u>N018057 004</u>	Nov 08, 1988
<u>AP</u>	HQ SPCLT PHARMA	<u>1MG/ML</u>	<u>A091062 001</u>	Apr 18, 2012
<u>AP</u>	MYLAN LABS LTD	<u>1MG/ML</u>	<u>A074656 001</u>	May 16, 2000
<u>AP</u>	PHARMACHEMIE BV	<u>1MG/ML</u>	<u>A075036 001</u>	Nov 07, 2000
<u>AP</u>	WEST-WARD PHARMS INT	<u>1MG/ML</u>		

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-95 (of 436)

CITALOPRAM HYDROBROMIDE

SOLUTION;ORAL

CITALOPRAM HYDROBROMIDE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE/5ML</u>	<u>A077812 001</u>	Aug 28, 2006
<u>AA</u>	HETERO LABS LTD III	<u>EQ 10MG BASE/5ML</u>	<u>A201450 001</u>	Dec 15, 2015
<u>AA</u>	SILARX	<u>EQ 10MG BASE/5ML</u>	<u>A077629 001</u>	Jun 15, 2006
<u>AA</u> !	WEST-WARD PHARMS INT	<u>EQ 10MG BASE/5ML</u>	<u>A077043 001</u>	Dec 13, 2004

TABLET;ORAL

CELEXA

<u>AB</u> +	FOREST LABS	<u>EQ 10MG BASE</u>	<u>N020822 001</u>	Apr 27, 2000
<u>AB</u> +		<u>EQ 20MG BASE</u>	<u>N020822 002</u>	Jul 17, 1998
<u>AB</u> +!		<u>EQ 40MG BASE</u>	<u>N020822 003</u>	Jul 17, 1998

CITALOPRAM HYDROBROMIDE

<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 10MG BASE</u>	<u>A077289 001</u>	Nov 30, 2006
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077289 002</u>	Nov 30, 2006
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077289 003</u>	Nov 30, 2006
<u>AB</u>	APOTEX INC	<u>EQ 10MG BASE</u>	<u>A077046 001</u>	Nov 24, 2004
<u>AB</u>	AUROBINDO	<u>EQ 10MG BASE</u>	<u>A077031 001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077031 002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077031 003</u>	Oct 28, 2004
<u>AB</u>	CIPLA LTD	<u>EQ 10MG BASE</u>	<u>A077044 001</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077044 002</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077044 003</u>	Nov 05, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>A077038 001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077038 002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077038 003</u>	Oct 28, 2004
<u>AB</u>	EPIC PHARMA	<u>EQ 10MG BASE</u>	<u>A077045 003</u>	Apr 29, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077045 002</u>	Apr 29, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077045 001</u>	Apr 29, 2005
<u>AB</u>	G AND W LABS INC	<u>EQ 10MG BASE</u>	<u>A077048 001</u>	Nov 16, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077048 002</u>	Nov 16, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077048 003</u>	Nov 16, 2004
<u>AB</u>	GLENMARK GENERICS	<u>EQ 10MG BASE</u>	<u>A077654 001</u>	Feb 27, 2009
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077654 002</u>	Feb 27, 2009
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077654 003</u>	Feb 27, 2009
<u>AB</u>	INVAGEN PHARMS	<u>EQ 10MG BASE</u>	<u>A077534 001</u>	Oct 03, 2006
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077534 002</u>	Oct 03, 2006
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077534 003</u>	Oct 03, 2006
<u>AB</u>	JUBILANT GENERICS	<u>EQ 10MG BASE</u>	<u>A205407 001</u>	Dec 23, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205407 002</u>	Dec 23, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205407 003</u>	Dec 23, 2015
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A077042 001</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077042 002</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077042 003</u>	Nov 05, 2004
<u>AB</u>	PLIVA	<u>EQ 10MG BASE</u>	<u>A077232 001</u>	Oct 31, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077232 002</u>	Oct 31, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077232 003</u>	Oct 31, 2005
<u>AB</u>	SUN PHARM INDNS INC	<u>EQ 10MG BASE</u>	<u>A077032 001</u>	Nov 12, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077032 002</u>	Nov 12, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077032 003</u>	Nov 12, 2004
<u>AB</u>	TORPHARM	<u>EQ 20MG BASE</u>	<u>A077046 002</u>	Nov 24, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077046 003</u>	Nov 24, 2004
<u>AB</u>	TORRENT PHARMS	<u>EQ 10MG BASE</u>	<u>A078216 001</u>	Mar 27, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078216 002</u>	Mar 27, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078216 003</u>	Mar 27, 2007

CITRIC ACID; GLUCONOLACTONE; MAGNESIUM CARBONATE

SOLUTION;IRRIGATION

RENACIDIN

+! UNITED GUARDIAN 6.602GM/100ML;198MG/100ML;3.177GM/100ML N019481 001 Oct 02, 1990

CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE

FOR SOLUTION;ORAL

PREPOPIK

+! FERRING PHARMS INC 12GM/PACKET;3.5GM/PACKET;10MG/PACKET N202535 001 Jul 16, 2012

SOLUTION;ORAL

CLENPIQ

+! FERRING PHARMS INC 12GM/160ML;3.5GM/160ML;10MG/160ML N209589 001 Nov 28, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-96 (of 436)

CITRIC ACID; UREA C-13

FOR SOLUTION, TABLET, FOR SOLUTION;ORAL  
 IDKIT:HP  
 +! EXALENZ BIOSCIENCE N/A, 4GM; 75MG, N/A

N021314 001 Dec 17, 2002

CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE

<u>AP</u>	!	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A076571 001</u> Apr 22, 2004
<u>AP</u>		MYLAN LABS LTD	<u>1MG/ML</u>	<u>A200510 001</u> Oct 06, 2011
<u>AP</u>		WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A075405 001</u> Feb 28, 2000

CLARITHROMYCIN

FOR SUSPENSION;ORAL

<u>BIAXIN</u>				
<u>AB</u>	+	ABBVIE	<u>125MG/5ML</u>	<u>N050698 001</u> Dec 23, 1993
<u>AB</u>	+!		<u>250MG/5ML</u>	<u>N050698 002</u> Dec 23, 1993

CLARITHROMYCIN

<u>AB</u>		SANDOZ	<u>125MG/5ML</u>	<u>A065283 002</u> Sep 04, 2007
<u>AB</u>			<u>250MG/5ML</u>	<u>A065283 003</u> Sep 04, 2007
<u>AB</u>		SUN PHARM INDs LTD	<u>125MG/5ML</u>	<u>A065382 001</u> Aug 30, 2007
<u>AB</u>			<u>250MG/5ML</u>	<u>A065382 002</u> Aug 30, 2007

TABLET;ORAL

<u>BIAXIN</u>				
<u>AB</u>	+!	ABBVIE	<u>250MG</u>	<u>N050662 001</u> Oct 31, 1991
<u>AB</u>	+!		<u>500MG</u>	<u>N050662 002</u> Oct 31, 1991

CLARITHROMYCIN

<u>AB</u>		ALLIED PHARMA INC	<u>250MG</u>	<u>A202710 001</u> Jun 10, 2013
<u>AB</u>			<u>500MG</u>	<u>A202710 002</u> Jun 10, 2013
<u>AB</u>		APOTEX CORP	<u>250MG</u>	<u>A065384 001</u> Aug 20, 2007
<u>AB</u>			<u>500MG</u>	<u>A065384 002</u> Aug 20, 2007
<u>AB</u>		AUROBINDO	<u>250MG</u>	<u>A065489 001</u> Jul 25, 2012
<u>AB</u>			<u>500MG</u>	<u>A065489 002</u> Jul 25, 2012
<u>AB</u>		HEC PHARM USA INC	<u>250MG</u>	<u>A203584 001</u> Sep 28, 2015
<u>AB</u>			<u>500MG</u>	<u>A203584 002</u> Sep 28, 2015
<u>AB</u>		SANDOZ	<u>250MG</u>	<u>A065144 001</u> Oct 18, 2005
<u>AB</u>			<u>500MG</u>	<u>A065136 001</u> Aug 25, 2005
<u>AB</u>		SUN PHARM INDs LTD	<u>250MG</u>	<u>A065174 001</u> Sep 24, 2004
<u>AB</u>			<u>500MG</u>	<u>A065174 002</u> Sep 24, 2004
<u>AB</u>		TEVA	<u>250MG</u>	<u>A065155 001</u> May 31, 2005
<u>AB</u>			<u>500MG</u>	<u>A065155 002</u> May 31, 2005
<u>AB</u>		WEST-WARD PHARMS INT	<u>250MG</u>	<u>A065178 002</u> May 25, 2004
<u>AB</u>			<u>500MG</u>	<u>A065178 001</u> May 25, 2004
<u>AB</u>		WOCKHARDT	<u>250MG</u>	<u>A065266 001</u> May 31, 2006
<u>AB</u>			<u>500MG</u>	<u>A065266 002</u> May 31, 2006

TABLET, EXTENDED RELEASE;ORAL

<u>CLARITHROMYCIN</u>				
<u>AB</u>		ACTAVIS LABS FL INC	<u>500MG</u>	<u>A065145 001</u> Jun 24, 2004
<u>AB</u>		ALLIED PHARMA INC	<u>500MG</u>	<u>A203243 001</u> Feb 29, 2016
<u>AB</u>		LUPIN LTD	<u>500MG</u>	<u>A202532 001</u> Sep 15, 2015
<u>AB</u>	!	MAYNE PHARMA	<u>500MG</u>	<u>A065154 001</u> May 18, 2005

CLEMASTINE FUMARATE

SYRUP;ORAL

<u>CLEMASTINE FUMARATE</u>				
<u>AA</u>	!	TEVA	<u>EQ 0.5MG BASE/5ML</u>	<u>A073399 001</u> Jun 30, 1994
<u>AA</u>		WOCKHARDT	<u>EQ 0.5MG BASE/5ML</u>	<u>A074863 001</u> Mar 13, 1998

TABLET;ORAL

CLEMASTINE FUMARATE		
!	TEVA	2.68MG

A073283 001 Jan 31, 1992

CLEVIDIPIINE

EMULSION; INTRAVENOUS

CLEVIPREX		
+	CHIESI USA INC	25MG/50ML (0.5MG/ML)
+		50MG/100ML (0.5MG/ML)
+		125MG/250ML (0.5MG/ML)

N022156 001 Aug 01, 2008  
 N022156 002 Aug 01, 2008  
 N022156 003 Nov 08, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PREScription DRUG PRODUCT LIST**

3-97 (of 436)

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

## CLEOCIN HYDROCHLORIDE

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>EO 75MG BASE</u>	<u>N050162</u> <u>001</u>
<u>AB</u>	+		<u>EO 150MG BASE</u>	<u>N050162</u> <u>002</u>
<u>AB</u>	+		<u>EO 300MG BASE</u>	<u>N050162</u> <u>003</u> Apr 14, 1988
<b>CLINDAMYCIN HYDROCHLORIDE</b>				
<u>AB</u>		AUROBINDO PHARMA	<u>EO 150MG BASE</u>	<u>A065442</u> <u>001</u> Aug 26, 2009
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A065442</u> <u>002</u> Aug 26, 2009
<u>AB</u>		EPIC PHARMA LLC	<u>EO 150MG BASE</u>	<u>A065194</u> <u>001</u> Mar 22, 2004
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A065194</u> <u>002</u> Mar 22, 2004
<u>AB</u>		G AND W LABS INC	<u>EO 150MG BASE</u>	<u>A063029</u> <u>001</u> Sep 20, 1989
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A063029</u> <u>002</u> Aug 05, 2005
<u>AB</u>		LANNETT	<u>EO 75MG BASE</u>	<u>A065243</u> <u>002</u> Aug 12, 2005
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A065243</u> <u>001</u> Aug 12, 2005
<u>AB</u>		SUN PHARM INDs LTD	<u>EO 150MG BASE</u>	<u>A065061</u> <u>001</u> Feb 02, 2001
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A065061</u> <u>002</u> Feb 02, 2001
<u>AB</u>		WATSON LABS	<u>EO 150MG BASE</u>	<u>A063083</u> <u>001</u> Jul 31, 1991
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A063083</u> <u>002</u> Mar 18, 2003
<u>AB</u>		ZYDUS PHARMS USA	<u>EO 75MG BASE</u>	<u>A065217</u> <u>001</u> Jan 31, 2005
<u>AB</u>			<u>EO 150MG BASE</u>	<u>A065217</u> <u>002</u> Jan 31, 2005
<u>AB</u>			<u>EO 300MG BASE</u>	<u>A065217</u> <u>003</u> Jan 31, 2005
<b>CLINDAMYCIN HYDROCHLORIDE</b>				
<u>AB</u>		LANNETT	<u>EO 150MG BASE</u>	<u>A065243</u> <u>003</u> Aug 12, 2005

## CLINDAMYCIN PALMITATE HYDROCHLORIDE

**FOR SOLUTION; ORAL**

## CLEOCIN

<u>AA</u>	!	PHARMACIA AND UPJOHN	<u>EQ 75MG BASE/5ML</u>	<u>A062644</u>	<u>001</u>	Apr 07, 1986
<u>CLINDAMYCIN PALMITATE HYDROCHLORIDE</u>						
<u>AA</u>		AMNEAL PHARMS	<u>EQ 75MG BASE/5ML</u>	<u>A203513</u>	<u>001</u>	Mar 13, 2014
<u>AA</u>		AUROBINDO PHARMA LTD	<u>EQ 75MG BASE/5ML</u>	<u>A202409</u>	<u>001</u>	Apr 30, 2013
<u>AA</u>		LYNE	<u>EQ 75MG BASE/5ML</u>	<u>A201821</u>	<u>001</u>	Aug 28, 2012
<u>AA</u>		MYLAN PHARMS INC	<u>EQ 75MG BASE/5ML</u>	<u>A203063</u>	<u>001</u>	May 25, 2016
<u>AA</u>		ORIT LABS LLC	<u>EQ 75MG BASE/5ML</u>	<u>A206958</u>	<u>001</u>	May 05, 2017
<u>AA</u>		PADDOCK LLC	<u>EQ 75MG BASE/5ML</u>	<u>A090902</u>	<u>001</u>	Jul 07, 2010

## CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

## CLINDAMYCIN PHOSPHATE

AT PERRIGO UK FINCO 1% A090785\_001 Mar 31, 2010  
EVOCLIN

AT +! MYLAN PHARMS INC 1% N050801\_001 Oct 22, 2004  
CREAM;VAGINAL

## CLEOCIN

**B** +! PHARMACIA AND  
UP TOWN

CL TNDAM

**CLINDAMICIN PHOSPHATE**      **AB FOUGERA PHARMS EQ 2% BASE**      **A065139 001 Dec 27, 2004**

+1 PERBTGO PHARMA

## GEL; TOPICAL

## CLEOCIN T

AB +! PHARMACIA AND EQ 1% BASE N050615 001 Jan 07, 1987  
UP JOHN

## CLINDAMYCIN PHOSPHATE

**b** FOUND

CLINDAGE II

BT +! PRECISION DERMAT EQ 1% BASE N050782 001 Nov 27, 2000  
INJECTABLE; INJECTION

## **CLEOCIN PHOSPHATE**

**P** PHARMACIA AND  
UP-SKIN

UPJOHN

AP +! EQ 150MG BASE/ML NU50441 001  
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

**THE PHARMACEUTICAL  
JOURNAL**

P + !

+!

CLINDAMYCIN PHOSPHATE

<u>AP</u>	ALVOGEN INC	<u>EO 150MG BASE/ML</u>	<u>A062800 001</u>	Jul 24, 1987
<u>AP</u>		<u>EO 150MG BASE/ML</u>	<u>A062801 001</u>	Jul 24, 1987

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-98 (of 436)

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 150MG BASE/ML</u>	<u>A062943 001</u>	Sep 29, 1988
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A065346 001</u>	Mar 29, 2007
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A065347 001</u>	May 09, 2007
<u>AP</u>	MYLAN LABS LTD	<u>EQ 150MG BASE/ML</u>	<u>A204748 001</u>	Oct 10, 2017
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A204749 001</u>	Oct 10, 2017
<u>AP</u>	SAGENT PHARMS	<u>EQ 150MG BASE/ML</u>	<u>A090108 001</u>	Sep 30, 2011
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A090109 001</u>	Sep 30, 2011
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 150MG BASE/ML</u>	<u>A062889 001</u>	Apr 25, 1988
<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>A065206 001</u>	Sep 24, 2004

CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER

<u>AP</u>	AKORN INC	<u>EQ 6MG BASE/ML</u>	<u>A203048 001</u>	Apr 04, 2013
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A203048 002</u>	Apr 04, 2013
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A203048 003</u>	Apr 04, 2013
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 6MG BASE/ML</u>	<u>A208084 001</u>	Jun 28, 2017
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A208084 002</u>	Jun 28, 2017
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A208084 003</u>	Jun 28, 2017
<u>AP</u>	SANDOZ INC	<u>EQ 6MG BASE/ML</u>	<u>A201692 001</u>	May 31, 2012
<u>AP</u>		<u>EQ 12MG BASE/ML</u>	<u>A201692 002</u>	May 31, 2012
<u>AP</u>		<u>EQ 18MG BASE/ML</u>	<u>A201692 003</u>	May 31, 2012

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% +! ABRAXIS PHARM LOTION;TOPICAL

EQ 900MG BASE/100ML

N050635 001 Dec 22, 1989

CLEOCIN T

<u>AB</u>	+! PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050600 001</u>	May 31, 1989
-----------	-------------------------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB</u>	FOUGERA PHARMS SOLUTION;IV (INFUSION)	<u>EQ 1% BASE</u>	<u>A065067 001</u>	Jan 31, 2002
	CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE			
+!	BAXTER HLTHCARE CORP	EQ 300MG BASE/50ML (EQ 6MG BASE/ML)	N208083 001	Apr 20, 2017
+!		EQ 600MG BASE/50ML (EQ 12MG BASE/ML)	N208083 002	Apr 20, 2017
+!		EQ 900MG BASE/50ML (EQ 18MG BASE/ML)	N208083 003	Apr 20, 2017

SOLUTION;TOPICAL

CLEOCIN T

<u>AT</u>	+! PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050537 001</u>	
-----------	-------------------------	-------------------	--------------------	--

CLINDA-DERM

<u>AT</u>	PADDOCK LLC	<u>EQ 1% BASE</u>	<u>A063329 001</u>	Sep 30, 1992
-----------	-------------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AT</u>	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<u>A065254 001</u>	Feb 14, 2006
<u>AT</u>	FOUGERA PHARMS INC	<u>EQ 1% BASE</u>	<u>A064159 001</u>	Jun 05, 1997
<u>AT</u>	G AND W LABS INC	<u>EQ 1% BASE</u>	<u>A062811 001</u>	Sep 01, 1988
<u>AT</u>	PERRIGO NEW YORK	<u>EQ 1% BASE</u>	<u>A064050 001</u>	Nov 30, 1995
<u>AT</u>	TARO PHARM IND	<u>EQ 1% BASE</u>	<u>A065184 001</u>	Mar 31, 2004
<u>AT</u>	TELIGENT PHARMA INC	<u>EQ 1% BASE</u>	<u>A206945 001</u>	Dec 30, 2016
<u>AT</u>	VINTAGE PHARMS	<u>EQ 1% BASE</u>	<u>A203343 001</u>	May 29, 2015
<u>AT</u>	WOCKHARDT	<u>EQ 1% BASE</u>	<u>A063304 001</u>	Jul 15, 1997

SUPPOSITORIY;VAGINAL

CLEOCIN

+!	PHARMACIA AND UPJOHN	100MG	N050767 001	Aug 13, 1999
----	----------------------	-------	-------------	--------------

SWAB;TOPICAL

CLEOCIN

<u>AT</u>	+ PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N050537 002</u>	Feb 22, 1994
-----------	------------------------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AT</u>	AKORN	<u>EQ 1% BASE</u>	<u>A065513 001</u>	Jun 17, 2010
<u>AT</u>	PERRIGO NEW YORK	<u>EQ 1% BASE</u>	<u>A065049 001</u>	May 25, 2000

CLINDETS

<u>AT</u>	PERRIGO CO	<u>EQ 1% BASE</u>	<u>A064136 001</u>	Sep 30, 1996
-----------	------------	-------------------	--------------------	--------------

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL;TOPICAL

CLINDAMYCIN PHOSPHATE AND TRETINOIN

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>1.2%:0.025%</u>	<u>A202564 001</u>	Jun 12, 2015
-----------	----------------------	--------------------	--------------------	--------------

ZIANA

<u>AB</u>	+! MEDICIS	<u>1.2%:0.025%</u>	<u>N050802 001</u>	Nov 07, 2006
-----------	------------	--------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-99 (of 436)

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL;TOPICAL  
 VELTIN  
 BX +! AQUA PHARMS LLC 1.2%;0.025% N050803 001 Jul 16, 2010

CLOBAZAM

SUSPENSION;ORAL  
 ONFI  
 +! LUNDBECK PHARMS LLC 2.5MG/ML N203993 001 Dec 14, 2012  
 TABLET;ORAL  
 ONFI  
 + LUNDBECK PHARMS LLC 10MG N202067 002 Oct 21, 2011  
 +! 20MG N202067 003 Oct 21, 2011

CLOBETASOL PROPIONATE

AEROSOL, FOAM;TOPICAL

CLOBETASOL PROPIONATE  
AB1 INGENUS PHARMS LLC 0.05% A206805 001 Jul 31, 2017  
AB1 PERRIGO ISRAEL 0.05% A077763 001 Mar 10, 2008  
OLUX  
AB1 +! MYLAN PHARMS INC 0.05% N021142 001 May 26, 2000  
CLOBETASOL PROPIONATE  
AB2 PERRIGO ISRAEL 0.05% A201402 001 Aug 14, 2012  
OLUX E  
AB2 +! MYLAN PHARMS INC 0.05% N022013 001 Jan 12, 2007  
 CREAM;TOPICAL

CLOBETASOL PROPIONATE  
AB1 FOUGERA PHARMS INC 0.05% A074392 001 Sep 30, 1996  
AB1 G AND W LABS INC 0.05% A074139 001 Aug 03, 1994  
AB1 MYLAN PHARMS INC 0.05% A075338 001 Feb 09, 2001  
AB1 TARO 0.05% A074249 001 Jul 08, 1996

CORMAX  
AB1 ! HI TECH PHARMA 0.05% A074220 001 May 16, 1997  
CLOBETASOL PROPIONATE (EMOLLIENT)  
AB2 ! FOUGERA PHARMS 0.05% A075430 001 May 26, 1999  
AB2 TARO 0.05% A075633 001 May 17, 2000  
AB2 TELIGENT PHARMA INC 0.05% A209411 001 Aug 21, 2017  
EMBELINE E  
AB2 HI TECH PHARMA 0.05% A075325 001 Dec 24, 1998

IMPOYZ  
 +! PROMIUS PHARMA LLC 0.025% N209483 001 Nov 28, 2017

GEL;TOPICAL  
CLOBETASOL PROPIONATE  
AB ! FOUGERA PHARMS 0.05% A075368 001 Feb 15, 2000  
AB PERRIGO CO 0.05% A075027 001 Oct 31, 1997  
AB TARO 0.05% A075279 001 May 28, 1999  
AB TELIGENT PHARMA INC 0.05% A208881 001 Mar 06, 2017

EMBELINE  
AB HI TECH PHARMA 0.05% A076141 001 Apr 12, 2002  
 LOTION;TOPICAL  
CLOBETASOL PROPIONATE  
AB ACTAVIS MID 0.05% A078223 001 Dec 04, 2008  
 ATLANTIC

AB LUPIN LTD 0.05% A209147 001 Sep 22, 2017  
AB TARO 0.05% A200302 001 Jul 02, 2012  
AB TELIGENT PHARMA INC 0.05% A208667 001 Nov 29, 2016

CLOBEX  
AB +! GALDERMA LABS LP 0.05% N021535 001 Jul 24, 2003  
 OINTMENT;TOPICAL  
CLOBETASOL PROPIONATE  
AB ! FOUGERA PHARMS 0.05% A074407 001 Feb 23, 1996  
AB G AND W LABS INC 0.05% A074089 001 Feb 16, 1994  
AB GLENMARK PHARMS 0.05% A208933 001 Mar 20, 2017  
AB MYLAN PHARMS INC 0.05% A075057 001 Aug 12, 1998  
AB TARO 0.05% A074248 001 Jul 12, 1996  
AB ZYDUS PHARMS USA 0.05% A210199 001 Oct 27, 2017  
 INC

EMBELINE  
AB HI TECH PHARMA 0.05% A074221 001 Mar 31, 1995  
 SHAMPOO;TOPICAL  
CLOBETASOL PROPIONATE  
AB ACTAVIS MID 0.05% A078854 001 Jun 07, 2011  
 ATLANTIC  
AB HI-TECH PHARMACAL 0.05% A209871 001 Oct 27, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-100 (of 436)

CLOBETASOL PROPIONATE

SHAMPOO;TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	PERRIGO ISRAEL	<u>0.05%</u>	<u>A090974 001</u> Aug 09, 2012
-----------	----------------	--------------	---------------------------------

CLOBEX

<u>AB</u>	+! GALDERMA LABS SOLUTION;TOPICAL	<u>0.05%</u>	<u>N021644 001</u> Feb 05, 2004
-----------	-----------------------------------	--------------	---------------------------------

CLOBETASOL PROPIONATE

<u>AT</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A075391 001</u> Feb 08, 1999
<u>AT</u>	G & W LABS INC	<u>0.05%</u>	<u>A074331 001</u> Dec 15, 1995
<u>AT</u>	MACLEODS PHARMS LTD	<u>0.05%</u>	<u>A209361 001</u> Oct 25, 2017
<u>AT</u>	NOVEL LABS INC	<u>0.05%</u>	<u>A206075 001</u> Nov 23, 2015
<u>AT</u>	TARO	<u>0.05%</u>	<u>A075224 001</u> Nov 16, 1998
<u>AT</u>		<u>0.05%</u>	<u>A075363 001</u> Dec 29, 2000
<u>AT</u>	TOLMAR	<u>0.05%</u>	<u>A076977 001</u> Aug 05, 2005
<u>AT</u>	WOCKHARDT BIO AG	<u>0.05%</u>	<u>A075205 001</u> Nov 13, 1998

EMBELINE

<u>AT</u>	+! HI TECH PHARMA SPRAY;TOPICAL	<u>0.05%</u>	<u>A074222 001</u> Dec 06, 1995
-----------	---------------------------------	--------------	---------------------------------

CLOBETASOL PROPIONATE

<u>AT</u>	AKORN	<u>0.05%</u>	<u>A207218 001</u> Apr 28, 2017
<u>AT</u>	PADDOCK LLC	<u>0.05%</u>	<u>A090898 001</u> Jun 16, 2011
<u>AT</u>	ZYDUS PHARMS USA INC	<u>0.05%</u>	<u>A206378 001</u> Feb 16, 2017

CLOBEX

<u>AT</u>	+! GALDERMA LABS LP	<u>0.05%</u>	<u>N021835 001</u> Oct 27, 2005
-----------	---------------------	--------------	---------------------------------

CLOCORTOLONE PIVALATE

CREAM;TOPICAL

CLODERM

+! PROMIUS PHARMA LLC

0.1%

N017765 001

CLOFARABINE

INJECTABLE;IV (INFUSION)

CLOFARABINE

<u>AP</u>	ABON PHARMS LLC	<u>20MG/20ML (1MG/ML)</u>	<u>A204029 001</u> May 09, 2017
<u>AP</u>	AMNEAL PHARMS CO	<u>20MG/20ML (1MG/ML)</u>	<u>A208857 001</u> Nov 06, 2017
<u>AP</u>	DR REDDYS LABS LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A205375 001</u> Nov 06, 2017
<u>AP</u>	MSN LABS PVT LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A209775 001</u> Dec 06, 2017
<u>AP</u>	MYLAN LABS LTD	<u>20MG/20ML (1MG/ML)</u>	<u>A208860 001</u> Nov 06, 2017

CLOLAR

<u>AP</u>	+! GENZYME	<u>20MG/20ML (1MG/ML)</u>	<u>N021673 001</u> Dec 28, 2004
-----------	------------	---------------------------	---------------------------------

CLOMIPHENE CITRATE

TABLET;ORAL

CLOMID

<u>AB</u>	+! SANOFI AVENTIS US	<u>50MG</u>	<u>N016131 002</u>
-----------	----------------------	-------------	--------------------

CLOMIPHENE CITRATE

<u>AB</u>	PAR PHARM	<u>50MG</u>	<u>A075528 001</u> Aug 30, 1999
-----------	-----------	-------------	---------------------------------

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

ANAFRANIL

<u>AB</u>	+! SPECGX LLC	<u>25MG</u>	<u>N019906 001</u> Dec 29, 1989
<u>AB</u>	+	<u>50MG</u>	<u>N019906 002</u> Dec 29, 1989
<u>AB</u>	+	<u>75MG</u>	<u>N019906 003</u> Dec 29, 1989

CLOMIPRAMINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>25MG</u>	<u>A074947 001</u> Apr 30, 1998
<u>AB</u>		<u>50MG</u>	<u>A074947 002</u> Apr 30, 1998
<u>AB</u>		<u>75MG</u>	<u>A074947 003</u> Apr 30, 1998
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A074364 001</u> Mar 29, 1996
<u>AB</u>		<u>25MG</u>	<u>A074953 001</u> Jun 25, 1997
<u>AB</u>		<u>50MG</u>	<u>A074364 002</u> Mar 29, 1996
<u>AB</u>		<u>50MG</u>	<u>A074953 002</u> Jun 25, 1997
<u>AB</u>		<u>75MG</u>	<u>A074364 003</u> Mar 29, 1996
<u>AB</u>		<u>75MG</u>	<u>A074953 003</u> Jun 25, 1997
<u>AB</u>	TARO	<u>25MG</u>	<u>A074694 001</u> Dec 31, 1996
<u>AB</u>		<u>50MG</u>	<u>A074694 002</u> Dec 31, 1996
<u>AB</u>		<u>75MG</u>	<u>A074694 003</u> Dec 31, 1996
<u>AB</u>	TEVA	<u>25MG</u>	<u>A074958 001</u> Aug 26, 1997
<u>AB</u>		<u>50MG</u>	<u>A074958 002</u> Aug 26, 1997
<u>AB</u>		<u>75MG</u>	<u>A074958 003</u> Aug 26, 1997
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A208961 001</u> Dec 27, 2017
<u>AB</u>		<u>50MG</u>	<u>A208961 002</u> Dec 27, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-101 (of 436)

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

AB 75MG A208961 003 Dec 27, 2017

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

<u>AB</u>	ACCORD HLTHCARE	<u>0 .5MG</u>	<u>A077147 001</u> May 02, 2005
<u>AB</u>		<u>1MG</u>	<u>A077147 002</u> May 02, 2005
<u>AB</u>		<u>2MG</u>	<u>A077147 003</u> May 02, 2005
<u>AB</u>	ACTAVIS ELIZABETH	<u>0 .5MG</u>	<u>A074869 001</u> Oct 31, 1996
<u>AB</u>		<u>1MG</u>	<u>A074869 002</u> Oct 31, 1996
<u>AB</u>		<u>2MG</u>	<u>A074869 003</u> Oct 31, 1996
<u>AB</u>	MYLAN	<u>0 .5MG</u>	<u>A075150 001</u> Oct 05, 1998
<u>AB</u>		<u>1MG</u>	<u>A075150 002</u> Oct 05, 1998
<u>AB</u>		<u>2MG</u>	<u>A075150 003</u> Oct 05, 1998
<u>AB</u>	PRINSTON INC	<u>0 .5MG</u>	<u>A077856 001</u> Jun 28, 2006
<u>AB</u>		<u>1MG</u>	<u>A077856 002</u> Jun 28, 2006
<u>AB</u>		<u>2MG</u>	<u>A077856 003</u> Jun 28, 2006
<u>AB</u>	SANDOZ	<u>0 .5MG</u>	<u>A074979 001</u> Aug 29, 1997
<u>AB</u>		<u>1MG</u>	<u>A074979 002</u> Aug 29, 1997
<u>AB</u>		<u>2MG</u>	<u>A074979 003</u> Aug 29, 1997
<u>AB</u>	SUN PHARM INDs INC	<u>0 .5MG</u>	<u>A075423 001</u> Apr 27, 2001
<u>AB</u>		<u>1MG</u>	<u>A075423 002</u> Apr 27, 2001
<u>AB</u>		<u>2MG</u>	<u>A075423 003</u> Apr 27, 2001
<u>AB</u>	TEVA	<u>0 .5MG</u>	<u>A074569 001</u> Sep 10, 1996
<u>AB</u>		<u>1MG</u>	<u>A074569 002</u> Sep 10, 1996
<u>AB</u>		<u>2MG</u>	<u>A074569 003</u> Sep 10, 1996
<u>AB</u>	WATSON LABS	<u>0 .5MG</u>	<u>A074964 001</u> Dec 30, 1997
<u>AB</u>		<u>1MG</u>	<u>A074964 002</u> Dec 30, 1997
<u>AB</u>		<u>2MG</u>	<u>A074964 003</u> Dec 30, 1997
<u>KLONOPIN</u>			
<u>AB</u>	+ ROCHE	<u>0 .5MG</u>	<u>N017533 001</u>
<u>AB</u>	+!	<u>1MG</u>	<u>N017533 002</u>
<u>AB</u>	+	<u>2MG</u>	<u>N017533 003</u>

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	BARR	<u>0 .125MG</u>	<u>A077194 001</u> Aug 10, 2005
<u>AB</u>		<u>0 .25MG</u>	<u>A077194 002</u> Aug 10, 2005
<u>AB</u>		<u>0 .5MG</u>	<u>A077194 003</u> Aug 10, 2005
<u>AB</u>		<u>1MG</u>	<u>A077194 004</u> Aug 10, 2005
<u>AB</u>		<u>2MG</u>	<u>A077194 005</u> Aug 10, 2005
<u>AB</u>	PAR PHARM	<u>0 .125MG</u>	<u>A077171 001</u> Aug 03, 2005
<u>AB</u>		<u>0 .25MG</u>	<u>A077171 002</u> Aug 03, 2005
<u>AB</u>		<u>0 .5MG</u>	<u>A077171 003</u> Aug 03, 2005
<u>AB</u>	!	<u>1MG</u>	<u>A077171 004</u> Aug 03, 2005
<u>AB</u>		<u>2MG</u>	<u>A077171 005</u> Aug 03, 2005
<u>AB</u>	SUN PHARM INDs INC	<u>0 .125MG</u>	<u>A078654 001</u> Aug 27, 2014
<u>AB</u>		<u>0 .25MG</u>	<u>A078654 002</u> Aug 27, 2014
<u>AB</u>		<u>0 .5MG</u>	<u>A078654 003</u> Aug 27, 2014
<u>AB</u>		<u>1MG</u>	<u>A078654 004</u> Aug 27, 2014
<u>AB</u>		<u>2MG</u>	<u>A078654 005</u> Aug 27, 2014

CLONIDINE

FILM, EXTENDED RELEASE; TRANSDERMAL

CATAPRES-TTS-1

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>0 .1MG/24HR</u>	<u>N018891 001</u> Oct 10, 1984
-----------	------------------------	--------------------	---------------------------------

CATAPRES-TTS-2

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>0 .2MG/24HR</u>	<u>N018891 002</u> Oct 10, 1984
-----------	------------------------	--------------------	---------------------------------

CATAPRES-TTS-3

<u>AB</u>	+! BOEHRINGER INGELHEIM	<u>0 .3MG/24HR</u>	<u>N018891 003</u> Oct 10, 1984
-----------	-------------------------	--------------------	---------------------------------

CLONIDINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>0 .1MG/24HR</u>	<u>A090873 001</u> May 06, 2014
<u>AB</u>		<u>0 .2MG/24HR</u>	<u>A090873 002</u> May 06, 2014
<u>AB</u>		<u>0 .3MG/24HR</u>	<u>A090873 003</u> May 06, 2014
<u>AB</u>	AVEVA	<u>0 .1MG/24HR</u>	<u>A076157 001</u> Aug 18, 2009
<u>AB</u>		<u>0 .2MG/24HR</u>	<u>A076157 002</u> Aug 18, 2009
<u>AB</u>		<u>0 .3MG/24HR</u>	<u>A076157 003</u> Aug 18, 2009
<u>AB</u>	MAYNE PHARMA	<u>0 .1MG/24HR</u>	<u>A079090 001</u> Aug 20, 2010
<u>AB</u>		<u>0 .2MG/24HR</u>	<u>A079090 002</u> Aug 20, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-102 (of 436)

CLONIDINE

FILM, EXTENDED RELEASE; TRANSDERMAL

CLONIDINE

<u>AB</u>		<u>0.3MG/24HR</u>	<u>A079090 003</u>	Aug 20, 2010
<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.1MG/24HR</u>	<u>A076166 001</u>	Jul 16, 2010
<u>AB</u>		<u>0.2MG/24HR</u>	<u>A076166 002</u>	Jul 16, 2010
<u>AB</u>		<u>0.3MG/24HR</u>	<u>A076166 003</u>	Jul 16, 2010

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CLONIDINE HYDROCHLORIDE

<u>AP</u>	EXELA PHARMA SCS LLC	<u>1MG/10ML (0.1MG/ML)</u>	<u>A203167 001</u>	Oct 29, 2013
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A203167 002</u>	Oct 29, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200673 001</u>	Jul 08, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200673 002</u>	Jul 08, 2011
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/10ML (0.1MG/ML)</u>	<u>A200300 001</u>	Jan 26, 2011
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A200300 002</u>	Jan 26, 2011
<u>AP</u>	LUITPOLD	<u>1MG/10ML (0.1MG/ML)</u>	<u>A091104 001</u>	Oct 08, 2009
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A091104 002</u>	Oct 08, 2009
<u>AP</u>	ZYDUS PHARMS USA INC	<u>1MG/10ML (0.1MG/ML)</u>	<u>A202601 001</u>	Feb 20, 2014
<u>AP</u>		<u>5MG/10ML (0.5MG/ML)</u>	<u>A202601 002</u>	Feb 20, 2014
<b>DURACLON</b>				
<u>AP</u>	+ MYLAN INSTITUTIONAL	<u>1MG/10ML (0.1MG/ML)</u>	<u>N020615 001</u>	Oct 02, 1996
<u>AP</u>	+!	<u>5MG/10ML (0.5MG/ML)</u>	<u>N020615 002</u>	Apr 27, 1999

TABLET; ORAL

CATAPRES

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>0.1MG</u>	<u>N017407 001</u>
<u>AB</u>	+		<u>0.2MG</u>	<u>N017407 002</u>
<u>AB</u>	+!		<u>0.3MG</u>	<u>N017407 003</u>

CLONIDINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A070974 001</u>	Dec 16, 1986
<u>AB</u>		<u>0.2MG</u>	<u>A070975 001</u>	Dec 16, 1986
<u>AB</u>		<u>0.3MG</u>	<u>A070976 001</u>	Dec 16, 1986
<u>AB</u>	ALEMBIC PHARMS LTD	<u>0.1MG</u>	<u>A091368 001</u>	Dec 06, 2011
<u>AB</u>		<u>0.2MG</u>	<u>A091368 002</u>	Dec 06, 2011
<u>AB</u>		<u>0.3MG</u>	<u>A091368 003</u>	Dec 06, 2011
<u>AB</u>	FRONTIDA BIOPHARM	<u>0.1MG</u>	<u>A070925 001</u>	Sep 04, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070924 001</u>	Sep 04, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070923 001</u>	Sep 04, 1987
<u>AB</u>	IMPAX LABS	<u>0.1MG</u>	<u>A078099 001</u>	Aug 27, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078099 002</u>	Aug 27, 2009
<u>AB</u>		<u>0.3MG</u>	<u>A078099 003</u>	Aug 27, 2009
<u>AB</u>	MYLAN	<u>0.1MG</u>	<u>A070317 002</u>	Jul 09, 1987
<u>AB</u>		<u>0.2MG</u>	<u>A070317 003</u>	Jun 09, 1987
<u>AB</u>		<u>0.3MG</u>	<u>A070317 001</u>	Jun 09, 1987
<u>AB</u>	PRINSTON INC	<u>0.1MG</u>	<u>A077901 001</u>	Mar 09, 2007
<u>AB</u>		<u>0.2MG</u>	<u>A077901 002</u>	Mar 09, 2007
<u>AB</u>		<u>0.3MG</u>	<u>A077901 003</u>	Mar 09, 2007
<u>AB</u>	SUN PHARM INDs INC	<u>0.1MG</u>	<u>A090329 001</u>	Jul 03, 2014
<u>AB</u>		<u>0.2MG</u>	<u>A090329 002</u>	Jul 03, 2014
<u>AB</u>		<u>0.3MG</u>	<u>A090329 003</u>	Jul 03, 2014
<u>AB</u>	UNICHEM	<u>0.1MG</u>	<u>A078895 001</u>	Aug 26, 2009
<u>AB</u>		<u>0.2MG</u>	<u>A078895 002</u>	Aug 26, 2009
<u>AB</u>		<u>0.3MG</u>	<u>A078895 003</u>	Aug 26, 2009
<u>AB</u>	YUNG SHIN PHARM	<u>0.1MG</u>	<u>A202297 001</u>	Jun 13, 2013
<u>AB</u>		<u>0.2MG</u>	<u>A202297 002</u>	Jun 13, 2013
<u>AB</u>		<u>0.3MG</u>	<u>A202297 003</u>	Jun 13, 2013
	CHARTWELL MOLECULES	0.1MG	A071785 002	Apr 05, 1988
		0.2MG	A071785 003	Apr 05, 1988

TABLET, EXTENDED RELEASE; ORAL

CLONIDINE HYDROCHLORIDE

<u>AB1</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A203320 001</u>	May 15, 2015
<u>AB1</u>	AJANTA PHARMA LTD	<u>0.1MG</u>	<u>A209686 001</u>	Nov 20, 2017
<u>AB1</u>	AMNEAL PHARMS NY	<u>0.1MG</u>	<u>A210052 001</u>	Nov 20, 2017
<u>AB1</u>	LUPIN LTD	<u>0.1MG</u>	<u>A209285 001</u>	Oct 23, 2017
<u>AB1</u>	XIAMEN LP PHARM CO	<u>0.1MG</u>	<u>A209757 001</u>	Nov 20, 2017
<b>CLONIDINE HYDROCHLORIDE</b>				
<u>AB1</u>	ANCHEM PHARMS	<u>0.1MG</u>	<u>A202984 001</u>	Sep 30, 2013
<b>KAPVAY</b>				
<u>AB1</u>	+! CONCORDIA PHARMS INC	<u>0.1MG</u>	<u>N022331 003</u>	Sep 28, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-103 (of 436)

CLONIDINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CLONIDINE HYDROCHLORIDE

<u>AB2</u>	ACTAVIS ELIZABETH	<u>0.1MG</u>	<u>A202792 001</u> May 15, 2015
------------	-------------------	--------------	---------------------------------

CLOPIDOGREL BISULFATE

TABLET; ORAL

CLOPIDOGREL BISULFATE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 75MG BASE</u>	<u>A202925 001</u> Mar 27, 2013
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202925 002</u> Mar 27, 2013
<u>AB</u>	ACME LABS	<u>EQ 75MG BASE</u>	<u>A078004 001</u> May 17, 2012
<u>AB</u>	AMNEAL PHARMS	<u>EQ 75MG BASE</u>	<u>A203751 001</u> Apr 11, 2014
<u>AB</u>	APOTEX INC	<u>EQ 75MG BASE</u>	<u>A076274 001</u> May 17, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076274 002</u> Mar 04, 2014
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 75MG BASE</u>	<u>A090540 001</u> May 17, 2012
<u>AB</u>	CSPC OUYI PHARM CO	<u>EQ 75MG BASE</u>	<u>A204359 001</u> Feb 02, 2017
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 75MG BASE</u>	<u>A076273 001</u> Jan 14, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 300MG BASE</u>	<u>A091023 001</u> May 17, 2012
<u>AB</u>	GATE PHARMS	<u>EQ 300MG BASE</u>	<u>A091216 001</u> May 17, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A202928 001</u> Feb 10, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 75MG BASE</u>	<u>A077665 001</u> May 17, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077665 002</u> May 17, 2012
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 75MG BASE</u>	<u>A204165 001</u> Sep 15, 2014
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204165 002</u> Sep 15, 2014
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 75MG BASE</u>	<u>A078133 001</u> Jun 10, 2013
<u>AB</u>	SUN PHARMA GLOBAL	<u>EQ 75MG BASE</u>	<u>A090494 001</u> May 17, 2012
<u>AB</u>	TEVA	<u>EQ 75MG BASE</u>	<u>A076999 001</u> May 17, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 300MG BASE</u>	<u>A090625 001</u> May 17, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A090844 001</u> May 17, 2012
<u>AB</u>	WOCKHARDT LTD	<u>EQ 75MG BASE</u>	<u>A202266 001</u> Aug 14, 2012
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202266 002</u> Nov 20, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 75MG BASE</u>	<u>A201686 001</u> Oct 10, 2012

PLAVIX

<u>AB</u> + SANOFI AVENTIS US	<u>EQ 75MG BASE</u>	<u>N020839 001</u> Nov 17, 1997
<u>AB</u> +!	<u>EQ 300MG BASE</u>	<u>N020839 002</u> Sep 20, 2007

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM

AUROLIFE PHARMA LLC	3.75MG	A072112 002 Aug 11, 2017
	7.5MG	A072112 003 Aug 11, 2017

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

<u>AB</u> MYLAN	<u>3.75MG</u>	<u>A071858 002</u> Jul 17, 1987
<u>AB</u>	<u>7.5MG</u>	<u>A071858 003</u> Jul 17, 1987
<u>AB</u>	<u>15MG</u>	<u>A071858 001</u> Jul 17, 1987
<u>AB</u> TARO PHARM	<u>3.75MG</u>	<u>A075731 003</u> Apr 27, 2000
<u>AB</u>	<u>7.5MG</u>	<u>A075731 002</u> Apr 27, 2000
<u>AB</u>	<u>15MG</u>	<u>A075731 001</u> Apr 27, 2000
<u>AB</u> GEN-XENE		
<u>AB</u> ALRA	<u>3.75MG</u>	<u>A071787 001</u> Apr 26, 1988
<u>AB</u>	<u>7.5MG</u>	<u>A071788 001</u> Apr 26, 1988
<u>AB</u>	<u>15MG</u>	<u>A071789 001</u> Apr 26, 1988
<u>AB</u> TRANXENE		
<u>AB</u> + RECORDATI RARE	<u>7.5MG</u>	<u>N017105 007</u>
<u>AB</u> +!	<u>15MG</u>	<u>N017105 008</u>

CLOTRIMAZOLE

CREAM; TOPICAL

CLOTRIMAZOLE

<u>AB</u> FOUGERA PHARMS	<u>1%</u>	<u>A078338 001</u> Sep 02, 2008
<u>AB</u> GLENMARK PHARMS	<u>1%</u>	<u>A090219 001</u> Aug 03, 2010
<u>AB</u> ! TARO	<u>1%</u>	<u>A072640 001</u> Aug 31, 1993

SOLUTION; TOPICAL

CLOTRIMAZOLE

<u>AT</u> ! TARO	<u>1%</u>	<u>A074580 001</u> Jul 29, 1996
<u>AT</u> TEVA	<u>1%</u>	<u>A073306 001</u> Feb 28, 1995

TROCHE/LOZENGE; ORAL

CLOTRIMAZOLE

<u>AB</u> PADDOCK LLC	<u>10MG</u>	<u>A076763 001</u> Oct 28, 2005
<u>AB</u> ! WEST-WARD PHARMS INT	<u>10MG</u>	<u>A076387 001</u> Jul 29, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-104 (of 436)

CLOZAPINE

SUSPENSION;ORAL

VERSACLOZ

+! TASMAN PHARMA

50MG/ML

N203479 001 Feb 06, 2013

TABLET;ORAL

CLOZAPINE

<u>AB</u>	ACCORD HLTHCARE	<u>25MG</u>	<u>A202873 001</u>	Nov 25, 2015
<u>AB</u>		<u>100MG</u>	<u>A202873 002</u>	Nov 25, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>25MG</u>	<u>A206433 001</u>	Nov 29, 2016
<u>AB</u>		<u>50MG</u>	<u>A206433 002</u>	Nov 29, 2016
<u>AB</u>		<u>100MG</u>	<u>A206433 003</u>	Nov 29, 2016
<u>AB</u>		<u>200MG</u>	<u>A206433 004</u>	Nov 29, 2016
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>12.5MG</u>	<u>A074949 003</u>	Jul 31, 2003
<u>AB</u>		<u>25MG</u>	<u>A074949 001</u>	Nov 26, 1997
<u>AB</u>		<u>50MG</u>	<u>A074949 004</u>	Apr 25, 2005
<u>AB</u>		<u>50MG</u>	<u>A076809 003</u>	Dec 16, 2005
<u>AB</u>		<u>100MG</u>	<u>A074949 002</u>	Nov 26, 1997
<u>AB</u>		<u>100MG</u>	<u>A076809 002</u>	Dec 16, 2005
<u>AB</u>		<u>200MG</u>	<u>A076809 001</u>	Dec 16, 2005
<u>AB</u>	MAYNE PHARMA	<u>25MG</u>	<u>A203807 001</u>	Sep 17, 2015
<u>AB</u>		<u>50MG</u>	<u>A203807 003</u>	Aug 22, 2017
<u>AB</u>		<u>100MG</u>	<u>A203807 002</u>	Sep 17, 2015
<u>AB</u>		<u>200MG</u>	<u>A203807 004</u>	Aug 22, 2017
<u>AB</u>	MYLAN	<u>12.5MG</u>	<u>A075417 003</u>	Apr 15, 2010
<u>AB</u>		<u>25MG</u>	<u>A075417 001</u>	May 27, 1999
<u>AB</u>		<u>50MG</u>	<u>A075417 004</u>	Apr 15, 2010
<u>AB</u>		<u>100MG</u>	<u>A075417 002</u>	May 27, 1999
<u>AB</u>		<u>200MG</u>	<u>A075417 005</u>	Apr 15, 2010
<u>AB</u>	SUN PHARM INDs INC	<u>25MG</u>	<u>A075713 001</u>	Nov 15, 2002
<u>AB</u>		<u>50MG</u>	<u>A075713 003</u>	Aug 19, 2005
<u>AB</u>		<u>100MG</u>	<u>A075713 002</u>	Nov 15, 2002
<u>AB</u>		<u>200MG</u>	<u>A075713 004</u>	Nov 07, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A209480 001</u>	Dec 06, 2017
<u>AB</u>		<u>50MG</u>	<u>A209480 002</u>	Dec 06, 2017
<u>AB</u>		<u>100MG</u>	<u>A209480 003</u>	Dec 06, 2017
<u>AB</u>		<u>200MG</u>	<u>A209480 004</u>	Dec 06, 2017

CLOZARIL

<u>AB</u>	+ HERITAGE LIFE	<u>25MG</u>	<u>N019758 001</u>	Sep 26, 1989
<u>AB</u>	+!	<u>100MG</u>	<u>N019758 002</u>	Sep 26, 1989

TABLET, ORALLY DISINTEGRATING;ORAL

CLOZAPINE

<u>AB</u>	BARR LABS INC	<u>25MG</u>	<u>A090308 001</u>	Nov 25, 2015
<u>AB</u>		<u>100MG</u>	<u>A090308 002</u>	Nov 25, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>25MG</u>	<u>A201824 002</u>	Sep 15, 2015
<u>AB</u>		<u>100MG</u>	<u>A201824 003</u>	Sep 15, 2015
<u>AB</u>	TEVA PHARMS USA	<u>150MG</u>	<u>A203039 001</u>	Nov 25, 2015
<u>AB</u>		<u>200MG</u>	<u>A203039 002</u>	Nov 25, 2015

FAZACLO ODT

<u>AB</u>	+ JAZZ PHARMS III	<u>25MG</u>	<u>N021590 001</u>	Feb 10, 2004
<u>AB</u>	+!	<u>100MG</u>	<u>N021590 002</u>	Feb 10, 2004
<u>AB</u>	+	<u>150MG</u>	<u>N021590 005</u>	Jul 09, 2010
<u>AB</u>	+	<u>200MG</u>	<u>N021590 006</u>	Jul 09, 2010
	+	12.5MG	N021590 004	May 30, 2007

COBICISTAT

TABLET;ORAL

TYBOST

+! GILEAD SCIENCES INC 150MG

N203094 001 Sep 24, 2014

COBICISTAT; DARUNAVIR ETHANOLATE

TABLET;ORAL

PREZCOBIX

+! JANSSEN PRODS

150MG;EQ 800MG BASE

N205395 001 Jan 29, 2015

COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

GENVOYA

+! GILEAD SCIENCES INC

150MG;150MG;200MG;EQ 10MG BASE

N207561 001 Nov 05, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-105 (of 436)

COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

STRIBILD

+! GILEAD SCIENCES INC 150MG;150MG;200MG;300MG

N203100 001 Aug 27, 2012

COBIMETINIB FUMARATE

TABLET;ORAL

COTELLIC

+! GENENTECH INC EQ 20MG BASE

N206192 001 Nov 10, 2015

COCAINE HYDROCHLORIDE

SOLUTION;NASAL

GORELTO

+! GENUS LIFESCIENCES 4%

N209963 001 Dec 14, 2017

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

<u>AA</u>	VINTAGE	<u>10MG/5ML;5MG/5ML;6.25MG/5ML</u>	<u>A040660 001</u>	Dec 07, 2006
<u>AA</u>	<u>PROMETH VC W/ CODEINE</u>	<u>10MG/5ML;5MG/5ML;6.25MG/5ML</u>	<u>A088764 001</u>	Oct 31, 1984
<u>AA</u>	! ACTAVIS MID ATLANTIC	<u>10MG/5ML;5MG/5ML;6.25MG/5ML</u>	<u>A040674 001</u>	Dec 23, 2014
<u>AA</u>	HI-TECH PHARMA CO	<u>10MG/5ML;5MG/5ML;6.25MG/5ML</u>	<u>A040674 001</u>	Dec 23, 2014
<u>AA</u>	<u>PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE</u>	<u>10MG/5ML;5MG/5ML;6.25MG/5ML</u>	<u>A200963 001</u>	Aug 26, 2015

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

<u>AA</u>	! ACTAVIS MID ATLANTIC	<u>10MG/5ML;6.25MG/5ML</u>	<u>A088763 001</u>	Oct 31, 1984
<u>AA</u>	AMNEAL PHARMS	<u>10MG/5ML;6.25MG/5ML</u>	<u>A200894 001</u>	Apr 24, 2013
<u>AA</u>	HI TECH PHARMA	<u>10MG/5ML;6.25MG/5ML</u>	<u>A040151 001</u>	Aug 26, 1997
<u>AA</u>	NOSTRUM LABS INC	<u>10MG/5ML;6.25MG/5ML</u>	<u>A090180 001</u>	Mar 17, 2010
<u>AA</u>	TRIS PHARMA INC	<u>10MG/5ML;6.25MG/5ML</u>	<u>A200386 001</u>	Jun 29, 2012
<u>AA</u>	WOCKHARDT BIO AG	<u>10MG/5ML;6.25MG/5ML</u>	<u>A088875 001</u>	Dec 17, 1984
	<u>PROMETHAZINE WITH CODEINE</u>			
<u>AA</u>	VINTAGE	<u>10MG/5ML;6.25MG/5ML</u>	<u>A040650 001</u>	Jan 31, 2006

CODEINE PHOSPHATE; PSEUDOEPHENDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

TRIACIN-C

+! STI PHARMA LLC 10MG/5ML;30MG/5ML;1.25MG/5ML

A088704 001 Mar 22, 1985

CODEINE SULFATE

TABLET;ORAL

CODEINE SULFATE

<u>AB</u>	LANNETT HOLDINGS INC	<u>15MG</u>	<u>A203046 001</u>	Jun 13, 2014
<u>AB</u>		<u>30MG</u>	<u>A203046 002</u>	Jun 13, 2014
<u>AB</u>		<u>60MG</u>	<u>A203046 003</u>	Jun 13, 2014
<u>AB</u>	! WEST-WARD PHARMS INT	<u>15MG</u>	<u>N022402 001</u>	Jul 16, 2009
<u>AB</u>	+	<u>30MG</u>	<u>N022402 002</u>	Jul 16, 2009
<u>AB</u>	++!	<u>60MG</u>	<u>N022402 003</u>	Jul 16, 2009

COLCHICINE

CAPSULE;ORAL

MITIGARE

+! HIKMA INTL PHARMS 0.6MG

N204820 001 Sep 26, 2014

TABLET;ORAL

COLCRYS

+! TAKEDA PHARMS USA 0.6MG

N022352 001 Jul 29, 2009

COLCHICINE; PROBENECID

TABLET;ORAL

COL-PROBENECID

<u>AB</u>	! WATSON LABS	<u>0.5MG;500MG</u>	<u>A084279 001</u>	
<u>AB</u>	<u>PROBENECID AND COLCHICINE</u>	<u>0.5MG;500MG</u>	<u>A040618 001</u>	May 13, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-106 (of 436)

COLESEVELAM HYDROCHLORIDE

FOR SUSPENSION;ORAL

WELCHOL

+ DAIICHI SANKYO 1.875GM/PACKET  
+! 3.75GM/PACKET

N022362 001 Oct 02, 2009  
N022362 002 Oct 02, 2009

TABLET;ORAL

WELCHOL

+! DAIICHI SANKYO 625MG

N021176 001 May 26, 2000

COlestipol Hydrochloride

GRANULE;ORAL

COlestid

AB + PHARMACIA UPJOHN 5GM/SCOOPFUL  
AB +! 5GM/PACKET

N017563 003 Sep 22, 1995  
N017563 004 Sep 22, 1995

COlestipol Hydrochloride

AB IMPAX LABS 5GM/SCOOPFUL  
AB 5GM/PACKET

A077277 001 May 02, 2006  
A077277 002 May 02, 2006

FLAVORED COlestid

+ PHARMACIA UPJOHN 5GM/PACKET  
+ 5GM/SCOOPFUL

N017563 001  
N017563 002

TABLET;ORAL

COlestid

AB +! PHARMACIA UPJOHN 1GM

N020222 001 Jul 19, 1994

COlestipol Hydrochloride

AB IMPAX LABS 1GM

A077510 001 Oct 24, 2006

COLISTIMETHATE SODIUM

INJECTABLE;INJECTION

COLISTIMETHATE SODIUM

AP EMCURE PHARMS LTD EQ 150MG BASE/VIAL  
AP FRESENIUS KABI USA EQ 150MG BASE/VIAL  
AP PADDOCK LLC EQ 150MG BASE/VIAL  
AP SAGENT PHARMS EQ 150MG BASE/VIAL  
AP X GEN PHARMS EQ 150MG BASE/VIAL  
AP XELLIA PHARMS APS EQ 150MG BASE/VIAL

A202359 001 Sep 28, 2012  
A065364 001 Apr 17, 2008  
A065177 001 Mar 19, 2004  
A201365 001 Feb 19, 2014  
A064216 001 Feb 26, 1999  
A205356 001 May 29, 2015

COLY-MYCIN M

AP +! PAR STERILE PRODUCTS EQ 150MG BASE/VIAL

N050108 002

COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS;OTIC

COLY-MYCIN S

+! ENDO PHARMS INC EQ 3MG BASE/ML;10MG/ML;EQ 3.3MG  
BASE/ML;0.5MG/ML

N050356 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE;IV (INFUSION)

VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER

+! CUMBERLAND PHARMS 20MG/100ML (0.2MG/ML)

N021697 002 Oct 08, 2008

COPANLISIB DIHYDROCHLORIDE

POWDER;IV (INFUSION)

ALIQOPA

+! BAYER HEALTHCARE 60MG/VIAL

N209936 001 Sep 14, 2017

COPPER

INTRAUTERINE DEVICE;INTRAUTERINE

PARAGARD T 380A

+! COOPERSURGICAL 309MG/COPPER

N018680 001 Nov 15, 1984

CORTICORELIN OVINE TRIFLUtATE

INJECTABLE;INJECTION

ACTHREL

+! FERRING EQ 0.1MG BASE/VIAL

N020162 001 May 23, 1996

CORTicotropin

INJECTABLE;INJECTION

H.P. ACTHAR GEL

+! MALLINCKRODT ARD 80 UNITS/ML

N008372 008

CORTISONE ACETATE

TABLET;ORAL

CORTISONE ACETATE

! HIKMA INTL PHARMS 25MG

A080776 002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-107 (of 436)

COSYNTROPIN

INJECTABLE; INJECTION

CORTROSYN

AP +! AMPHASTAR PHARMS 0.25MG/VIAL

N016750 001

INC

COSYNTROPIN

AP MYLAN INSTITUTIONAL 0.25MG/VIAL

A090574 001 Dec 17, 2009

AP SANDOZ 0.25MG/VIAL

A202147 001 Jun 29, 2012

CRISABOROLE

OINTMENT; TOPICAL

EUCRISA

+! ANACOR PHARMS INC 2%

N207695 001 Dec 14, 2016

CRIZOTINIB

CAPSULE; ORAL

XALKORI

+! PF PRISM CV 200MG

N202570 001 Aug 26, 2011

+! 250MG

N202570 002 Aug 26, 2011

CROFTELEMER

TABLET, DELAYED RELEASE; ORAL

FULYZAQ

+! NAPO PHARMS INC 125MG

N202292 001 Dec 31, 2012

CROMOLYN SODIUM

CONCENTRATE; ORAL

CROMOLYN SODIUM

AA AILEX PHARMS LLC 100MG/5ML

A209264 001 Oct 16, 2017

AA MICRO LABS LTD 100MG/5ML

A202745 001 Apr 04, 2013

AA RISING PHARMS INC 100MG/5ML

A202583 001 Oct 27, 2011

GASTROCROM

AA +! MYLAN SPECIALITY LP 100MG/5ML

N020479 001 Feb 29, 1996

SOLUTION; INHALATION

CROMOLYN SODIUM

AN AILEX PHARMS LLC 10MG/ML

A209453 001 Oct 16, 2017

AN MYLAN SPECIALITY LP 10MG/ML

A074209 001 Apr 26, 1994

AN ! TEVA PHARMS 10MG/ML

A075271 001 Jan 18, 2000

AN WOCKHARDT BIO AG 10MG/ML

A075346 001 Oct 25, 1999

SOLUTION/DROPS; OPHTHALMIC

CROMOLYN SODIUM

AT ! AKORN 4%

A074706 001 Apr 29, 1998

AT ALCON 4%

A075282 001 Jun 16, 1999

CROTAMITON

CREAM; TOPICAL

EURAX

+! RANBAXY 10%

N006927 001

LOTION; TOPICAL

CROTAN

AT MARNEL PHARMS 10%

A087204 001

EURAX

AT +! RANBAXY 10%

N009112 003

CUPRIC CHLORIDE

INJECTABLE; INJECTION

CUPRIC CHLORIDE IN PLASTIC CONTAINER

+! HOSPIRA EQ 0.4MG COPPER/ML

N018960 001 Jun 26, 1986

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN

AP ! LUITPOLD 1MG/ML

A080737 001

AP MYLAN LABS LTD 1MG/ML

A204829 001 Jun 05, 2017

AP SOMERSET THERAPS 1MG/ML

A206503 001 Dec 11, 2015

LLC

AP WEST-WARD PHARMS 1MG/ML

A080515 002

INT

VIBISONE

AP ! FRESENIUS KABI USA 1MG/ML

A080557 003

SPRAY, METERED; NASAL

NASCOBAL

+! ENDO PHARMS INC 0.5MG/SPRAY

N021642 001 Jan 31, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-108 (of 436)

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

AMRIX

+ TEVA PHARMS INTL	15MG	N021777 001 Feb 01, 2007
+!	30MG	N021777 002 Feb 01, 2007

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>5MG</u>	<u>A071611 002</u>	Feb 03, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A071611 003</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A071611 001</u>	May 03, 1989
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078643 001</u>	Sep 26, 2008
<u>AB</u>		<u>10MG</u>	<u>A078643 002</u>	Sep 26, 2008
<u>AB</u>	FRONTIDA BIOPHARM	<u>5MG</u>	<u>A073541 002</u>	Apr 06, 2006
<u>AB</u>		<u>10MG</u>	<u>A073541 001</u>	May 23, 1995
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A090478 001</u>	Jul 23, 2010
<u>AB</u>		<u>10MG</u>	<u>A090478 002</u>	Jul 23, 2010
<u>AB</u>	JUBILANT CADISTA	<u>5MG</u>	<u>A077563 001</u>	Apr 19, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A077563 003</u>	Aug 25, 2017
<u>AB</u>		<u>10MG</u>	<u>A077563 002</u>	Apr 19, 2006
<u>AB</u>	KVK TECH	<u>5MG</u>	<u>A078048 001</u>	Feb 28, 2011
<u>AB</u>		<u>10MG</u>	<u>A078048 002</u>	Feb 28, 2011
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A073144 002</u>	Feb 03, 2006
<u>AB</u>		<u>7.5MG</u>	<u>A073144 003</u>	Mar 25, 2013
<u>AB</u>	!	<u>10MG</u>	<u>A073144 001</u>	May 30, 1991
<u>AB</u>	ORIT LABS LLC	<u>5MG</u>	<u>A078218 002</u>	Jun 19, 2015
<u>AB</u>		<u>10MG</u>	<u>A078218 001</u>	Apr 18, 2008
<u>AB</u>	OXFORD PHARMS	<u>5MG</u>	<u>A077209 002</u>	Feb 03, 2006
<u>AB</u>		<u>10MG</u>	<u>A077209 001</u>	Oct 04, 2005
<u>AB</u>	PLIVA	<u>10MG</u>	<u>A074421 001</u>	Sep 29, 1995
<u>AB</u>	PRINSTON INC	<u>5MG</u>	<u>A077797 001</u>	Feb 28, 2007
<u>AB</u>		<u>10MG</u>	<u>A077797 002</u>	Feb 28, 2007
<u>AB</u>	RUBICON RES PVT LTD	<u>5MG</u>	<u>A208170 001</u>	May 31, 2017
<u>AB</u>		<u>7.5MG</u>	<u>A208170 002</u>	May 31, 2017
<u>AB</u>		<u>10MG</u>	<u>A208170 003</u>	May 31, 2017
<u>AB</u>	SUN PHARM INDs LTD	<u>5MG</u>	<u>A078722 001</u>	May 12, 2008
<u>AB</u>		<u>7.5MG</u>	<u>A078722 002</u>	May 12, 2008
<u>AB</u>		<u>10MG</u>	<u>A078722 003</u>	May 12, 2008

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPENTOLATE

<u>AT</u>	AKORN	<u>1%</u>	<u>A040164 001</u>	Jan 13, 1997
-----------	-------	-----------	--------------------	--------------

CYCLOGYL

<u>AT</u>	! NOVARTIS PHARMS CORP	<u>0.5%</u>	<u>A084109 001</u>	
-----------	------------------------	-------------	--------------------	--

<u>AT</u>	!	<u>1%</u>	<u>A084110 001</u>	
-----------	---	-----------	--------------------	--

CYCLOPENTOLATE HYDROCHLORIDE

<u>AT</u>	AKORN INC	<u>0.5%</u>	<u>A205937 001</u>	Dec 09, 2015
-----------	-----------	-------------	--------------------	--------------

PENTOLAIR

<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>A040075 001</u>	Apr 29, 1994
-----------	-----------------	-----------	--------------------	--------------

CYCLOGYL

!	NOVARTIS PHARMS CORP	2%	A084108 001	
---	----------------------	----	-------------	--

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOMYDRIL

!	NOVARTIS PHARMS CORP	0.2%;1%	A084300 001	
---	----------------------	---------	-------------	--

CYCLOPHOSPHAMIDE

CAPSULE; ORAL

CYCLOPHOSPHAMIDE

+	WEST-WARD PHARMS INT	25MG	N203856 001	Sep 16, 2013
+!		50MG	N203856 002	Sep 16, 2013

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

<u>AP</u>	! BAXTER HLTHCARE	<u>500MG/VIAL</u>	<u>A040745 001</u>	May 21, 2008
<u>AP</u>	!	<u>1GM/VIAL</u>	<u>A040745 002</u>	May 21, 2008
<u>AP</u>	!	<u>2GM/VIAL</u>	<u>A040745 003</u>	May 21, 2008
<u>AP</u>	JIANGSU HENGRUI MED	<u>500MG/VIAL</u>	<u>A204555 001</u>	Oct 31, 2014
<u>AP</u>		<u>1GM/VIAL</u>	<u>A204555 002</u>	Oct 31, 2014
<u>AP</u>		<u>2GM/VIAL</u>	<u>A204555 003</u>	Oct 31, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-109 (of 436)

CYCLOSERINE

CAPSULE;ORAL  
 SEROMYCIN  
 ! PURDUE GMP 250MG A060593 001

CYCLOSPORINE

CAPSULE;ORAL

CYCLOSPORINE

<u>AB1</u>	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A065110 003</u>	Mar 29, 2005
<u>AB1</u>		<u>50MG</u>	<u>A065110 001</u>	Mar 29, 2005
<u>AB1</u>		<u>100MG</u>	<u>A065110 002</u>	Mar 29, 2005
<u>AB1</u>	MAYNE PHARMA	<u>25MG</u>	<u>A065044 002</u>	Dec 20, 2000
<u>AB1</u>		<u>100MG</u>	<u>A065044 001</u>	Dec 20, 2000
<u>AB1</u>	SANDOZ	<u>25MG</u>	<u>A065017 002</u>	Jan 13, 2000
<u>AB1</u>		<u>100MG</u>	<u>A065017 001</u>	Jan 13, 2000

GENGRAF

<u>AB1</u>	ABBVIE	<u>25MG</u>	<u>A065003 001</u>	May 12, 2000
<u>AB1</u>		<u>50MG</u>	<u>A065003 002</u>	May 12, 2000
<u>AB1</u>		<u>100MG</u>	<u>A065003 003</u>	May 12, 2000

NEORAL

<u>AB1</u> +	NOVARTIS	<u>25MG</u>	<u>N050715 001</u>	Jul 14, 1995
<u>AB1</u> +!		<u>100MG</u>	<u>N050715 002</u>	Jul 14, 1995

CYCLOSPORINE

<u>AB2</u>	APOTEX	<u>25MG</u>	<u>A065040 001</u>	May 09, 2002
<u>AB2</u>		<u>100MG</u>	<u>A065040 002</u>	May 09, 2002

SANDIMMUNE

<u>AB2</u> +	NOVARTIS	<u>25MG</u>	<u>N050625 001</u>	Mar 02, 1990
<u>AB2</u> +!		<u>100MG</u>	<u>N050625 002</u>	Mar 02, 1990
BX +		50MG	N050625 003	Nov 23, 1992

EMULSION;OPHTHALMIC

RESTASIS

+! ALLERGAN 0.05%

N050790 001 Dec 23, 2002

RESTASIS MULTIDOSE

+! ALLERGAN 0.05%

N050790 002 Oct 27, 2016

INJECTABLE;INJECTION

CYCLOSPORINE

<u>AP</u>	LUITPOLD	<u>50MG/ML</u>	<u>A065151 001</u>	Oct 07, 2003
<u>AP</u>	WEST-WARD PHARMS INT	<u>50MG/ML</u>	<u>A065004 001</u>	Oct 29, 1999

SANDIMMUNE

<u>AP</u> +!	NOVARTIS	<u>50MG/ML</u>	<u>N050573 001</u>	Nov 14, 1983
SOLUTION;ORAL				

CYCLOSPORINE

<u>AB1</u>	ABBVIE	<u>100MG/ML</u>	<u>A065025 001</u>	Mar 03, 2000
<u>AB1</u>	IVAX SUB TEVA PHARMS	<u>100MG/ML</u>	<u>A065078 001</u>	Mar 25, 2005
<u>AB1</u>	MAYNE PHARMA	<u>100MG/ML</u>	<u>A065054 001</u>	Dec 18, 2001

NEORAL

<u>AB1</u> +!	NOVARTIS	<u>100MG/ML</u>	<u>N050716 001</u>	Jul 14, 1995
---------------	----------	-----------------	--------------------	--------------

CYCLOSPORINE

<u>AB2</u>	WOCKHARDT BIO AG	<u>100MG/ML</u>	<u>A065133 001</u>	Sep 17, 2004
------------	------------------	-----------------	--------------------	--------------

SANDIMMUNE

<u>AB2</u> +!	NOVARTIS	<u>100MG/ML</u>	<u>N050574 001</u>	Nov 14, 1983
---------------	----------	-----------------	--------------------	--------------

CYPROHEPTADINE HYDROCHLORIDE

SYRUP;ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	BIO-PHARM INC	<u>2MG/5ML</u>	<u>A204823 001</u>	Dec 27, 2016
<u>AA</u> !	LYNE	<u>2MG/5ML</u>	<u>A040668 001</u>	Jun 28, 2006
<u>AA</u>	PHARM ASSOC	<u>2MG/5ML</u>	<u>A091295 001</u>	Mar 28, 2013
<u>AA</u>	SILARK PHARMS INC	<u>2MG/5ML</u>	<u>A203191 001</u>	Jul 13, 2017

TABLET;ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>	AILEX PHARMS LLC	<u>4MG</u>	<u>A207555 001</u>	Jan 31, 2017
<u>AA</u>	APEX PHARMS INC	<u>4MG</u>	<u>A207783 001</u>	Dec 29, 2016
<u>AA</u>	APPCO PHARMA LLC	<u>4MG</u>	<u>A206553 001</u>	Nov 29, 2016
<u>AA</u>	COREPHARMA	<u>4MG</u>	<u>A040537 001</u>	Sep 30, 2003
<u>AA</u>	INGENUS PHARMS NJ	<u>4MG</u>	<u>A205087 001</u>	Sep 23, 2015
<u>AA</u> !	IVAX SUB TEVA PHARMS	<u>4MG</u>	<u>A087056 001</u>	
<u>AA</u>	PAR PHARM	<u>4MG</u>	<u>A087129 001</u>	
<u>AA</u>	SANTOS BIOTECH	<u>4MG</u>	<u>A040644 001</u>	May 30, 2006
<u>AA</u>	ZYDUS PHARMS USA	<u>4MG</u>	<u>A208938 001</u>	May 19, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-110 (of 436)

CYPROHEPTADINE HYDROCHLORIDE

TABLET;ORAL

**CYPROHEPTADINE HYDROCHLORIDE**

INC

CYSTEAMINE BITARTRATE

CAPSULE;ORAL

CYSTAGON

+ MYLAN	EQ 50MG BASE
+!	EQ 150MG BASE

N020392 001	Aug 15, 1994
N020392 002	Aug 15, 1994

CAPSULE, DELAYED RELEASE;ORAL

PROCYSBI

+ HORIZON PHARMA USA	EQ 25MG BASE
+!	EQ 75MG BASE

N203389 001	Apr 30, 2013
N203389 002	Apr 30, 2013

CYSTEAMINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CYSTARAN

+! LEADIANT BIOSCI INC	EQ 0.44% BASE
------------------------	---------------

N200740 001	Oct 02, 2012
-------------	--------------

CYTARABINE

INJECTABLE;INJECTION

**CYTARABINE**

<b>AP</b> ! FRESENIUS KABI USA	<b><u>100MG/ML</u></b>
<b>AP</b>	<b><u>20MG/ML</u></b>
<b>AP</b> ! HOSPIRA	<b><u>20MG/ML</u></b>
<b>AP</b> !	<b><u>20MG/ML</u></b>
<b>AP</b> !	<b><u>20MG/ML</u></b>
<b>AP</b>	<b><u>100MG/ML</u></b>
<b>AP</b>	<b><u>20MG/ML</u></b>
<b>AP</b>	<b><u>20MG/ML</u></b>
<b>AP</b>	<b><u>100MG/ML</u></b>
<b>AP</b>	<b><u>20MG/ML</u></b>
<b>AP</b>	<b><u>20MG/ML</u></b>
<b>AP</b>	<b><u>100MG/ML</u></b>
<b>AP</b>	<b><u>20MG/ML</u></b>
<b>AP</b>	<b><u>100MG/VIAL</u></b>
WEST-WARD PHARMS INT	
!	500MG/VIAL
!	1GM/VIAL
!	2GM/VIAL

<b>A076512 001</b>	Jan 15, 2004
<b>A206190 001</b>	Nov 09, 2017
<b>A071868 001</b>	Jun 04, 1990
<b>A072168 001</b>	Aug 31, 1990
<b>A072945 001</b>	Feb 28, 1994
<b>A075383 001</b>	Nov 22, 1999
<b>A200914 001</b>	Dec 13, 2011
<b>A200915 001</b>	Dec 13, 2011
<b>A201784 001</b>	Jan 30, 2012
<b>A200916 001</b>	Dec 13, 2011
A071471 001	Aug 02, 1989
A071472 001	Aug 02, 1989
A074245 001	Aug 31, 1994
A074245 002	Aug 31, 1994

INJECTABLE, LIPOSOMAL;INJECTION

DEPOCYT

+! PACIRA PHARMS INC	10MG/ML
----------------------	---------

N021041 001	Apr 01, 1999
-------------	--------------

CYTARABINE; DAUNORUBICIN

POWDER;IV (INFUSION)

VYXEOS

+! CELATOR PHARMS	100MG;44MG
-------------------	------------

N209401 001	Aug 03, 2017
-------------	--------------

DABIGATRAN ETEXILATE MESYLATE

CAPSULE;ORAL

PRADAXA

+ BOEHRINGER INGELHEIM	EQ 75MG BASE
+	EQ 110MG BASE
+!	EQ 150MG BASE

N022512 001	Oct 19, 2010
N022512 003	Nov 20, 2015
N022512 002	Oct 19, 2010

DABRAFENIB MESYLATE

CAPSULE;ORAL

TAFINLAR

+ NOVARTIS PHARMS CORP	EQ 50MG BASE
+!	EQ 75MG BASE

N202806 001	May 29, 2013
N202806 002	May 29, 2013

DACARBAZINE

INJECTABLE;INJECTION

**DACARBAZINE**

<b>AP</b> ! FRESENIUS KABI USA	<b><u>200MG/VIAL</u></b>
<b>AP</b>	<b><u>200MG/VIAL</u></b>
<b>AP</b>	<b><u>200MG/VIAL</u></b>
<b>AP</b> !	<b><u>500MG/VIAL</u></b>
<b>AP</b>	<b><u>200MG/VIAL</u></b>
<b>AP</b> !	<b><u>500MG/VIAL</u></b>
FRESENIUS KABI USA	100MG/VIAL

<b>A075371 002</b>	Aug 27, 1999
<b>A075940 001</b>	Oct 18, 2001
<b>A075259 002</b>	Aug 27, 1998
<b>A075259 001</b>	Sep 22, 2000
<b>A075812 001</b>	Jun 15, 2001
<b>A075812 002</b>	Oct 31, 2002
A075371 001	Aug 27, 1999

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-111 (of 436)

DACLATASVIR DIHYDROCHLORIDE

TABLET; ORAL

DAKLINZA

+ BRISTOL-MYERS SQUIBB	EQ 30MG BASE	N206843 001 Jul 24, 2015
+!	EQ 60MG BASE	N206843 002 Jul 24, 2015
+	EQ 90MG BASE	N206843 003 Apr 13, 2016

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

<u>AP</u> +! RECORDATI RARE	<u>0.5MG/VIAL</u>	<u>N050682 001</u>
<u>AP</u> LUITPOLD PHARMS INC	<u>0.5MG/VIAL</u>	<u>A202562 001</u> Aug 23, 2013
<u>AP</u> MYLAN LABS LTD	<u>0.5MG/VIAL</u>	<u>A203385 001</u> Nov 09, 2017

DALBAVANCIN HYDROCHLORIDE

POWDER; IV (INFUSION)

DALVANCE

+! ALLERGAN SALES LLC	EQ 500MG BASE/VIAL	N021883 001 May 23, 2014
-----------------------	--------------------	--------------------------

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL

AMPYRA

<u>AB</u> +! ACORDA	<u>10MG</u>	<u>N022250 001</u> Jan 22, 2010
<u>AB</u> ACTAVIS LABS FL INC	<u>10MG</u>	<u>A206836 001</u> Jan 23, 2017
<u>AB</u> AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A206811 001</u> Jan 23, 2017

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; IV (INFUSION)

SYNERCID

+! KING PHARMS	350MG/VIAL;150MG/VIAL	N050748 001 Sep 21, 1999
----------------	-----------------------	--------------------------

DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

+ PFIZER INC	2,500IU/0.2ML (12,500IU/ML)	N020287 001 Dec 22, 1994
+	5,000IU/0.2ML (25,000IU/ML)	N020287 003 Mar 18, 1996
+	7,500IU/0.3ML (25,000IU/ML)	N020287 005 Apr 04, 2002
+	10,000IU/ML (10,000IU/ML)	N020287 004 Jan 30, 1998
+	12,500IU/0.5ML (25,000IU/ML)	N020287 009 May 01, 2007
+	15,000IU/0.6ML (25,000IU/ML)	N020287 010 May 01, 2007
+	18,000IU/0.72ML (25,000IU/ML)	N020287 011 May 01, 2007
+!	95,000IU/3.8ML (25,000IU/ML)	N020287 006 Apr 04, 2002

DANAZOL

CAPSULE; ORAL

DANAZOL

<u>AB</u> BARR	<u>50MG</u>	<u>A074582 003</u> May 29, 1998
<u>AB</u>	<u>100MG</u>	<u>A074582 002</u> May 29, 1998
<u>AB</u> !	<u>200MG</u>	<u>A074582 001</u> Aug 09, 1996
<u>AB</u> LANNETT	<u>50MG</u>	<u>A077246 002</u> Apr 19, 2007
<u>AB</u>	<u>100MG</u>	<u>A077246 003</u> Apr 19, 2007
<u>AB</u>	<u>200MG</u>	<u>A077246 001</u> Sep 28, 2005

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUM

<u>AB</u> + PAR STERILE PRODUCTS	<u>25MG</u>	<u>N017443 001</u>
<u>AB</u> +	<u>50MG</u>	<u>N017443 003</u>
<u>AB</u> +!	<u>100MG</u>	<u>N017443 002</u>

DANTROLENE SODIUM

<u>AB</u> ELITE LABS INC	<u>25MG</u>	<u>A076686 001</u> Oct 24, 2005
<u>AB</u>	<u>50MG</u>	<u>A076686 002</u> Oct 24, 2005
<u>AB</u>	<u>100MG</u>	<u>A076686 003</u> Oct 24, 2005
<u>AB</u> IMPAX LABS	<u>25MG</u>	<u>A076856 001</u> Mar 01, 2005
<u>AB</u>	<u>50MG</u>	<u>A076856 002</u> Mar 01, 2005
<u>AB</u>	<u>100MG</u>	<u>A076856 003</u> Mar 01, 2005

FOR SUSPENSION; INTRAVENOUS

RYANODEX

+! EAGLE PHARMS	250MG/VIAL	N205579 001 Jul 22, 2014
-----------------	------------	--------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-112 (of 436)

DANTROLENE SODIUM

INJECTABLE; INJECTION

DANTRIUM

<u>AP</u>	+!	PAR STERILE PRODUCTS	<u>20MG/VIAL</u>	<u>N018264 001</u>
<u>AP</u>		HIKMA PHARMS	<u>20MG/VIAL</u>	<u>A204762 001</u> Jun 19, 2017
<u>AP</u>		MYLAN INSTITUTIONAL	<u>20MG/VIAL</u>	<u>A205239 001</u> Feb 18, 2016
<u>AP</u>		<u>REVONTO</u>	<u>20MG/VIAL</u>	<u>A078378 001</u> Jul 24, 2007

DAPAGLIFLOZIN PROPANEDIOL

TABLET; ORAL

FARXIGA

+ ASTRAZENECA AB	EQ 5MG BASE	N202293 001 Jan 08, 2014
+!	EQ 10MG BASE	N202293 002 Jan 08, 2014

DAPAGLIFLOZIN PROPYANEDIOL; METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

XIGDUO XR

+ ASTRAZENECA AB	EQ 2.5MG BASE;1GM	N205649 005 Jul 28, 2017
+	EQ 5MG BASE;500MG	N205649 001 Oct 29, 2014
+	EQ 5MG BASE;1GM	N205649 002 Oct 29, 2014
+	EQ 10MG BASE;500MG	N205649 003 Oct 29, 2014
+!	EQ 10MG BASE;1GM	N205649 004 Oct 29, 2014

DAPAGLIFLOZIN PROPYANEDIOL; SAXagliptin HYDROCHLORIDE

TABLET; ORAL

QTERN

+! ASTRAZENECA AB	EQ 10MG BASE;EQ 5MG BASE	N209091 001 Feb 27, 2017
-------------------	--------------------------	--------------------------

DAPSONE

GEL; TOPICAL

ACZONE

<u>AB</u>	+! ALLERGAN	<u>5%</u>	<u>N021794 001</u> Jul 07, 2005
<u>AB</u>	TARO	<u>5%</u>	<u>A209506 001</u> Oct 16, 2017
	ACZONE		
+!	ALLERGAN INC	7.5%	N207154 001 Feb 24, 2016

TABLET; ORAL

DAPSONE

<u>AB</u>	ACTAVIS LLC	<u>25MG</u>	<u>A204380 001</u> Mar 23, 2017
<u>AB</u>		<u>100MG</u>	<u>A204380 002</u> Mar 23, 2017
<u>AB</u>	ALVOGEN	<u>25MG</u>	<u>A205429 001</u> Jan 07, 2016
<u>AB</u>		<u>100MG</u>	<u>A205429 002</u> Jan 07, 2016
<u>AB</u>	JACOBUS	<u>25MG</u>	<u>A086841 001</u>
<u>AB</u>	!	<u>100MG</u>	<u>A086842 001</u>
<u>AB</u>	NOSTRUM LABS INC	<u>25MG</u>	<u>A203887 001</u> May 06, 2016
<u>AB</u>		<u>100MG</u>	<u>A203887 002</u> May 06, 2016
<u>AB</u>	NOVITIUM PHARMA	<u>25MG</u>	<u>A206505 001</u> Dec 01, 2016
<u>AB</u>		<u>100MG</u>	<u>A206505 002</u> Dec 01, 2016
<u>AB</u>	VIRTUS PHARMS	<u>25MG</u>	<u>A204074 001</u> May 10, 2016
<u>AB</u>		<u>100MG</u>	<u>A204074 002</u> May 10, 2016

DAPTOMYCIN

POWDER; INTRAVENOUS

CUBICIN

<u>AP</u>	+! CUBIST PHARMS LLC	<u>500MG/VIAL</u>	<u>N021572 002</u> Sep 12, 2003
<u>AP</u>	<u>DAPTOMYCIN</u>		
<u>AP</u>	CRANE PHARMS LLC	<u>500MG/VIAL</u>	<u>A206005 001</u> Jun 15, 2016
<u>AP</u>	HOSPIRA INC	<u>500MG/VIAL</u>	<u>A202857 001</u> Sep 12, 2014
<u>AP</u>	TEVA PARENTERAL	<u>500MG/VIAL</u>	<u>A091039 001</u> Mar 25, 2016
	CUBICIN RF		
+!	CUBIST PHARMS LLC	500MG/VIAL	N021572 003 Jul 06, 2016
	POWDER; IV (INFUSION)		
	DAPTOMYCIN		
+!	SAGENT PHARMS	350MG/VIAL	N208385 001 Sep 12, 2017
+!	XELLIA PHARMS APS	350MG/VIAL	N209949 001 Oct 20, 2017

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

DARIFENACIN

<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 7.5MG BASE</u>	<u>A207302 001</u> Jul 28, 2017
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>A207302 002</u> Jul 28, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-113 (of 436)

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

**DARIFENACIN HYDROBROMIDE**

<b>AB</b>	ALEMBIC PHARMS LTD	<u>EQ 7.5MG BASE</u>	<b>A207681 001</b>	Dec 08, 2017
<b>AB</b>		<u>EQ 15MG BASE</u>	<b>A207681 002</b>	Dec 08, 2017
<b>AB</b>	ANCHEN PHARMS	<u>EQ 7.5MG BASE</u>	<b>A091190 001</b>	Mar 13, 2015
<b>AB</b>		<u>EQ 15MG BASE</u>	<b>A091190 002</b>	Mar 13, 2015
<b>AB</b>	AUROBINDO PHARMA LTD	<u>EQ 7.5MG BASE</u>	<b>A206743 001</b>	Sep 19, 2016
<b>AB</b>		<u>EQ 15MG BASE</u>	<b>A206743 002</b>	Sep 19, 2016
<b>AB</b>	CIPPLA LTD	<u>EQ 7.5MG BASE</u>	<b>A207664 001</b>	Sep 01, 2016
<b>AB</b>		<u>EQ 15MG BASE</u>	<b>A207664 002</b>	Sep 01, 2016
<b>AB</b>	JUBILANT GENERICS	<u>EQ 7.5MG BASE</u>	<b>A205550 001</b>	Oct 12, 2016
<b>AB</b>		<u>EQ 15MG BASE</u>	<b>A205550 002</b>	Oct 12, 2016
<b>AB</b>	TORRENT PHARMS LTD	<u>EQ 7.5MG BASE</u>	<b>A205209 001</b>	Nov 17, 2016
<b>AB</b>		<u>EQ 15MG BASE</u>	<b>A205209 002</b>	Nov 17, 2016
<b><u>ENABLEX</u></b>				
<b>AB</b>	+ APIL	<u>EQ 7.5MG BASE</u>	<b>N021513 001</b>	Dec 22, 2004
<b>AB</b>	+!	<u>EQ 15MG BASE</u>	<b>N021513 002</b>	Dec 22, 2004

DARUNAVIR ETHANOLATE

SUSPENSION; ORAL

PREZISTA

+! JANSSEN PRODS

TABLET; ORAL

EQ 100MG BASE/ML

N202895 001 Dec 16, 2011

**DARUNAVIR ETHANOLATE**

<b>AB</b>	TEVA PHARMS USA	<u>EQ 600MG BASE</u>	<b>A202118 001</b>	Nov 21, 2017
<b><u>PREZISTA</u></b>				
<b>AB</b>	+ JANSSEN PRODS	<u>EQ 600MG BASE</u>	<b>N021976 002</b>	Feb 25, 2008
	+	EQ 75MG BASE	N021976 004	Dec 18, 2008
	+	EQ 150MG BASE	N021976 005	Dec 18, 2008
	+!	EQ 800MG BASE	N021976 006	Nov 09, 2012

DASABUVIR SODIUM ; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, TABLET; ORAL

VIEKIRA PAK (COPACKAGED)

+! ABBVIE INC

EQ 250MG BASE,N/A,N/A,N/A;  
N/A,12.5MG,75MG,50MG

N206619 001 Dec 19, 2014

DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET, EXTENDED RELEASE; ORAL

VIEKIRA XR

+! ABBVIE INC

EQ 200MG BASE;8.33MG;50MG;33.33MG

N208624 001 Jul 22, 2016

DASATINIB

TABLET; ORAL

**DASATINIB**

<b>AB</b>	APOTEX INC	<u>20MG</u>	<b>A202103 001</b>	Jun 10, 2016
<b>AB</b>		<u>50MG</u>	<b>A202103 002</b>	Jun 10, 2016
<b>AB</b>		<u>70MG</u>	<b>A202103 003</b>	Jun 10, 2016
<b>AB</b>		<u>100MG</u>	<b>A202103 004</b>	Jun 10, 2016
<b><u>SPRYCEL</u></b>				
<b>AB</b>	+ BRISTOL MYERS SQUIBB	<u>20MG</u>	<b>N021986 001</b>	Jun 28, 2006
<b>AB</b>	+	<u>50MG</u>	<b>N021986 002</b>	Jun 28, 2006
<b>AB</b>	+	<u>70MG</u>	<b>N021986 003</b>	Jun 28, 2006
<b>AB</b>	+!	<u>100MG</u>	<b>N021986 004</b>	May 30, 2008
	+	80MG	N021986 005	Oct 28, 2010
	+	140MG	N021986 006	Oct 28, 2010

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

**CERUBIDINE**

<b>AP</b>	! WEST-WARD PHARMS INT	<u>EQ 20MG BASE/VIAL</u>	<b>A064103 001</b>	Feb 03, 1995
<b><u>DAUNORUBICIN HYDROCHLORIDE</u></b>				
<b>AP</b>	FRESENIUS KABI USA	<u>EQ 20MG BASE/VIAL</u>	<b>A065000 001</b>	May 25, 1999
<b>AP</b>	TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<b>A065035 001</b>	Jan 24, 2000
<b>AP</b>	+! WEST-WARD PHARMS INT	<u>EQ 5MG BASE/ML</u>	<b>N050731 001</b>	Jan 30, 1998
	FRESENIUS KABI USA	EQ 5MG BASE/VIAL	A065034 001	Nov 20, 2001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-114 (of 436)

DECITABINE

INJECTABLE; INTRAVENOUS

DACOGEN

AP +! OTSUKA PHARM CO LTD 50MG/VIAL

N021790 001 May 02, 2006

DECITABINE

AP ACCORD HLTHCARE 50MG/VIAL

A203475 001 Feb 27, 2017

AP CHEMI SPA 50MG/VIAL

A206033 001 Sep 22, 2017

AP CIPLA LTD 50MG/VIAL

A208601 001 Nov 16, 2017

AP DR REDDYS LABS LTD 50MG/VIAL

A203131 001 Jul 11, 2013

AP PHARMASCIENCE INC 50MG/VIAL

A204607 001 May 31, 2017

AP SANDOZ INC 50MG/VIAL

A202969 001 Aug 28, 2014

POWDER; INTRAVENOUS

DECITABINE

+! SUN PHARMA GLOBAL 50MG/VIAL

N205582 001 Jan 28, 2014

DEFERASIROX

GRANULE; ORAL

JADENU SPRINKLE

+ NOVARTIS PHARMS CORP

90MG

N207968 001 May 18, 2017

+ 180MG

N207968 002 May 18, 2017

+! 360MG

N207968 003 May 18, 2017

TABLET; ORAL

JADENU

+ NOVARTIS PHARMS CORP

90MG

N206910 001 Mar 30, 2015

+ 180MG

N206910 002 Mar 30, 2015

+! 360MG

N206910 003 Mar 30, 2015

TABLET, FOR SUSPENSION; ORAL

DEFERASIROX

AB ACTAVIS ELIZABETH 125MG

A203560 001 Jan 26, 2016

AB 250MG

A203560 002 Jan 26, 2016

AB 500MG

A203560 003 Jan 26, 2016

EXJADE

AB + NOVARTIS 125MG

N021882 001 Nov 02, 2005

AB + 250MG

N021882 002 Nov 02, 2005

AB +! 500MG

N021882 003 Nov 02, 2005

DEFERIPRONE

SOLUTION; ORAL

FERRIPROX

+! APOPHARMA INC 100MG/ML

N208030 001 Sep 09, 2015

TABLET; ORAL

FERRIPROX

+! APOPHARMA INC 500MG

N021825 001 Oct 14, 2011

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

AP FRESENIUS KABI USA 500MG/VIAL

A078718 001 Sep 15, 2009

AP 2GM/VIAL

A078718 002 Sep 15, 2009

AP GLAND PHARMA LTD 500MG/VIAL

A207384 001 Sep 29, 2017

AP 2GM/VIAL

A207384 002 Sep 29, 2017

AP HOSPIRA 500MG/VIAL

A076019 001 Mar 17, 2004

AP 2GM/VIAL

A076019 002 Mar 17, 2004

AP WEST-WARD PHARMS INT 500MG/VIAL

A078086 001 May 30, 2007

AP 2GM/VIAL

A078086 002 May 30, 2007

DESFERAL

AP +! NOVARTIS 500MG/VIAL

N016267 001

AP +! 2GM/VIAL

N016267 002 May 25, 2000

DEFIBROTIDE SODIUM

SOLUTION; IV (INFUSION)

DEFITELIO

+! JAZZ PHARMS INC 200MG/2.5ML (80MG/ML)

N208114 001 Mar 30, 2016

DEFLAZACORT

SUSPENSION; ORAL

EMFLAZA

+! PTC THERAP 22.75MG/ML

N208685 001 Feb 09, 2017

TABLET; ORAL

EMFLAZA

+ PTC THERAP 6MG

N208684 001 Feb 09, 2017

+ 18MG

N208684 002 Feb 09, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-115 (of 436)

DEFLAZACORT

TABLET;ORAL  
 EMFLAZA  
 +  
 +!

30MG	N208684 003	Feb 09, 2017
36MG	N208684 004	Feb 09, 2017

DEGARELIX ACETATE

POWDER;SUBCUTANEOUS  
 FIRMAGON  
 + FERRING  
 +!

EQ 80MG BASE/VIAL	N022201 001	Dec 24, 2008
EQ 120MG BASE/VIAL	N022201 002	Dec 24, 2008

DELAFLOXACIN MEGLUMINE

POWDER;IV (INFUSION)

BAXDELA  
 +! MELINTA

EQ 300MG BASE/VIAL	N208611 001	Jun 19, 2017
--------------------	-------------	--------------

TABLET;ORAL

BAXDELA  
 +! MELINTA

EQ 450MG BASE	N208610 001	Jun 19, 2017
---------------	-------------	--------------

DELAVIDINE MESYLATE

TABLET;ORAL  
 DESCRIPTOR

+ VIIV HLTHCARE	100MG	N020705 001	Apr 04, 1997
+!	200MG	N020705 002	Jul 14, 1999

DEMECTOCYCLINE HYDROCHLORIDE

TABLET;ORAL

DEMECTOCYCLINE HYDROCHLORIDE

<u>AB</u>	AKORN	<u>150MG</u>	<u>A065389 001</u>	Dec 01, 2008
<u>AB</u>		<u>300MG</u>	<u>A065389 002</u>	Dec 01, 2008
<u>AB</u>	AMNEAL PHARM	<u>150MG</u>	<u>A065425 001</u>	Feb 27, 2008
<u>AB</u>	!	<u>300MG</u>	<u>A065425 002</u>	Feb 27, 2008
<u>AB</u>	BARR	<u>150MG</u>	<u>A065171 001</u>	Dec 13, 2004
<u>AB</u>		<u>300MG</u>	<u>A065171 002</u>	Dec 13, 2004
<u>AB</u>	EPIC PHARMA LLC	<u>150MG</u>	<u>A065447 001</u>	Aug 18, 2015
<u>AB</u>		<u>300MG</u>	<u>A065447 002</u>	Aug 18, 2015

DEOXYCHOLIC ACID

SOLUTION;SUBCUTANEOUS  
 KYBELLA

+! KYTHERA BIOPHARMS	20MG/2ML (10MG/ML)	N206333 001	Apr 29, 2015
----------------------	--------------------	-------------	--------------

DESFLURANE

LIQUID;INHALATION  
 SUPRANE

+! BAXTER HLTHCARE	99.9%	N020118 001	Sep 18, 1992
--------------------	-------	-------------	--------------

DESIPRAMINE HYDROCHLORIDE

TABLET;ORAL

DESIPRAMINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS TOTOWA	<u>10MG</u>	<u>A074430 001</u>	Feb 09, 1996
<u>AB</u>		<u>25MG</u>	<u>A071601 001</u>	Jun 05, 1987
<u>AB</u>		<u>50MG</u>	<u>A071588 001</u>	Jun 05, 1987
<u>AB</u>		<u>75MG</u>	<u>A071602 001</u>	Oct 05, 1987
<u>AB</u>		<u>100MG</u>	<u>A071766 001</u>	Oct 05, 1987
<u>AB</u>		<u>150MG</u>	<u>A074430 002</u>	Feb 09, 1996
<u>AB</u>	AMNEAL PHARMS CO	<u>10MG</u>	<u>A208105 001</u>	Mar 17, 2016
<u>AB</u>		<u>25MG</u>	<u>A208105 002</u>	Mar 17, 2016
<u>AB</u>		<u>50MG</u>	<u>A208105 003</u>	Mar 17, 2016
<u>AB</u>		<u>75MG</u>	<u>A208105 004</u>	Mar 17, 2016
<u>AB</u>		<u>100MG</u>	<u>A208105 005</u>	Mar 17, 2016
<u>AB</u>		<u>150MG</u>	<u>A208105 006</u>	Mar 17, 2016
<u>AB</u>	COREPHARMA	<u>10MG</u>	<u>A205153 001</u>	Oct 28, 2016
<u>AB</u>		<u>25MG</u>	<u>A205153 002</u>	Oct 28, 2016
<u>AB</u>		<u>50MG</u>	<u>A205153 003</u>	Oct 28, 2016
<u>AB</u>		<u>75MG</u>	<u>A205153 004</u>	Oct 28, 2016
<u>AB</u>		<u>100MG</u>	<u>A205153 005</u>	Oct 28, 2016
<u>AB</u>		<u>150MG</u>	<u>A205153 006</u>	Oct 28, 2016
<u>AB</u>	HERITAGE PHARMS INC	<u>10MG</u>	<u>A207433 001</u>	May 05, 2016
<u>AB</u>		<u>25MG</u>	<u>A207433 002</u>	May 05, 2016
<u>AB</u>		<u>50MG</u>	<u>A207433 003</u>	May 05, 2016
<u>AB</u>		<u>75MG</u>	<u>A207433 004</u>	May 05, 2016
<u>AB</u>		<u>100MG</u>	<u>A207433 005</u>	May 05, 2016
<u>AB</u>		<u>150MG</u>	<u>A207433 006</u>	May 05, 2016
<u>AB</u>	INGENUS PHARMS LLC	<u>10MG</u>	<u>A204963 001</u>	Dec 26, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-116 (of 436)

DESIPRAMINE HYDROCHLORIDE

TABLET;ORAL

DESIPRAMINE HYDROCHLORIDE

<u>AB</u>		<u>25MG</u>	<u>A204963 002</u>	Dec 26, 2017
<u>AB</u>		<u>50MG</u>	<u>A204963 003</u>	Dec 26, 2017
<u>AB</u>		<u>75MG</u>	<u>A204963 004</u>	Dec 26, 2017
<u>AB</u>		<u>100MG</u>	<u>A204963 005</u>	Dec 26, 2017
<u>AB</u>		<u>150MG</u>	<u>A204963 006</u>	Dec 26, 2017
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A072099 001</u>	May 24, 1988
<u>AB</u>		<u>25MG</u>	<u>A072100 001</u>	May 24, 1988
<u>AB</u>		<u>50MG</u>	<u>A072101 001</u>	May 24, 1988
<u>AB</u>		<u>75MG</u>	<u>A072102 001</u>	Jun 20, 1988
<u>AB</u>		<u>100MG</u>	<u>A072103 001</u>	Jun 20, 1988
<u>AB</u>		<u>150MG</u>	<u>A072104 001</u>	Jun 20, 1988

NORPRAMIN

<u>AB</u>	+	US PHARM HOLDINGS	<u>10MG</u>	<u>N014399 007</u>	Feb 11, 1982
<u>AB</u>	+		<u>25MG</u>	<u>N014399 001</u>	
<u>AB</u>	+		<u>50MG</u>	<u>N014399 003</u>	
<u>AB</u>	+		<u>75MG</u>	<u>N014399 004</u>	
<u>AB</u>	+!		<u>100MG</u>	<u>N014399 005</u>	
<u>AB</u>	+		<u>150MG</u>	<u>N014399 006</u>	

DESIPRAMINE HYDROCHLORIDE

ANI PHARMS INC	25MG	A071803 002	Dec 08, 1987
	50MG	A071803 003	Dec 08, 1987
	75MG	A071803 004	Dec 08, 1987
	150MG	A071803 005	May 29, 1997

DESIKUDIN RECOMBINANT

INJECTABLE;SUBCUTANEOUS

IPIVASK

+!	VALEANT PHARMS NORTH	15MG/VIAL
----	----------------------	-----------

N021271 001 Apr 04, 2003

DESLORATADINE

SOLUTION;ORAL

CLARINEX

<u>AA</u>	+!	MERCK SHARP DOHME	<u>0 . 5MG /ML</u>	<u>N021300 001</u>	Sep 01, 2004
<u>AA</u>		TARO PHARM	<u>0 . 5MG /ML</u>	<u>A202936 001</u>	May 26, 2016
<u>AA</u>		TARO PHARM IND	<u>0 . 5MG /ML</u>	<u>A202592 001</u>	Jun 30, 2015

TABLET;ORAL

CLARINEX

<u>AB</u>	+!	MERCK SHARP DOHME	<u>5MG</u>	<u>N021165 001</u>	Dec 21, 2001
<u>AB</u>		DESLORATADINE			
<u>AB</u>		BELCHER PHARMS	<u>5MG</u>	<u>A078355 001</u>	Apr 19, 2012
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG</u>	<u>A078365 001</u>	Mar 08, 2011
<u>AB</u>		LUPIN PHARMS	<u>5MG</u>	<u>A078352 001</u>	Oct 25, 2010
<u>AB</u>		MYLAN PHARMS INC	<u>5MG</u>	<u>A078351 001</u>	Feb 10, 2012
<u>AB</u>		ORCHID HLTHCARE	<u>5MG</u>	<u>A078357 001</u>	Feb 19, 2010
<u>AB</u>		PERRIGO R AND D	<u>5MG</u>	<u>A078361 001</u>	Dec 22, 2011
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A078364 001</u>	Dec 03, 2010
<u>AB</u>		SUN PHARM IND	<u>5MG</u>	<u>A078359 001</u>	Nov 16, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

CLARINEX

<u>AB</u>	+	MERCK SHARP DOHME	<u>2 . 5MG</u>	<u>N021312 002</u>	Jul 14, 2005
<u>AB</u>	+!		<u>5MG</u>	<u>N021312 001</u>	Jun 26, 2002
<u>AB</u>		DESLORATADINE			

<u>AB</u>	REDDYS	<u>2 . 5MG</u>	<u>A078367 001</u>	Jul 12, 2010
<u>AB</u>		<u>5MG</u>	<u>A078367 002</u>	Jul 12, 2010

DESLORATADINE; PSEUDOEPHENDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CLARINEX D 24 HOUR

<u>AB</u>	+!	MERCK SHARP DOHME	<u>5MG;240MG</u>	<u>N021605 001</u>	Mar 03, 2005
<u>AB</u>		DESLORATADINE AND PSEUDOEPHENDRINE SULFATE 24 HOUR			
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG;240MG</u>	<u>A078366 001</u>	Apr 26, 2011

CLARINEX-D 12 HOUR

+!	MERCK SHARP DOHME	2 . 5MG;120MG
----	-------------------	---------------

N021313 001 Feb 01, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-117 (of 436)

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

AP +! FERRING PHARMS INC 0.004MG/ML

N018938 001 Mar 30, 1984

DESMOPRESSIN ACETATE

AP SAGENT PHARMS 0.004MG/ML

A204695 001 Aug 22, 2017

AP 0.004MG/ML

A204751 001 Aug 22, 2017

AP SUN PHARM INDUS LTD 0.004MG/ML

A091280 001 Jan 25, 2013

SOLUTION; NASAL

DDAVP

AB +! FERRING PHARMS INC 0.01%

N017922 001

DESMOPRESSIN ACETATE

AB SUN PHARM INDUS 0.01%

A077212 001 Apr 12, 2012

SPRAY, METERED; NASAL

DDAVP (NEEDS NO REFRIGERATION)

AB +! FERRING PHARMS INC 0.01MG/SPRAY

N017922 003 Aug 07, 1996

DESMOPRESSIN ACETATE

AB ! BAUSCH AND LOMB 0.01MG/SPRAY

A074830 001 Jan 25, 1999

DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)

AB APOTEX INC 0.01MG/SPRAY

A076703 001 Jan 27, 2005

AB SUN PHARMA GLOBAL 0.01MG/SPRAY

A078271 001 Dec 23, 2013

AB ZYDUS PHARMS USA 0.01MG/SPRAY

A091345 001 Oct 03, 2017

INC

MINIRIN

AB +! FERRING 0.01MG/SPRAY

N021333 001 Sep 16, 2002

NOCTIVA

+ AVADEL SPECIET 0.00083MG/SPRAY

N201656 001 Mar 03, 2017

+! STIMATE (NEEDS NO REFRIGERATION) 0.00166MG/SPRAY

N201656 002 Mar 03, 2017

+! FERRING PHARMS INC 0.15MG/SPRAY

N020355 002 Oct 24, 2007

TABLET; ORAL

DDAVP

AB + FERRING PHARMS INC 0.1MG

N019955 001 Sep 06, 1995

AB +! 0.2MG

N019955 002 Sep 06, 1995

DESMOPRESSIN ACETATE

AB ACTAVIS LABS FL INC 0.1MG

A076470 001 Jul 01, 2005

AB 0.2MG

A076470 002 Jul 01, 2005

AB APOTEX INC 0.1MG

A077414 001 Mar 07, 2006

AB 0.2MG

A077414 002 Mar 07, 2006

AB GLENMARK PHARMS LTD 0.1MG

A201831 001 May 28, 2015

AB 0.2MG

A201831 002 May 28, 2015

AB HERITAGE PHARMA 0.1MG

A207880 001 May 26, 2017

AB 0.2MG

A207880 002 May 26, 2017

AB IMPAX LABS INC 0.1MG

A077122 001 Jan 25, 2006

AB 0.2MG

A077122 002 Jan 25, 2006

AB MYLAN PHARMS INC 0.1MG

A200653 001 Jun 27, 2014

AB 0.2MG

A200653 002 Jun 27, 2014

DESOGESTREL; ETHINYLN DROSTROL

TABLET; ORAL-28

BEKYREE

AB LUPIN LTD 0.15MG, N/A; 0.02MG, 0.01MG

A202226 001 Aug 12, 2015

CYCLESSA

AB +! ASPEN GLOBAL INC 0.1MG, 0.125MG, 0.15MG; 0.025MG, 0.025MG, 0.

N021090 001 Dec 20, 2000

025MG

DESOGEN

AB ORGANON USA INC 0.15MG; 0.03MG

N020071 002 Dec 10, 1992

DESOGESTREL AND ETHINYLN DROSTROL

AB ACCORD HLTHCARE 0.15MG, N/A; 0.02MG, 0.01MG

A209170 001 Jun 05, 2017

AB AUROBINDO PHARMA LTD 0.15MG, N/A; 0.02MG, 0.01MG

A206853 001 Mar 22, 2017

AB ! DURAMED PHARMS BARR 0.15MG; 0.03MG

A075256 002 Aug 12, 1999

AB MAYNE PHARMA 0.15MG, N/A; 0.02MG, 0.01MG

A076916 001 Dec 29, 2008

AB 0.1MG, 0.125MG, 0.15MG; 0.025MG, 0.025MG, 0.

A077182 001 Jan 24, 2006

025MG

AB MYLAN LABS LTD 0.15MG, N/A; 0.02MG, 0.01MG

A202296 001 Aug 30, 2013

AB 0.15MG; 0.03MG

A202085 001 May 20, 2015

AB NOVAST LABS LTD 0.15MG; 0.03MG

A091234 001 Jul 12, 2013

AB WATSON LABS 0.15MG; 0.03MG

A076915 001 Jul 29, 2005

EMOQUETTE

AB VINTAGE PHARMS LLC 0.15MG; 0.03MG

A076675 001 Feb 25, 2011

ENSKYCE

AB LUPIN LTD 0.15MG; 0.03MG

A201887 001 Mar 07, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-118 (of 436)

DESOGESTREL; ETHINYL ESTRADIOL

TABLET;ORAL-28

ISIBLOOM

AB LABS LEON FARMA 0.15MG;0.03MG A202789\_001 Aug 12, 2015

KALLIGA

AB AUROBINDO PHARMA LTD 0.15MG;0.03MG A207081\_001 May 17, 2017

KARIVA

AB ! BARR 0.15MG,N/A;0.02MG,0.01MG A075863\_001 Apr 05, 2002

KIMIDESS

AB VINTAGE PHARMS 0.15MG,N/A;0.02MG,0.01MG A076681\_001 Apr 30, 2015

PIMTREA

AB NOVAST LABS LTD 0.15MG,N/A;0.02MG,0.01MG A091247\_001 Aug 01, 2013

VELIVET

AB DURAMED PHARMS BARR 0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG A076455\_001 Feb 24, 2004

VIORELE

AB GLENMARK GENERICS 0.15MG,N/A;0.02MG,0.01MG A091346\_001 Apr 02, 2012

VOLNEA

AB LABS LEON FARMA 0.15MG,N/A;0.02MG,0.01MG A202689\_001 Sep 09, 2016

DESONIDE

AEROSOL, FOAM;TOPICAL

VERDESO

+! AQUA PHARMS 0.05% N021978 001 Sep 19, 2006

CREAM;TOPICAL

DESONIDE

AB G AND W LABS INC 0.05% A074027\_001 Sep 28, 1992

AB GLENMARK PHARMS 0.05% A209729\_001 Jul 24, 2017

AB +! PERRIGO NEW YORK 0.05% N017010\_001

AB TARO 0.05% A073548\_001 Jun 30, 1992

DESOWEN

AB GALDERMA LABS LP 0.05% N019048\_001 Dec 14, 1984

GEL;TOPICAL

DESONATE

+! BAYER HLTHCARE 0.05% N021844 001 Oct 20, 2006

LOTION;TOPICAL

DESONIDE

AB FOUGERA PHARMS 0.05% A075860\_001 Mar 19, 2002

AB GLENMARK PHARMS 0.05% A209494\_001 Sep 26, 2017

AB TARO PHARM 0.05% A202161\_001 Oct 31, 2014

AB TELIGENT PHARMA INC 0.05% A207855\_001 Sep 28, 2017

DESOWEN

AB ! GALDERMA LABS LP 0.05% A072354\_001 Jan 24, 1992

OINTMENT;TOPICAL

DESONIDE

AB FOUGERA PHARMS 0.05% A075751\_001 Mar 12, 2001

AB GLENMARK PHARMS LTD 0.05% A209996\_001 Sep 15, 2017

AB HI-TECH PHARMACAL 0.05% A208836\_001 Mar 27, 2017

AB +! PERRIGO NEW YORK 0.05% N017426\_001

AB TARO 0.05% A074254\_001 Aug 03, 1994

DESOWEN

AB GALDERMA LABS LP 0.05% A071425\_001 Jun 15, 1988

DESOXIMETASONE

CREAM;TOPICAL

DESOXIMETASONE

AB ACTAVIS MID ATLANTIC 0.25% A205082\_001 Sep 04, 2015

AB AKORN 0.05% A203787\_001 Jan 06, 2017

AB 0.25% A203234\_001 Jun 12, 2015

AB FOUGERA PHARMS 0.25% A078369\_001 Jun 29, 2010

AB LUPIN ATLANTIS 0.05% A208163\_001 Jan 10, 2017

AB 0.25% A208164\_001 Jan 09, 2017

AB PERRIGO NEW YORK 0.25% A076510\_001 Jul 01, 2003

TOPICORT

AB ! TARO 0.05% A073210\_001 Nov 30, 1990

AB ! 0.25% A073193\_001 Nov 30, 1990

GEL;TOPICAL

DESOXIMETASONE

AB AKORN 0.05% A090727\_001 Mar 10, 2011

AB PERRIGO NEW YORK 0.05% A077552\_001 Jan 09, 2006

AB RISING PHARMS INC 0.05% A204675\_001 Aug 12, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-119 (of 436)

DESOXIMETASONE

GEL;TOPICAL

**TOPICORT**

**AB** ! TARO **0.05%**

**A074904 001** Jul 14, 1998

OINTMENT;TOPICAL

**DESOXIMETASONE**

<b>AB</b>	ACTAVIS MID ATLANTIC	<b>0.25%</b>
<b>AB</b>	AKORN	<b>0.25%</b>
<b>AB</b>	FOUGERA PHARMS	<b>0.25%</b>
<b>AB</b>	G AND W LABS INC	<b>0.25%</b>
<b>AB</b>	GLENMARK GENERICS	<b>0.25%</b>
<b>AB</b>	LUPIN ATLANTIS	<b>0.05%</b>
<b>AB</b>		<b>0.25%</b>
<b>AB</b>	NOVEL LABS INC	<b>0.25%</b>
<b>AB</b>	PERRIGO ISRAEL	<b>0.25%</b>
<b>AB</b>	RISING PHARMS INC	<b>0.25%</b>
<b>AB</b>	TELIGENT PHARMA INC	<b>0.25%</b>
<b>AB</b>	ZYDUS PHARMS USA INC	<b>0.25%</b>

<b>A204965 001</b>	Nov 07, 2016
<b>A201005 001</b>	Apr 24, 2014
<b>A078657 001</b>	Sep 28, 2012
<b>A206740 001</b>	Dec 23, 2016
<b>A202838 001</b>	Sep 20, 2013
<b>A208044 001</b>	Dec 12, 2016
<b>A208104 001</b>	Dec 01, 2016
<b>A206792 001</b>	May 10, 2016
<b>A077770 001</b>	Apr 20, 2015
<b>A204272 001</b>	Nov 30, 2016
<b>A208101 001</b>	Feb 25, 2016
<b>A205206 001</b>	Sep 19, 2017

**TOPICORT**

**AB** +! TARO **0.05%**

**N018594 001** Jan 17, 1985  
**A074286 001** Jun 07, 1996

SPRAY;TOPICAL

**DESOXIMETASONE**

**AT** PERRIGO ISRAEL **0.25%**

**A206441 001** Jan 20, 2017

**TOPICORT**

**AT** +! TARO **0.25%**

**N204141 001** Apr 11, 2013

DESVENLAFAXINE

TABLET, EXTENDED RELEASE;ORAL  
 DESVENLAFAXINE

BC	+!	ALEMBIC PHARMS LTD	50MG
BC	+		100MG
KHEDEZLA			
BC		OSMOTICA PHARM CORP	50MG
BC			100MG

<b>N204150 001</b>	Mar 04, 2013
<b>N204150 002</b>	Mar 04, 2013
<b>N204683 001</b>	Jul 10, 2013
<b>N204683 002</b>	Jul 10, 2013

DESVENLAFAXINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL  
 DESVENLAFAXINE

+!	SUN PHARMA GLOBAL	EQ 50MG BASE
+!		EQ 100MG BASE

<b>N205583 001</b>	Jan 28, 2014
<b>N205583 002</b>	Jan 28, 2014

DESVENLAFAXINE SUCCINATE

TABLET, EXTENDED RELEASE;ORAL

**DESVENLAFAXINE SUCCINATE**

<b>AB</b>	ACTAVIS LABS FL	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	ALEMBIC PHARMS LTD	<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	LUPIN LTD	<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	MYLAN PHARMS INC	<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	SANDOZ INC	<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	WEST-WARD PHARMS INT	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	ZYDUS PHARMS USA INC	<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>PRISTIQ</b>		
<b>AB</b>	+ WYETH PHARMS INC	<b>EQ 25MG BASE</b>
<b>AB</b>	+	<b>EQ 50MG BASE</b>
<b>AB</b>	+	<b>EQ 100MG BASE</b>

<b>A204065 001</b>	Jul 29, 2016
<b>A204065 002</b>	Jul 29, 2016
<b>A204065 003</b>	Jul 29, 2016
<b>A204003 001</b>	Jun 29, 2015
<b>A204003 002</b>	Jun 29, 2015
<b>A204172 001</b>	Jun 29, 2015
<b>A204172 002</b>	Jun 29, 2015
<b>A204095 001</b>	Jun 29, 2015
<b>A204095 002</b>	Jun 29, 2015
<b>A204028 001</b>	Jun 29, 2015
<b>A204028 002</b>	Jun 29, 2015
<b>A204082 002</b>	Aug 28, 2017
<b>A204082 001</b>	Feb 16, 2016
<b>A204083 001</b>	Feb 16, 2016
<b>A204020 001</b>	Oct 11, 2017
<b>A204020 002</b>	Oct 11, 2017
<b>N021992 003</b>	Aug 20, 2014
<b>N021992 001</b>	Feb 29, 2008
<b>N021992 002</b>	Feb 29, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-120 (of 436)

DEUTETRABENAZINE

TABLET;ORAL

AUSTEDO

+ TEVA BRANDED PHARM	6MG	N208082 001 Apr 03, 2017
+	9MG	N208082 002 Apr 03, 2017
+!	12MG	N208082 003 Apr 03, 2017

DEXAMETHASONE

CONCENTRATE;ORAL

DEXAMETHASONE INTENSOL

! WEST-WARD PHARMS	1MG/ML	A088252 001 Sep 01, 1983
INT		

ELIXIR;ORAL

DEXAMETHASONE

<b>AA</b>	LYNE	<b>0 .5MG/5ML</b>	<b>A090891 001</b> Jul 12, 2011
<b>AA</b> !	STI PHARMA LLC	<b>0 .5MG/5ML</b>	<b>A084754 001</b>
<b>AA</b>	VINTAGE PHARMS	<b>0 .5MG/5ML</b>	<b>A091188 001</b> May 11, 2011
<b>AA</b>	WOCKHARDT BIO AG	<b>0 .5MG/5ML</b>	<b>A088254 001</b> Jul 27, 1983

IMPLANT;INTRAVITREAL

OZURDEX

+! ALLERGAN	0 .7MG	N022315 001 Jun 17, 2009
-------------	--------	--------------------------

SOLUTION;ORAL

DEXAMETHASONE

! WEST-WARD PHARMS	0 .5MG/5ML	A088248 001 Sep 01, 1983
INT		

SUSPENSION/DROPS;OPHTHALMIC

MAXIDEX

+! NOVARTIS PHARMS CORP	0 .1%	N013422 001
-------------------------	-------	-------------

TABLET;ORAL

DEXAMETHASONE

<b>AB</b>	ECR	<b>1 .5MG</b>	<b>A040700 001</b> Aug 15, 2008
<b>AB</b>	LARKEN LABS INC	<b>1 .5MG</b>	<b>A201270 001</b> Jul 17, 2017
<b>AB</b>	WEST-WARD PHARMS INT	<b>1 .5MG</b>	<b>A084610 001</b>
BP	PAR PHARM	0 .5MG	A088148 001 Apr 28, 1983
BP		0 .75MG	A088160 001 Apr 28, 1983
BP		4MG	A088238 001 Apr 28, 1983
BP		6MG	A088481 001 Nov 28, 1983
BP	WEST-WARD PHARMS INT	0 .5MG	A084611 001
BP		0 .75MG	A084613 001
BP		1MG	A088306 001 Sep 15, 1983
BP		2MG	A087916 001 Aug 26, 1982
BP		4MG	A084612 001
BP !		6MG	A088316 001 Sep 15, 1983
BP	XSPIRE PHARMA	1 .5MG	A088237 001 Apr 28, 1983

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE;INJECTION

DEXAMETHASONE SODIUM PHOSPHATE

<b>AP</b>	AUROBINDO PHARMA LTD	<b>EQ 4MG PHOSPHATE/ML</b>	<b>A206781 001</b> Dec 01, 2015
<b>AP</b>	FRESENIUS KABI USA	<b>EQ 4MG PHOSPHATE/ML</b>	<b>A084916 001</b>
<b>AP</b>		<b>EQ 4MG PHOSPHATE/ML</b>	<b>A203129 001</b> Sep 30, 2015
<b>AP</b> !		<b>EQ 10MG PHOSPHATE/ML</b>	<b>A040572 001</b> Apr 22, 2005
<b>AP</b> !	LUITPOLD	<b>EQ 4MG PHOSPHATE/ML</b>	<b>A087440 001</b> Jul 21, 1982
<b>AP</b>	MYLAN LABS LTD	<b>EQ 4MG PHOSPHATE/ML</b>	<b>A040803 001</b> Aug 29, 2008
<b>AP</b>		<b>EQ 10MG PHOSPHATE/ML</b>	<b>A040802 001</b> Aug 29, 2008
<b>AP</b>	WEST-WARD PHARMS INT	<b>EQ 4MG PHOSPHATE/ML</b>	<b>A084282 001</b>
<b>AP</b> !		<b>EQ 10MG PHOSPHATE/ML</b>	<b>A087702 001</b> Sep 07, 1982
<b>DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE</b>			
<b>AP</b> !	FRESENIUS KABI USA	<b>EQ 10MG PHOSPHATE/ML</b>	<b>A040491 001</b> Apr 11, 2003
SOLUTION/DROPS;OPHTHALMIC, OTIC			
<b>DEXAMETHASONE SODIUM PHOSPHATE</b>			
<b>AT</b>	BAUSCH AND LOMB	<b>EQ 0 .1% PHOSPHATE</b>	<b>A040069 001</b> Jul 26, 1996
<b>AT</b> !	SANDOZ INC	<b>EQ 0 .1% PHOSPHATE</b>	<b>A088771 001</b> Jan 16, 1985

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-121 (of 436)

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

**MAXITROL**

<b>AT</b>	+!	NOVARTIS PHARMS CORP	<u>0.1%:EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<b>N050065 002</b>
<b>NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE</b>				
<b>AT</b>		BAUSCH AND LOMB	<u>0.1%:EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<b>A064063 001</b> Jul 25, 1994
<b>AT</b>		PERRIGO CO TENNESSEE	<u>0.1%:EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<b>A062938 001</b> Jul 31, 1989
SUSPENSION/DROPS;OPHTHALMIC				
<b>DEXASPORIN</b>				
<b>AT</b>		BAUSCH AND LOMB	<u>0.1%:EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<b>A064135 001</b> Sep 13, 1995
<b>MAXITROL</b>				
<b>AT</b>	+!	NOVARTIS PHARMS CORP	<u>0.1%:EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<b>N050023 002</b>
<b>AT</b>		SANDOZ INC	<u>0.1%:EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<b>A062341 001</b> May 22, 1984

DEXAMETHASONE; TOBRAMYCIN

OINTMENT;OPHTHALMIC

**TOBRADEX**

+!	NOVARTIS PHARMS CORP	<u>0.1%;0.3%</u>	
<b>TOBRADEX</b>			

TOBRADEX

<b>AB</b>	+!	NOVARTIS PHARMS CORP	<u>0.1%;0.3%</u>	<b>N050592 001</b> Aug 18, 1988
<b>TOBRAMYCIN AND DEXAMETHASONE</b>				
<b>AB</b>		BAUSCH AND LOMB	<u>0.1%;0.3%</u>	<b>A064134 001</b> Oct 27, 1999
		TOBRADEX ST		

+!	NOVARTIS PHARMS CORP	<u>0.05%;0.3%</u>
----	----------------------	-------------------

N050818 001 Feb 13, 2009

DEXCHLORPHENIRAMINE MALEATE

SYRUP;ORAL

DEXCHLORPHENIRAMINE MALEATE

!	WOCKHARDT BIO AG	<u>2MG/5ML</u>
---	------------------	----------------

A088251 001 Mar 23, 1984

DEXLANSOPRAZOLE

CAPSULE, DELAYED RELEASE;ORAL

**DEXILANT**

<b>AB</b>	+!	TAKEDA PHARMS USA	<u>60MG</u>	<b>N022287 002</b> Jan 30, 2009
<b>DEXLANSOPRAZOLE</b>				
<b>AB</b>		PAR PHARM INC	<u>60MG</u>	<b>A202294 001</b> Apr 19, 2017
		DEXILANT		

+	TAKEDA PHARMS USA	<u>30MG</u>
---	-------------------	-------------

N022287 001 Jan 30, 2009

DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

**DEXMEDETOMIDINE HYDROCHLORIDE**

<b>AP</b>		ACCORD HLTHCARE	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A204023 001</b> Feb 09, 2016
<b>AP</b>		ACTAVIS INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A204686 001</b> Oct 17, 2016
<b>AP</b>		AKORN INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A202585 001</b> Nov 24, 2014
<b>AP</b>		AUROBINDO PHARMA LTD	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A205867 001</b> Mar 17, 2016
<b>AP</b>		FRESENIUS KABI USA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A201072 001</b> Sep 18, 2015
<b>AP</b>		JIANGSU HENGRIUI MED	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A209065 001</b> Sep 19, 2017
<b>AP</b>		LUITPOLD PHARMS INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A203773 001</b> May 12, 2017
<b>AP</b>		MYLAN INSTITUTIONAL	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A202881 001</b> Aug 18, 2014
<b>AP</b>		PAR STERILE PRODUCTS	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A203972 001</b> Aug 18, 2014
<b>AP</b>		SANDOZ INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A091465 001</b> Jun 14, 2016
<b>AP</b>		SUN PHARM INDNS INC	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A202126 001</b> Aug 20, 2015
<b>AP</b>		TEVA PHARMS USA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A205272 001</b> Nov 28, 2017
<b>AP</b>		WEST-WARD PHARMS INT	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>A205046 001</b> Apr 26, 2017

**PRECEDEX**

<b>AP</b>	+!	HOSPIRA	<u>EQ 200MCG BASE/2ML (EQ 100MCG BASE/ML)</u>	<b>N021038 001</b> Dec 17, 1999
	+!		<u>EQ 80MCG BASE/20ML (EQ 4MCG BASE/ML)</u>	N021038 004 Nov 14, 2014
	+!		<u>EQ 200MCG BASE/50ML (EQ 4MCG BASE/ML)</u>	N021038 002 Mar 13, 2013
	+!		<u>EQ 400MCG BASE/100ML (EQ 4MCG BASE/ML)</u>	N021038 003 Mar 13, 2013

SOLUTION;IV (INFUSION)

DEXMEDETOMIDINE HYDROCHLORIDE

+!	HQ SPCLT PHARMA	<u>EQ 1MG BASE/10ML (EQ 100MCG BASE/ML)</u>	N206628 002 Oct 21, 2015
+!		<u>EQ 400MCG BASE/4ML (EQ 100MCG BASE/ML)</u>	N206628 001 Oct 21, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-122 (of 436)

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	IMPAK LABS INC	<u>5MG</u>	<u>A079108 001</u>	Aug 05, 2015
<u>AB</u>		<u>10MG</u>	<u>A079108 002</u>	Aug 05, 2015
<u>AB</u>		<u>15MG</u>	<u>A079108 003</u>	May 19, 2014
<u>AB</u>		<u>20MG</u>	<u>A079108 004</u>	Dec 21, 2015
<u>AB</u>		<u>25MG</u>	<u>A203614 001</u>	Jul 05, 2017
<u>AB</u>		<u>30MG</u>	<u>A079108 005</u>	Nov 21, 2013
<u>AB</u>		<u>35MG</u>	<u>A203614 002</u>	Jul 05, 2017
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>15MG</u>	<u>A078992 003</u>	Nov 18, 2013
<u>AB</u>		<u>30MG</u>	<u>A078992 004</u>	Nov 18, 2013
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A204266 001</u>	Aug 25, 2015
<u>AB</u>		<u>10MG</u>	<u>A204266 002</u>	Aug 25, 2015
<u>AB</u>		<u>15MG</u>	<u>A204266 003</u>	Aug 25, 2015
<u>AB</u>		<u>20MG</u>	<u>A204266 004</u>	Dec 21, 2015
<u>AB</u>		<u>30MG</u>	<u>A202580 001</u>	Aug 28, 2013
<u>AB</u>		<u>40MG</u>	<u>A204266 007</u>	Aug 25, 2015
<u>AB</u>	PAR PHARM INC	<u>5MG</u>	<u>A202842 001</u>	Nov 30, 2016
<u>AB</u>		<u>10MG</u>	<u>A202842 002</u>	Nov 30, 2016
<u>AB</u>		<u>15MG</u>	<u>A202842 003</u>	Nov 30, 2016
<u>AB</u>		<u>20MG</u>	<u>A202842 004</u>	Nov 30, 2016
<u>AB</u>		<u>25MG</u>	<u>A202842 005</u>	Nov 30, 2016
<u>AB</u>		<u>30MG</u>	<u>A202842 006</u>	Nov 30, 2016
<u>AB</u>		<u>35MG</u>	<u>A202842 007</u>	Nov 30, 2016
<u>AB</u>		<u>40MG</u>	<u>A202842 008</u>	Nov 30, 2016
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A078908 001</u>	Nov 19, 2013
<u>AB</u>		<u>10MG</u>	<u>A078908 002</u>	Nov 19, 2013
<u>AB</u>		<u>15MG</u>	<u>A078908 004</u>	May 19, 2014
<u>AB</u>		<u>20MG</u>	<u>A078908 003</u>	Nov 19, 2013
<u>AB</u>		<u>25MG</u>	<u>A202731 001</u>	Jul 05, 2017
<u>AB</u>		<u>30MG</u>	<u>A202731 003</u>	May 19, 2014
<u>AB</u>		<u>35MG</u>	<u>A202731 004</u>	Jul 05, 2017
<u>AB</u>		<u>40MG</u>	<u>A202731 002</u>	Nov 19, 2013
<b>FOCALIN XR</b>				
<u>AB</u>	+ NOVARTIS	<u>5MG</u>	<u>N021802 001</u>	May 26, 2005
<u>AB</u>	+	<u>10MG</u>	<u>N021802 002</u>	May 26, 2005
<u>AB</u>	+	<u>15MG</u>	<u>N021802 004</u>	Aug 01, 2006
<u>AB</u>	+	<u>20MG</u>	<u>N021802 003</u>	May 26, 2005
<u>AB</u>	+	<u>25MG</u>	<u>N021802 008</u>	Apr 21, 2011
<u>AB</u>	+	<u>30MG</u>	<u>N021802 005</u>	Oct 23, 2009
<u>AB</u>	+	<u>35MG</u>	<u>N021802 007</u>	Apr 21, 2011
<u>AB</u>	+!	<u>40MG</u>	<u>N021802 006</u>	Aug 11, 2010

TABLET;ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI INC	<u>2.5MG</u>	<u>A206931 001</u>	Dec 04, 2015
<u>AB</u>		<u>5MG</u>	<u>A206931 002</u>	Dec 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A206931 003</u>	Dec 04, 2015
<u>AB</u>	LANNETT CO INC	<u>2.5MG</u>	<u>A209468 001</u>	Sep 25, 2017
<u>AB</u>		<u>5MG</u>	<u>A209468 002</u>	Sep 25, 2017
<u>AB</u>		<u>10MG</u>	<u>A209468 003</u>	Sep 25, 2017
<u>AB</u>	NOVEL LABS INC	<u>2.5MG</u>	<u>A204534 001</u>	Dec 04, 2015
<u>AB</u>		<u>5MG</u>	<u>A204534 002</u>	Dec 04, 2015
<u>AB</u>		<u>10MG</u>	<u>A204534 003</u>	Dec 04, 2015
<u>AB</u>	RHODES PHARMS	<u>2.5MG</u>	<u>A208756 001</u>	Nov 20, 2017
<u>AB</u>		<u>5MG</u>	<u>A208756 002</u>	Nov 20, 2017
<u>AB</u>		<u>10MG</u>	<u>A208756 003</u>	Nov 20, 2017
<u>AB</u>	SUN PHARM INDUSTRIES	<u>2.5MG</u>	<u>A201231 001</u>	Sep 24, 2015
<u>AB</u>		<u>5MG</u>	<u>A201231 002</u>	Sep 24, 2015
<u>AB</u>		<u>10MG</u>	<u>A201231 003</u>	Sep 24, 2015
<u>AB</u>	TEVA PHARMS	<u>2.5MG</u>	<u>A077107 003</u>	Jan 29, 2007
<u>AB</u>		<u>5MG</u>	<u>A077107 001</u>	Jan 29, 2007
<u>AB</u>		<u>10MG</u>	<u>A077107 002</u>	Jan 29, 2007
<u>AB</u>	TRIS PHARMA INC	<u>2.5MG</u>	<u>A207901 001</u>	Aug 26, 2016
<u>AB</u>		<u>5MG</u>	<u>A207901 002</u>	Aug 26, 2016
<u>AB</u>		<u>10MG</u>	<u>A207901 003</u>	Aug 26, 2016
<b>FOCALIN</b>				
<u>AB</u>	+ NOVARTIS	<u>2.5MG</u>	<u>N021278 001</u>	Nov 13, 2001
<u>AB</u>	+	<u>5MG</u>	<u>N021278 002</u>	Nov 13, 2001
<u>AB</u>	+!	<u>10MG</u>	<u>N021278 003</u>	Nov 13, 2001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-123 (of 436)

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXRAZOXANE HYDROCHLORIDE

<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A207321 001</u>	Nov 28, 2016
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 250MG BASE/VIAL</u>	<u>A200752 001</u>	Oct 19, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A200752 002</u>	Oct 19, 2011
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 250MG BASE/VIAL</u>	<u>A076068 001</u>	Sep 28, 2004
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A076068 002</u>	Sep 28, 2004
	<u>ZINECARD</u>			
<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>EQ 250MG BASE/VIAL</u>	<u>N020212 001</u>	May 26, 1995
<u>AP</u>	+! TOTECT	<u>EQ 500MG BASE/VIAL</u>	<u>N020212 002</u>	May 26, 1995
	+! CLINIGEN HLTHCARE	EQ 500MG BASE/VIAL	N022025 001	Sep 06, 2007

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE

<u>AB</u>	+ IMPAX LABS INC	<u>5MG</u>	<u>N017078 001</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N017078 002</u>	
<u>AB</u>	+!	<u>15MG</u>	<u>N017078 003</u>	

DEXTROAMPHETAMINE SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>5MG</u>	<u>A203901 001</u>	Nov 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A203901 002</u>	Nov 30, 2012
<u>AB</u>		<u>15MG</u>	<u>A203901 003</u>	Nov 30, 2012
<u>AB</u>	MAYNE PHARMA	<u>5MG</u>	<u>A076137 001</u>	Jan 18, 2002
<u>AB</u>		<u>10MG</u>	<u>A076137 002</u>	Jan 18, 2002
<u>AB</u>		<u>15MG</u>	<u>A076137 003</u>	Jan 18, 2002
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A206735 001</u>	Jan 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A206735 002</u>	Jan 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A206735 003</u>	Jan 27, 2016
<u>AB</u>	NESHER PHARMS	<u>5MG</u>	<u>A209111 001</u>	Jun 27, 2017
<u>AB</u>		<u>10MG</u>	<u>A209111 002</u>	Jun 27, 2017
<u>AB</u>		<u>15MG</u>	<u>A209111 003</u>	Jun 27, 2017
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A076353 001</u>	May 06, 2003
<u>AB</u>		<u>10MG</u>	<u>A076353 002</u>	May 06, 2003
<u>AB</u>		<u>15MG</u>	<u>A076353 003</u>	May 06, 2003
<u>AB</u>	VINTAGE PHARMS	<u>5MG</u>	<u>A205673 001</u>	Oct 31, 2017
<u>AB</u>		<u>10MG</u>	<u>A205673 002</u>	Oct 31, 2017
<u>AB</u>		<u>15MG</u>	<u>A205673 003</u>	Oct 31, 2017

SOLUTION; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	! OUTLOOK PHARMS	<u>5MG/5ML</u>	<u>A040776 001</u>	Jan 29, 2008
<u>AA</u>	TRIS PHARMA INC	<u>5MG/5ML</u>	<u>A203644 001</u>	May 29, 2013

TABLET; ORAL

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202893 001</u>	Jul 31, 2013
<u>AA</u>		<u>10MG</u>	<u>A202893 002</u>	Jul 31, 2013
<u>AA</u>	AVANTHI INC	<u>5MG</u>	<u>A203548 001</u>	Nov 23, 2015
<u>AA</u>		<u>10MG</u>	<u>A203548 002</u>	Nov 23, 2015
<u>AA</u>	BARR	<u>5MG</u>	<u>A040361 001</u>	Jan 31, 2001
<u>AA</u>	!	<u>10MG</u>	<u>A040361 002</u>	Jan 31, 2001
<u>AA</u>	MIKART	<u>5MG</u>	<u>A090533 002</u>	Oct 25, 2011
<u>AA</u>		<u>10MG</u>	<u>A090533 004</u>	Oct 25, 2011
<u>AA</u>	NESHER PHARMS	<u>5MG</u>	<u>A206588 001</u>	Mar 28, 2016
<u>AA</u>		<u>10MG</u>	<u>A206588 002</u>	Mar 28, 2016
<u>AA</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204330 001</u>	Mar 16, 2016
<u>AA</u>		<u>10MG</u>	<u>A204330 002</u>	Mar 16, 2016
<u>AA</u>	SPECGX LLC	<u>5MG</u>	<u>A040436 001</u>	Jan 29, 2002
<u>AA</u>		<u>10MG</u>	<u>A040436 002</u>	Jan 29, 2002
<u>AA</u>	SUNRISE PHARM INC	<u>5MG</u>	<u>A210059 001</u>	Oct 18, 2017
<u>AA</u>		<u>10MG</u>	<u>A210059 002</u>	Oct 18, 2017
	MIKART	2.5MG	A090533 001	Oct 25, 2011
		7.5MG	A090533 003	Oct 25, 2011
		15MG	A090533 005	Oct 25, 2011
		20MG	A090533 006	Oct 25, 2011
		30MG	A090533 007	Oct 25, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-124 (of 436)

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH W/ DEXTROMETHORPHAN

AA G AND W LABS INC 15MG/5ML; 6.25MG/5ML A088762 001 Oct 31, 1984

PROMETHAZINE DM

AA ! VINTAGE 15MG/5ML; 6.25MG/5ML A040649 001 Feb 14, 2006

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

AA HI TECH PHARMA 15MG/5ML; 6.25MG/5ML A040027 001 Jul 31, 1996

PROMETHAZINE W/ DEXTROMETHORPHAN

AA WOCKHARDT BIO AG 15MG/5ML; 6.25MG/5ML A088864 001 Jan 04, 1985

DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE

CAPSULE; ORAL

DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE

AB ACTAVIS ELIZABETH 20MG;10MG A202934 001 Oct 10, 2017

NUEDEXTA

AB +! AVANIR PHARMS 20MG;10MG N021879 001 Oct 29, 2010

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

AP +! B BRAUN 10GM/100ML N019626 004 Feb 02, 1988

AP +! BAXTER HLTHCARE 10GM/100ML N016694 001

AP +! ICU MEDICAL INC 10GM/100ML N018080 001

DEXTROSE 20% IN PLASTIC CONTAINER

AP +! BAXTER HLTHCARE 20GM/100ML N017521 004

AP +! ICU MEDICAL INC 20GM/100ML N018564 001 Mar 23, 1982

DEXTROSE 30% IN PLASTIC CONTAINER

AP +! BAXTER HLTHCARE 30GM/100ML N017521 003

AP +! ICU MEDICAL INC 30GM/100ML N019345 001 Jan 26, 1985

DEXTROSE 40% IN PLASTIC CONTAINER

AP +! BAXTER HLTHCARE 40GM/100ML N017521 002

AP +! ICU MEDICAL INC 40GM/100ML N018562 001 Mar 23, 1982

DEXTROSE 5% IN PLASTIC CONTAINER

AP +! B BRAUN 50MG/ML N016730 002

AP +! 5GM/100ML N016730 001

AP +! 5GM/100ML N019626 002 Feb 02, 1988

AP +! BAXTER HLTHCARE 50MG/ML N016673 003 Oct 30, 1985

AP +! 50MG/ML N020179 002 Dec 07, 1992

AP +! 5GM/100ML N016673 001

AP +! 5GM/100ML N020179 001 Dec 07, 1992

AP +! FRESENIUS KABI USA 50MG/ML A207449 001 Oct 21, 2016

AP +! HOSPIRA 50MG/ML N019222 001 Jul 13, 1984

AP +! 5GM/100ML N019466 001 Jul 15, 1985

AP +! 5GM/100ML N019479 001 Sep 17, 1985

AP +! ICU MEDICAL INC 50MG/ML N016367 002

DEXTROSE 50% IN PLASTIC CONTAINER

AP +! BAXTER HLTHCARE 50GM/100ML N017521 001

AP +! 50GM/100ML N020047 001 Jul 02, 1991

AP +! ICU MEDICAL INC 50GM/100ML N018563 001 Mar 23, 1982

DEXTROSE 70% IN PLASTIC CONTAINER

AP +! BAXTER HLTHCARE 70GM/100ML N017521 006 Mar 26, 1982

AP +! 70GM/100ML N020047 003 Jul 02, 1991

AP +! ICU MEDICAL INC 70GM/100ML N018561 001 Mar 23, 1982

AP +! 70GM/100ML N019893 001 Dec 26, 1989

DEXTROSE 25%

+! HOSPIRA 250MG/ML

N019445 002 Nov 23, 1998

DEXTROSE 50%

+ HOSPIRA 500MG/ML

N019445 003 Sep 03, 2014

DEXTROSE 50% IN PLASTIC CONTAINER

+ HOSPIRA 500MG/ML

N019445 001 Jun 03, 1986

DEXTROSE 60% IN PLASTIC CONTAINER

+! BAXTER HLTHCARE 60GM/100ML

N017521 005 Mar 26, 1982

DEXTROSE; MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 5GM/100ML;32MG/100ML;128MG/100ML;234MG/100ML N017385 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-125 (of 436)

DEXTROSE; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;21MG/100ML;128MG/100ML;234MG/100ML N017610 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;31MG/100ML;130MG/100ML;26MG/100ML;320MG/100ML N019873 001 Jun 10, 1993

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 5GM/100ML;31MG/100ML;141MG/100ML;20MG/100ML;12MG/100ML;260MG/100ML N017484 001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION

TONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;30MG/100ML;141MG/100ML;15MG/100ML;260MG/100ML;25MG/100ML N019513 001 May 08, 1986

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;30MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML N017609 001

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

AP + BAXTER HLTHCARE 5GM/100ML;150MG/100ML N017634 001

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER

AP + BAXTER HLTHCARE 5GM/100ML;224MG/100ML N017634 003

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

AP + BAXTER HLTHCARE 5GM/100ML;300MG/100ML N017634 002

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER

AP B BRAUN 5GM/100ML;150MG/100ML N019699 004 Sep 29, 1989

POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER

AP B BRAUN 5GM/100ML;300MG/100ML N019699 006 Sep 29, 1989

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

AP ICU MEDICAL INC 5GM/100ML;224MG/100ML N018371 003

DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5GM/100ML;75MG/100ML

POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML;149MG/100ML

POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER

ICU MEDICAL INC 5GM/100ML;298MG/100ML

N017634 001

N017634 003

N017634 002

N019699 004 Sep 29, 1989

N019699 006 Sep 29, 1989

N018371 003

N017634 004

N018371 001

N018371 002

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ

AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;200MG/100ML N018037 006 Apr 13, 1982

5GM/100ML;150MG/100ML;200MG/100ML N018037 007 Apr 13, 1982

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)

AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;200MG/100ML N018037 004

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ

AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;200MG/100ML N018037 008 Apr 13, 1982

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)

AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;200MG/100ML N018037 001

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ

AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;200MG/100ML N018037 005 Apr 13, 1982

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ

AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;200MG/100ML N018037 009 Apr 13, 1982

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ

AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;200MG/100ML N018037 002

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)

AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;200MG/100ML N018037 003

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;330MG/100ML N018629 005 Mar 23, 1982

5GM/100ML;150MG/100ML;330MG/100ML N018629 002 Mar 23, 1982

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-126 (of 436)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEO IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;330MG/100ML	N018629	003	Mar 23, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEO IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;330MG/100ML	N018629	004	Mar 23, 1982
AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;330MG/100ML	N018629	006	Mar 23, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEO IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;330MG/100ML	N018629	007	Mar 23, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEO IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;330MG/100ML	N018629	008	Mar 23, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEO IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;330MG/100ML	N018629	001	Mar 23, 1982
<u>DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEO (K) IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;450MG/100ML	N018008	010	
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;75MG/100ML;200MG/100ML	N019630	008	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;75MG/100ML;330MG/100ML	N019630	014	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;75MG/100ML;450MG/100ML	N019630	020	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;75MG/100ML;900MG/100ML	N019630	026	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;150MG/100ML;200MG/100ML	N019630	010	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;150MG/100ML;330MG/100ML	N019630	016	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;150MG/100ML;450MG/100ML	N019630	022	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;150MG/100ML;900MG/100ML	N019630	028	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;300MG/100ML;200MG/100ML	N019630	012	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;300MG/100ML;330MG/100ML	N019630	018	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;300MG/100ML;450MG/100ML	N019630	024	Feb 17, 1988
<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP B BRAUN 5GM/100ML;300MG/100ML;900MG/100ML	N019630	030	Feb 17, 1988
<u>POTASSIUM CHLORIDE 10MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;450MG/100ML	N018008	005	Apr 28, 1982
AP 5GM/100ML;150MG/100ML;450MG/100ML	N018008	006	Apr 28, 1982
AP ICU MEDICAL INC 5GM/100ML;74.5MG/100ML;450MG/100ML	N018362	005	Mar 28, 1988
AP 5GM/100ML;74.5MG/100ML;450MG/100ML	N018362	009	Jul 05, 1983
<u>POTASSIUM CHLORIDE 10MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;75MG/100ML;900MG/100ML	N019308	004	Apr 05, 1985
AP 5GM/100ML;150MG/100ML;900MG/100ML	N019308	002	Apr 05, 1985
AP ICU MEDICAL INC 5GM/100ML;74.5MG/100ML;900MG/100ML	N019691	002	Mar 24, 1988
AP 5GM/100ML;149MG/100ML;900MG/100ML	N019691	004	Mar 24, 1988
<u>POTASSIUM CHLORIDE 15MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP ICU MEDICAL INC 5GM/100ML;224MG/100ML;450MG/100ML	N018362	006	Mar 28, 1988
<u>POTASSIUM CHLORIDE 15MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP ICU MEDICAL INC 5GM/100ML;224MG/100ML;900MG/100ML	N019691	006	Mar 24, 1988
<u>POTASSIUM CHLORIDE 20MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;450MG/100ML	N018008	007	Apr 28, 1982
AP ICU MEDICAL INC 5GM/100ML;149MG/100ML;450MG/100ML	N018362	010	Jul 05, 1983
AP 5GM/100ML;298MG/100ML;450MG/100ML	N018362	007	Mar 28, 1988
<u>POTASSIUM CHLORIDE 20MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;150MG/100ML;900MG/100ML	N019308	005	Apr 05, 1985
AP 5GM/100ML;300MG/100ML;900MG/100ML	N019308	003	Apr 05, 1985
AP ICU MEDICAL INC 5GM/100ML;149MG/100ML;900MG/100ML	N019691	005	Mar 24, 1988
AP 5GM/100ML;298MG/100ML;900MG/100ML	N019691	008	Mar 24, 1988
<u>POTASSIUM CHLORIDE 30MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;450MG/100ML	N018008	008	Apr 28, 1982
AP ICU MEDICAL INC 5GM/100ML;224MG/100ML;450MG/100ML	N018362	002	
<u>POTASSIUM CHLORIDE 30MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;224MG/100ML;900MG/100ML	N019308	006	Apr 05, 1985
AP ICU MEDICAL INC 5GM/100ML;224MG/100ML;900MG/100ML	N019691	007	Mar 24, 1988
<u>POTASSIUM CHLORIDE 40MEO IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE 5GM/100ML;300MG/100ML;450MG/100ML	N018008	009	Apr 28, 1982
AP ICU MEDICAL INC 5GM/100ML;298MG/100ML;450MG/100ML	N018362	003	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-127 (of 436)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

**POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER**

<b>AP</b>	BAXTER HLTHCARE	<u>5GM/100ML;300MG/100ML;900MG/100ML</u>	<b>N019308 007</b>	Apr 05, 1985
<b>AP</b>	ICU MEDICAL INC	<u>5GM/100ML;298MG/100ML;900MG/100ML</u>	<b>N019691 009</b>	Mar 24, 1988

**POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER**

<b>AP</b>	BAXTER HLTHCARE	<u>5GM/100ML;150MG/100ML;450MG/100ML</u>	<b>N018008 004</b>	
<b>AP</b>	ICU MEDICAL INC	<u>5GM/100ML;74.5MG/100ML;450MG/100ML</u>	<b>N018362 008</b>	Mar 28, 1988
<b>AP</b>		<u>5GM/100ML;149MG/100ML;450MG/100ML</u>	<b>N018362 004</b>	Mar 28, 1988

**POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER**

<b>AP</b>	BAXTER HLTHCARE	<u>5GM/100ML;150MG/100ML;900MG/100ML</u>	<b>N019308 001</b>	Apr 05, 1985
<b>AP</b>	ICU MEDICAL INC	<u>5GM/100ML;74.5MG/100ML;900MG/100ML</u>	<b>N019691 001</b>	Mar 24, 1988
<b>AP</b>		<u>5GM/100ML;149MG/100ML;900MG/100ML</u>	<b>N019691 003</b>	Mar 24, 1988

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;37MG/100ML;200MG/100ML	N019630 031	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;37MG/100ML;450MG/100ML	N019630 037	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;37MG/100ML;900MG/100ML	N019630 043	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;110MG/100ML	N019630 001	Feb 17, 1988
---------	----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;200MG/100ML	N019630 007	Feb 17, 1988
---------	----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;330MG/100ML	N019630 013	Feb 17, 1988
---------	----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;450MG/100ML	N019630 019	Feb 17, 1988
---------	----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;900MG/100ML	N019630 025	Feb 17, 1988
---------	----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;75MG/100ML;200MG/100ML	N019630 032	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;75MG/100ML;450MG/100ML	N019630 038	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;75MG/100ML;900MG/100ML	N019630 044	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN	3.3GM/100ML;75MG/100ML;300MG/100ML	N019630 049	May 07, 1992
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;75MG/100ML;110MG/100ML	N019630 002	Feb 17, 1988
---------	----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;110MG/100ML;200MG/100ML	N019630 033	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;110MG/100ML;450MG/100ML	N019630 039	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;110MG/100ML;900MG/100ML	N019630 045	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN	3.3GM/100ML;110MG/100ML;300MG/100ML	N019630 050	May 07, 1992
---------	-------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;110MG/100ML;110MG/100ML	N019630 003	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;110MG/100ML;200MG/100ML	N019630 009	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;110MG/100ML;330MG/100ML	N019630 015	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;110MG/100ML;450MG/100ML	N019630 021	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;110MG/100ML;900MG/100ML	N019630 027	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;150MG/100ML;200MG/100ML	N019630 034	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;150MG/100ML;450MG/100ML	N019630 040	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;150MG/100ML;900MG/100ML	N019630 046	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN	3.3GM/100ML;150MG/100ML;300MG/100ML	N019630 051	May 07, 1992
---------	-------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;150MG/100ML;110MG/100ML	N019630 004	Feb 17, 1988
---------	-----------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;220MG/100ML;200MG/100ML	N019630 035	Feb 17, 1988
---------	------------------------------------	-------------	--------------

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;220MG/100ML;450MG/100ML	N019630 041	Feb 17, 1988
---------	------------------------------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-128 (of 436)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER  
 B BRAUN 10GM/100ML;220MG/100ML;900MG/100ML N019630 047 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 B BRAUN 3.3GM/100ML;220MG/100ML;300MG/100ML N019630 052 May 07, 1992  
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;220MG/100ML;110MG/100ML N019630 005 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;220MG/100ML;200MG/100ML N019630 011 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;220MG/100ML;330MG/100ML N019630 017 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;220MG/100ML;450MG/100ML N019630 023 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;220MG/100ML;900MG/100ML N019630 029 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER  
 B BRAUN 10GM/100ML;300MG/100ML;200MG/100ML N019630 036 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER  
 B BRAUN 10GM/100ML;300MG/100ML;450MG/100ML N019630 042 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER  
 B BRAUN 10GM/100ML;300MG/100ML;900MG/100ML N019630 048 Feb 17, 1988  
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 B BRAUN 3.3GM/100ML;300MG/100ML;300MG/100ML N019630 053 May 07, 1992  
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER  
 B BRAUN 5GM/100ML;300MG/100ML;110MG/100ML N019630 006 Feb 17, 1988  
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER  
 + ICU MEDICAL INC 5GM/100ML;74.5MG/100ML;225MG/100ML N018365 002 Jul 05, 1983  
 + 5GM/100ML;149MG/100ML;225MG/100ML N018365 006 Mar 28, 1988  
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;74.5MG/100ML;300MG/100ML N018876 001 Jan 17, 1986  
 5GM/100ML;149MG/100ML;300MG/100ML N018876 006 Mar 28, 1988  
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;224MG/100ML;225MG/100ML N018365 008 Mar 28, 1988  
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;224MG/100ML;300MG/100ML N018876 007 Mar 28, 1988  
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER  
 + ICU MEDICAL INC 5GM/100ML;149MG/100ML;225MG/100ML N018365 001  
 + 5GM/100ML;298MG/100ML;225MG/100ML N018365 009 Mar 28, 1988  
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;298MG/100ML;300MG/100ML N018876 008 Mar 28, 1988  
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;149MG/100ML;300MG/100ML N018876 002 Jan 17, 1986  
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;224MG/100ML;225MG/100ML N018365 003 Jul 05, 1983  
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;224MG/100ML;300MG/100ML N018876 003 Jan 17, 1986  
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;298MG/100ML;225MG/100ML N018365 004 Jul 05, 1983  
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;298MG/100ML;300MG/100ML N018876 004 Mar 28, 1988  
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;74.5MG/100ML;225MG/100ML N018365 005 Mar 28, 1988  
 5GM/100ML;149MG/100ML;225MG/100ML N018365 007 Mar 28, 1988  
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER  
 ICU MEDICAL INC 5GM/100ML;74.5MG/100ML;300MG/100ML N018876 005 Mar 28, 1988  
 5GM/100ML;149MG/100ML;300MG/100ML N018876 009 Mar 28, 1988

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>	
AP B BRAUN 2.5GM/100ML;450MG/100ML	N019631 004 Feb 24, 1988
AP +! BAXTER HLTHCARE 2.5GM/100ML;450MG/100ML	N016697 001
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>	
AP B BRAUN 5GM/100ML;200MG/100ML	N019631 007 Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>	
AP B BRAUN 5GM/100ML;330MG/100ML	N019631 008 Feb 24, 1988
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>	
AP B BRAUN 5GM/100ML;450MG/100ML	N019631 009 Feb 24, 1988
AP ICU MEDICAL INC 5GM/100ML;450MG/100ML	N017607 001
<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>	
AP B BRAUN 5GM/100ML;900MG/100ML	N019631 010 Feb 24, 1988
AP +! ICU MEDICAL INC 5GM/100ML;900MG/100ML	N017585 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-129 (of 436)

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;200MG/100ML</u>		N016689 001
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;330MG/100ML</u>		N016687 001
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;450MG/100ML</u>		N016683 001
<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP BAXTER HLTHCARE	<u>5GM/100ML;900MG/100ML</u>		N016678 001
DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;110MG/100ML		N019631 011 Feb 24, 1988
DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;200MG/100ML		N019631 012 Feb 24, 1988
DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;330MG/100ML		N019631 013 Feb 24, 1988
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;450MG/100ML		N019631 014 Feb 24, 1988
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML;900MG/100ML		N019631 015 Feb 24, 1988
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML;110MG/100ML		N019631 001 Feb 24, 1988
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML;330MG/100ML		N019631 003 Feb 24, 1988
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML;900MG/100ML		N019631 005 Feb 24, 1988
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
B BRAUN	3.3GM/100ML;300MG/100ML		N019631 016 Jan 19, 1990
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML		N019631 006 Feb 24, 1988
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER			
+! ICU MEDICAL INC	<u>5GM/100ML;225MG/100ML</u>		
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER			
+! ICU MEDICAL INC	<u>5GM/100ML;300MG/100ML</u>		N017799 001

DIATRIZOATE MEGLUMINE

SOLUTION; URETHRAL

CYSTOGRAPHIN			
+! BRACCO	30%		N010040 018
CYSTOGRAPHIN DILUTE			
+! BRACCO	18%		N010040 022 Nov 09, 1982

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

RENOGRAFIN-76			
+! BRACCO	66%;10%		N010040 001

SOLUTION; ORAL, RECTAL

<b>GASTROGRAFIN</b>			
---------------------	--	--	--

<b>AA +! BRACCO</b>	<u><b>66%;10%</b></u>		<b>N011245 003</b>
---------------------	-----------------------	--	--------------------

<b>MD-GASTROVIEW</b>			
----------------------	--	--	--

<b>AA LIEBEL-FLARSHEIM</b>	<u><b>66%;10%</b></u>		<b>A087388 001</b>
----------------------------	-----------------------	--	--------------------

DIATRIZOATE MEGLUMINE; IODIPAMIDE MEGLUMINE

SOLUTION; INTRAUTERINE

SINOGRAPHIN			
+! BRACCO	52.7%;26.8%		N011324 002

DIAZEPAM

CONCENTRATE; ORAL

<b>DIAZEPAM</b>			
-----------------	--	--	--

<b>AA LANNETT HOLDINGS</b>	<u><b>5MG/ML</b></u>		<b>A204433 001</b>
----------------------------	----------------------	--	--------------------

<b>INC</b>			
------------	--	--	--

<b>DIAZEPAM INTENSOL</b>			
--------------------------	--	--	--

<b>AA ! WEST-WARD PHARMS</b>	<u><b>5MG/ML</b></u>		<b>A071415 001</b>
------------------------------	----------------------	--	--------------------

<b>INT</b>			
------------	--	--	--

GEL; RECTAL

<b>DIASTAT</b>			
----------------	--	--	--

+! VALEANT PHARMS	2.5MG/0.5ML (5MG/ML)		
-------------------	----------------------	--	--

NORTH			
-------	--	--	--

<b>DIASTAT ACUDIAL</b>			
------------------------	--	--	--

+! VALEANT PHARMS	10MG/2ML (5MG/ML)		
-------------------	-------------------	--	--

NORTH			
-------	--	--	--

+!	20MG/4ML (5MG/ML)		
----	-------------------	--	--

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-130 (of 436)

DIAZEPAM

INJECTABLE; INJECTION

DIAZEPAM

! HOSPIRA	10MG/2ML (5MG/ML)	A072079 001 Dec 20, 1988
!	50MG/10ML (5MG/ML)	A071583 001 Oct 13, 1987

SOLUTION; ORAL

DIAZEPAM

AA LANNETT HOLDINGS

INC

5MG/5ML

A206477 001 Jun 24, 2016

AA ! WEST-WARD PHARMS

INT

5MG/5ML

A070928 001 Apr 03, 1987

TABLET; ORAL

DIAZEPAM

AB BARR

2MG

A070152 001 Nov 01, 1985

AB

10MG

A070154 001 Nov 01, 1985

AB IVAX SUB TEVA

PHARMS

2MG

A071307 001 Dec 10, 1986

AB

5MG

A071321 001 Dec 10, 1986

AB

10MG

A071322 001 Dec 10, 1986

AB MAYNE PHARMA

2MG

A071134 001 Feb 03, 1987

AB

5MG

A071135 001 Feb 03, 1987

AB

10MG

A071136 001 Feb 03, 1987

AB MYLAN

2MG

A070325 002 Sep 04, 1985

AB

5MG

A070325 003 Sep 04, 1985

AB VINTAGE PHARMS

2MG

A077749 001 Mar 31, 2006

AB

5MG

A077749 002 Mar 31, 2006

AB

10MG

A077749 003 Mar 31, 2006

VALIUM

AB + ROCHE

2MG

N013263 002

AB +

5MG

N013263 004

AB +!

10MG

N013263 006

DIAZEPAM

DAVA PHARMS INC

2MG

A070228 002 Sep 26, 1985

5MG

A070228 003 Sep 26, 1985

DIAZOXIDE

SUSPENSION; ORAL

PROGLYCEM

+! TEVA BRANDED PHARM

50MG/ML

N017453 001

DICHLORPHENAMIDE

TABLET; ORAL

KEVEYIS

+! STRONGBRIDGE US

50MG

N011366 002 Aug 07, 2015

DICLOFENAC

CAPSULE; ORAL

ZORVOLEX

+ IROKO PHARMS LLC

18MG

N204592 001 Oct 18, 2013

+!

35MG

N204592 002 Oct 18, 2013

DICLOFENAC EPOLAMINE

PATCH; TOPICAL

FLECTOR

+! INST BIOCHEM

1.3%

N021234 001 Jan 31, 2007

DICLOFENAC POTASSIUM

CAPSULE; ORAL

DICLOFENAC POTASSIUM

AB BIONPHARMA INC

25MG

A204648 001 Feb 23, 2016

ZIPSOR

AB +! DEPOMED INC

25MG

N022202 001 Jun 16, 2009

FOR SOLUTION; ORAL

CAMBIA

AB +! DEPOMED INC

50MG

N022165 001 Jun 17, 2009

DICLOFENAC POTASSIUM

AB PAR FORM

50MG

A202964 001 May 02, 2016

TABLET; ORAL

DICLOFENAC POTASSIUM

AB APOTEX

50MG

A076561 001 Mar 18, 2004

AB ! MYLAN

50MG

A075463 001 Jul 26, 1999

AB SANDOZ

50MG

A075229 001 Nov 20, 1998

AB TEVA

50MG

A075219 001 Aug 06, 1998

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PREScription DRUG PRODUCT LIST**

3-131 (of 436)

## DICLOFENAC SODIUM

GEL; TOPICAL

## DICLOFENAC SODIUM

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>3%</u>	<u>A206493</u> <u>001</u>	Dec 02, 2015
<u>AB</u>	AMNEAL PHARMS	<u>1%</u>	<u>A208077</u> <u>001</u>	Mar 18, 2016
<u>AB</u>	GLENMARK PHARMS LTD	<u>3%</u>	<u>A208301</u> <u>001</u>	Sep 13, 2016
<u>AB</u>	TARO	<u>3%</u>	<u>A206298</u> <u>001</u>	Apr 28, 2016
<b><u>DICLOFENAC SODIUM</u></b>				
<u>AB</u>	TOLMAR	<u>3%</u>	<u>A200936</u> <u>001</u>	Oct 28, 2013
<b><u>SOLARAZE</u></b>				
<u>AB</u>	+! FOUGERA PHARMS	<u>3%</u>	<u>N021005</u> <u>001</u>	Oct 16, 2000
<b><u>VOLTAREN</u></b>				
<u>AB</u>	+! GLAXOSMITHKLINE CONS	<u>1%</u>	<u>N022122</u> <u>001</u>	Oct 17, 2007
SOLUTION; INTRAVENOUS				
DYLOJECT				
+!	JAVELIN PHARMS INC	37.5MG/ML (37.5MG/ML)	N022396 001	Dec 23, 2014

## SOLUTION; TOPICAL

## DICLOFENAC

AMNEAL PHARMACEUTICALS

## AMNEAL PHAR

<u>AT</u>	ANDA REPOSITORY	<u>1.5%</u>	<u>A202393</u>	<u>001</u>	Nov 24, 2014
<u>AT</u>	APOTEX INC	<u>1.5%</u>	<u>A202027</u>	<u>001</u>	May 27, 2014
<u>AT</u>	LUPIN LTD	<u>1.5%</u>	<u>A204132</u>	<u>001</u>	Aug 20, 2015
<u>AT</u>	NOVEL LABS INC	<u>1.5%</u>	<u>A205878</u>	<u>001</u>	Dec 09, 2015
<u>AT</u>	RICONPHARMA LLC	<u>1.5%</u>	<u>A206715</u>	<u>001</u>	Aug 07, 2017
<u>AT</u>	TARO	<u>1.5%</u>	<u>A203818</u>	<u>001</u>	Nov 26, 2014
<u>AT</u>	TELIGENT PHARMA INC	<u>1.5%</u>	<u>A202769</u>	<u>001</u>	Jul 08, 2015
<u>AT</u>	WATSON LABS INC	<u>1.5%</u>	<u>A202852</u>	<u>001</u>	Nov 24, 2014
PENNSAID					
+!	HORIZON PHARMA	2%	N204623 001 Jan 16, 2014		
SOLUTION/DROPS;OPHTHALMIC					

DICLOFEN

**TAKORN**

**AKORN**  
**ALTAIR DYNAMIC IN**

ALTAIRE PHA  
TAMBOURIN

<u>AT</u>	BAUSCH AND LOMB	<u>0.1%</u>	<u>A078792</u>	<u>001</u>	Dec 28, 2007
<u>AT</u>	RISING PHARMS INC	<u>0.1%</u>	<u>A078553</u>	<u>001</u>	Dec 28, 2007
<u>AT</u>	SANDOZ INC	<u>0.1%</u>	<u>A078031</u>	<u>001</u>	Feb 06, 2008

**VOLTAREN**

**AT +! NOVARTIS**  
**TABLET, DELAYED RELEASE; ORAL**  
**DIGOXINIC SODIUM**

## DICLOFENAC SODIUM

<u>AB</u>	ACTAVIS ELIZABETH	<u>50MG</u>	<u>A074514</u>	<u>001</u>	Mar 26, 1996
<u>AB</u>		<u>75MG</u>	<u>A074514</u>	<u>002</u>	Mar 26, 1996
<u>AB</u>	CARLSBAD	<u>25MG</u>	<u>A075185</u>	<u>002</u>	Nov 13, 1998
<u>AB</u>		<u>50MG</u>	<u>A075185</u>	<u>003</u>	Nov 13, 1998
<u>AB</u>		<u>75MG</u>	<u>A075185</u>	<u>001</u>	Nov 13, 1998
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A075281</u>	<u>002</u>	Feb 12, 2002
<u>AB</u>		<u>75MG</u>	<u>A075281</u>	<u>003</u>	Feb 12, 2002
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A074376</u>	<u>001</u>	Sep 28, 1995
<u>AB</u>		<u>50MG</u>	<u>A074376</u>	<u>002</u>	Sep 28, 1995
<u>AB</u>		<u>75MG</u>	<u>A074394</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	UNIQUE PHARM LABS	<u>25MG</u>	<u>A090066</u>	<u>001</u>	Dec 01, 2010
<u>AB</u>		<u>50MG</u>	<u>A090066</u>	<u>002</u>	Dec 01, 2010
<u>AB</u>		<u>75MG</u>	<u>A077863</u>	<u>003</u>	Jun 08, 2007

TABLET, EXTENDED RELEASE; ORAL

## DICLOFENAC SODIUM

<u>AB</u>	!	DEXCEL LTD	<u>100MG</u>	<u>A076201</u>	<u>001</u>	Nov 06, 2002
<u>AB</u>		MYLAN	<u>100MG</u>	<u>A076152</u>	<u>001</u>	Dec 13, 2001
<u>AB</u>		VPNA	<u>100MG</u>	<u>A075492</u>	<u>001</u>	Feb 11, 2000

## DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE; QDAT

ARTHRITIS

**AB** + GD SEARLE LLC **50MG ; 0 .2MG** **N020607 001** Dec 24, 1997  
**AB** + ! **75MG ; 0 .2MG** **N020607 002** Dec 24, 1997

## DICLOFENAC SODIUM AND MISOPROSTOL

<u><a href="#">AB</a></u>	ACTAVIS LABS FL INC	<u>50MG ; 0.2MG</u>	<u><a href="#">A201089</a></u>	<u><a href="#">001</a></u>	Jul 09, 2012
<u><a href="#">AB</a></u>		<u>75MG ; 0.2MG</u>	<u><a href="#">A201089</a></u>	<u><a href="#">002</a></u>	Jul 09, 2012
<u><a href="#">AB</a></u>	AMNEAL PHARMS	<u>50MG ; 0.2MG</u>	<u><a href="#">A203995</a></u>	<u><a href="#">001</a></u>	Nov 25, 2016
<u><a href="#">AB</a></u>		<u>75MG ; 0.2MG</u>	<u><a href="#">A203995</a></u>	<u><a href="#">002</a></u>	Nov 25, 2016
<u><a href="#">AB</a></u>	EXELA HOLDINGS	<u>50MG ; 0.2MG</u>	<u><a href="#">A200540</a></u>	<u><a href="#">001</a></u>	Mar 14, 2014
<u><a href="#">AB</a></u>		<u>75MG ; 0.2MG</u>	<u><a href="#">A200540</a></u>	<u><a href="#">002</a></u>	Mar 14, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-132 (of 436)

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM AND MISOPROSTOL

<u>AB</u>	SANDOZ	<u>50MG;0.2MG</u>
<u>AB</u>		<u>75MG;0.2MG</u>

<u>A200158 001</u>	May 09, 2013
<u>A200158 002</u>	May 09, 2013

DICLOXAECILLIN SODIUM

CAPSULE;ORAL

DICLOXAECILLIN SODIUM

<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>
<u>AB</u>	!	<u>EQ 500MG BASE</u>
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>
<u>AB</u>		<u>EQ 500MG BASE</u>
	SANDOZ	EQ 125MG BASE

<u>A061454 001</u>	
<u>A061454 003</u>	
<u>A062286 001</u>	Jun 03, 1982
<u>A062286 002</u>	Jun 03, 1982
A061454 002	

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

BENTYL

<u>AB</u>	+!	FOREST LABS INC	<u>10MG</u>
-----------	----	-----------------	-------------

N007409 003 Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>	LANNETT	<u>10MG</u>
<u>AB</u>	MYLAN	<u>10MG</u>
<u>AB</u>	WATSON LABS	<u>10MG</u>

<u>A084285 001</u>	
<u>A040319 001</u>	Sep 07, 1999
<u>A085082 001</u>	Jun 19, 1986

INJECTABLE;INJECTION

BENTYL

<u>AP</u>	+!	FOREST LABS INC	<u>10MG/ML</u>
-----------	----	-----------------	----------------

N008370 001 Oct 15, 1984

BENTYL PRESERVATIVE FREE

<u>AP</u>	+!	FOREST LABS INC	<u>10MG/ML</u>
-----------	----	-----------------	----------------

N008370 002 Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

<u>AP</u>	LUITPOLD PHARMS INC	<u>10MG/ML</u>
<u>AP</u>	DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE)	
<u>AP</u>	WEST-WARD PHARMS INT	<u>10MG/ML</u>

<u>A208353 001</u>	Feb 17, 2017
<u>A040465 001</u>	Jun 30, 2003

SYRUP;ORAL

DICYCLOMINE HYDROCHLORIDE

! MIKART

10MG/5ML

A040169 001 Mar 24, 2005

TABLET;ORAL

BENTYL

<u>AB</u>	+!	FOREST LABS INC	<u>20MG</u>
-----------	----	-----------------	-------------

N007409 001 Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>	LANNETT	<u>20MG</u>
<u>AB</u>	MYLAN	<u>20MG</u>
<u>AB</u>	WATSON LABS	<u>20MG</u>

<u>A040230 001</u>	Feb 26, 1999
<u>A040317 001</u>	Sep 07, 1999
<u>A085223 001</u>	Jul 30, 1986

DIDANOSINE

CAPSULE, DELAYED REL PELLETS;ORAL

DIDANOSINE

<u>AB</u>	AUROBINDO PHARMA	<u>125MG</u>
<u>AB</u>		<u>200MG</u>
<u>AB</u>		<u>250MG</u>
<u>AB</u>		<u>400MG</u>
<u>AB</u>	BARR	<u>200MG</u>
<u>AB</u>		<u>250MG</u>
<u>AB</u>		<u>400MG</u>

<u>A090094 001</u>	Sep 24, 2008
<u>A090094 002</u>	Sep 24, 2008
<u>A090094 003</u>	Sep 24, 2008
<u>A090094 004</u>	Sep 24, 2008
<u>A077167 001</u>	Dec 03, 2004
<u>A077167 002</u>	Dec 03, 2004
<u>A077167 003</u>	Dec 03, 2004

VIDEX EC

<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>125MG</u>
<u>AB</u>	+	<u>200MG</u>
<u>AB</u>	+	<u>250MG</u>
<u>AB</u>	++!	<u>400MG</u>

<u>N021183 001</u>	Oct 31, 2000
<u>N021183 002</u>	Oct 31, 2000
<u>N021183 003</u>	Oct 31, 2000
<u>N021183 004</u>	Oct 31, 2000

FOR SOLUTION;ORAL

VIDEX

! BRISTOL-MYERS SQUIBB

10MG/ML

N020156 001 Oct 09, 1991

DIENOGENEST; ESTRADIOL VALERATE

TABLET;ORAL

NATAZIA

! BAYER HLTHCARE

N/A, 2MG, 3MG, N/A, N/A; 3MG, 2MG, 2MG, 1MG, N/A

N022252 001 May 06, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-133 (of 436)

DIETHYLPROMION HYDROCHLORIDE

TABLET;ORAL

DIETHYLPROMION HYDROCHLORIDE

<u>AA</u>	AVANTHI INC	<u>25MG</u>	<u>A201212 001</u> Dec 22, 2010
<u>AA</u>	LANNETT HOLDINGS INC	<u>25MG</u>	<u>A200177 001</u> Jul 18, 2011

TENUATE

<u>AA</u>	+!	ACTAVIS LABS UT INC	<u>25MG</u>	<u>N011722 002</u>
-----------	----	---------------------	-------------	--------------------

TABLET, EXTENDED RELEASE;ORAL

DIETHYLPROMION HYDROCHLORIDE

<u>AB</u>	LANNETT HOLDINGS INC	<u>75MG</u>	<u>A091680 001</u> Oct 24, 2011
-----------	----------------------	-------------	---------------------------------

TENUATE DOSPAN

<u>AB</u>	+!	ACTAVIS LABS UT INC	<u>75MG</u>	<u>N012546 001</u>
-----------	----	---------------------	-------------	--------------------

DIFLORASONE DIACETATE

CREAM;TOPICAL

DIFLORASONE DIACETATE

BX	!	FOUGERA PHARMS	0.05%	A076263 001 Dec 20, 2002
BX	!	TARO	0.05%	A075508 001 Apr 24, 2000

OINTMENT;TOPICAL

DIFLORASONE DIACETATE

<u>AB</u>	AKORN	<u>0.05%</u>	<u>A206572 001</u> Jul 24, 2015
<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A075374 001</u> Apr 27, 1999
<u>AB</u>	RICONPHARMA LLC	<u>0.05%</u>	<u>A207440 001</u> Feb 27, 2017
<u>AB</u>	! TARO	<u>0.05%</u>	<u>A075331 001</u> May 14, 1999

DIFLUNISAL

TABLET;ORAL

DIFLUNISAL

<u>AB</u>	HERITAGE PHARMA	<u>500MG</u>	<u>A202845 001</u> Mar 08, 2012
<u>AB</u>	! TEVA	<u>500MG</u>	<u>A073673 001</u> Jul 31, 1992
<u>AB</u>	ZYDUS PHARMS USA INC	<u>500MG</u>	<u>A203547 001</u> Jun 16, 2017

DIFLUPREDNATE

EMULSION;OPHTHALMIC

DUREZOL

+!	NOVARTIS PHARMS CORP	0.05%	N022212 001 Jun 23, 2008
----	----------------------	-------	--------------------------

DIGOXIN

ELIXIR;ORAL

DIGOXIN

+!	WEST-WARD PHARMS INT	0.05MG/ML	N021648 001 Aug 26, 2004
----	----------------------	-----------	--------------------------

INJECTABLE;INJECTION

DIGOXIN

<u>AP</u>	SANDOZ INC	<u>0.25MG/ML</u>	<u>A040481 001</u> Aug 21, 2003
<u>AP</u>	WEST-WARD PHARMS INT	<u>0.25MG/ML</u>	<u>A083391 001</u>

LANOXIN

<u>AP</u>	+! COVIS PHARMA BV	<u>0.25MG/ML</u>	<u>N009330 002</u>
	LANOXIN PEDIATRIC		

+!	COVIS PHARMA BV	0.1MG/ML	N009330 004
----	-----------------	----------	-------------

TABLET;ORAL

DIGOXIN

<u>AB</u>	HIKMA INTL PHARMS	<u>0.125MG</u>	<u>A077002 002</u> Oct 30, 2007
<u>AB</u>		<u>0.25MG</u>	<u>A077002 001</u> Oct 30, 2007

<u>AB</u>	IMPAK LABS	<u>0.125MG</u>	<u>A078556 001</u> Jul 20, 2009
<u>AB</u>		<u>0.25MG</u>	<u>A078556 002</u> Jul 20, 2009

<u>AB</u>	MYLAN PHARMS INC	<u>0.125MG</u>	<u>A040282 001</u> Dec 23, 1999
<u>AB</u>		<u>0.25MG</u>	<u>A040282 002</u> Dec 23, 1999

<u>AB</u>	STEVENS J	<u>0.125MG</u>	<u>A076268 001</u> Jul 26, 2002
<u>AB</u>		<u>0.25MG</u>	<u>A076268 002</u> Jul 26, 2002

<u>AB</u>	SUN PHARM INDs INC	<u>0.125MG</u>	<u>A076363 001</u> Jan 31, 2003
<u>AB</u>		<u>0.25MG</u>	<u>A076363 002</u> Jan 31, 2003

<u>AB</u>		<u>0.125MG</u>	<u>N020405 002</u> Sep 30, 1997
<u>AB</u>		<u>0.25MG</u>	<u>N020405 004</u> Sep 30, 1997

+!	CONCORDIA PHARMS INC	0.0625MG	N020405 001 Sep 30, 1997
+		0.1875MG	N020405 003 Sep 30, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-134 (of 436)

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

<u>AP</u>	+!	VALEANT	<u>1MG/ML</u>	<u>N005929 001</u>
		<u>DIHYDROERGOTAMINE MESYLATE</u>		
<u>AP</u>		HIKMA PHARMS	<u>1MG/ML</u>	<u>A206621 001</u> Sep 15, 2017
<u>AP</u>		PADDOCK LLC	<u>1MG/ML</u>	<u>A040475 001</u> Apr 28, 2003
<u>AP</u>		WEST-WARD PHARMS	<u>1MG/ML</u>	<u>A040453 001</u> Jun 09, 2003
		INT		
		SPRAY, METERED;NASAL		
		MIGRANAL		
	+!	VALEANT	0.5MG/INH	N020148 001 Dec 08, 1997

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB2</u>		APOTEX	<u>120MG</u>	<u>A074943 003</u> Dec 19, 2000
<u>AB2</u>			<u>180MG</u>	<u>A074943 002</u> Dec 19, 2000
<u>AB2</u>			<u>240MG</u>	<u>A074943 001</u> Aug 06, 1998
<u>AB2</u>		MYLAN	<u>120MG</u>	<u>A075124 002</u> Mar 18, 1998
<u>AB2</u>			<u>180MG</u>	<u>A075124 003</u> Mar 18, 1998
<u>AB2</u>	!		<u>240MG</u>	<u>A075124 001</u> Mar 18, 1998
		<u>CARDIZEM CD</u>		
<u>AB3</u>	+	VALEANT INTL	<u>120MG</u>	<u>N020062 001</u> Aug 10, 1992
<u>AB3</u>	+		<u>180MG</u>	<u>N020062 002</u> Dec 27, 1991
<u>AB3</u>	+		<u>240MG</u>	<u>N020062 003</u> Dec 27, 1991
<u>AB3</u>	+		<u>300MG</u>	<u>N020062 004</u> Dec 27, 1991
<u>AB3</u>	+		<u>360MG</u>	<u>N020062 005</u> Aug 24, 1999
		<u>CARTIA XT</u>		
<u>AB3</u>		ACTAVIS LABS FL INC	<u>120MG</u>	<u>A074752 002</u> Jul 09, 1998
<u>AB3</u>			<u>180MG</u>	<u>A074752 001</u> Jul 09, 1998
<u>AB3</u>			<u>240MG</u>	<u>A074752 003</u> Jul 09, 1998
<u>AB3</u>			<u>300MG</u>	<u>A074752 004</u> Jul 09, 1998
		<u>DILTIAZEM HYDROCHLORIDE</u>		
<u>AB3</u>		ACTAVIS ELIZABETH	<u>360MG</u>	<u>A202463 001</u> Dec 07, 2012
<u>AB3</u>		PAR PHARM	<u>120MG</u>	<u>A074984 001</u> Dec 20, 1999
<u>AB3</u>			<u>180MG</u>	<u>A074984 002</u> Dec 20, 1999
<u>AB3</u>			<u>240MG</u>	<u>A074984 003</u> Dec 20, 1999
<u>AB3</u>			<u>300MG</u>	<u>A074984 004</u> Dec 20, 1999
<u>AB3</u>		SUN PHARM IND LTD	<u>120MG</u>	<u>A203023 001</u> Jun 08, 2017
<u>AB3</u>			<u>180MG</u>	<u>A203023 002</u> Jun 08, 2017
<u>AB3</u>			<u>240MG</u>	<u>A203023 003</u> Jun 08, 2017
<u>AB3</u>			<u>300MG</u>	<u>A203023 004</u> Jun 08, 2017
<u>AB3</u>			<u>360MG</u>	<u>A203023 005</u> Jun 08, 2017
<u>AB3</u>		SUN PHARMA GLOBAL	<u>120MG</u>	<u>A090492 001</u> Oct 28, 2011
<u>AB3</u>			<u>180MG</u>	<u>A090492 002</u> Oct 28, 2011
<u>AB3</u>			<u>240MG</u>	<u>A090492 003</u> Oct 28, 2011
<u>AB3</u>			<u>300MG</u>	<u>A090492 004</u> Oct 28, 2011
<u>AB3</u>			<u>360MG</u>	<u>A090492 005</u> Oct 28, 2011
<u>AB3</u>		VALEANT PHARMS NORTH	<u>120MG</u>	<u>A075116 001</u> Dec 23, 1999
<u>AB3</u>			<u>180MG</u>	<u>A075116 002</u> Dec 23, 1999
<u>AB3</u>			<u>240MG</u>	<u>A075116 003</u> Dec 23, 1999
<u>AB3</u>			<u>300MG</u>	<u>A075116 004</u> Dec 23, 1999
<u>AB3</u>		ZYDUS PHARMS USA INC	<u>120MG</u>	<u>A206534 001</u> Aug 08, 2017
<u>AB3</u>			<u>180MG</u>	<u>A206534 002</u> Aug 08, 2017
<u>AB3</u>			<u>240MG</u>	<u>A206534 003</u> Aug 08, 2017
<u>AB3</u>			<u>300MG</u>	<u>A206534 004</u> Aug 08, 2017
<u>AB3</u>			<u>360MG</u>	<u>A206534 005</u> Aug 08, 2017
<u>AB4</u>		SANDOZ	<u>120MG</u>	<u>A091022 001</u> Sep 28, 2012
<u>AB4</u>			<u>180MG</u>	<u>A091022 002</u> Sep 28, 2012
<u>AB4</u>			<u>240MG</u>	<u>A091022 003</u> Sep 28, 2012
<u>AB4</u>			<u>300MG</u>	<u>A091022 004</u> Sep 28, 2012
<u>AB4</u>			<u>360MG</u>	<u>A091022 005</u> Sep 28, 2012
<u>AB4</u>			<u>420MG</u>	<u>A091022 006</u> Sep 28, 2012
<u>AB4</u>		SUN PHARMA GLOBAL	<u>120MG</u>	<u>A090421 001</u> Nov 15, 2010
<u>AB4</u>			<u>180MG</u>	<u>A090421 002</u> Nov 15, 2010
<u>AB4</u>			<u>240MG</u>	<u>A090421 003</u> Nov 15, 2010
<u>AB4</u>			<u>300MG</u>	<u>A090421 004</u> Nov 15, 2010
<u>AB4</u>			<u>360MG</u>	<u>A090421 005</u> Nov 15, 2010
<u>AB4</u>		ZYDUS PHARMS USA INC	<u>120MG</u>	<u>A206641 001</u> Aug 11, 2017
<u>AB4</u>			<u>180MG</u>	<u>A206641 002</u> Aug 11, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-135 (of 436)

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB4</u>		<u>240MG</u>	<u>A206641</u> <u>003</u>	Aug 11, 2017
<u>AB4</u>		<u>300MG</u>	<u>A206641</u> <u>004</u>	Aug 11, 2017
<u>AB4</u>		<u>360MG</u>	<u>A206641</u> <u>005</u>	Aug 11, 2017
<u>AB4</u>		<u>420MG</u>	<u>A206641</u> <u>006</u>	Aug 11, 2017

DILTZAC

<u>AB4</u>	APOTEX INC	<u>120MG</u>	<u>A076395</u> <u>001</u>	Feb 01, 2006
<u>AB4</u>		<u>180MG</u>	<u>A076395</u> <u>002</u>	Feb 01, 2006
<u>AB4</u>		<u>240MG</u>	<u>A076395</u> <u>003</u>	Feb 01, 2006
<u>AB4</u>		<u>300MG</u>	<u>A076395</u> <u>004</u>	Feb 01, 2006
<u>AB4</u>		<u>360MG</u>	<u>A076395</u> <u>005</u>	Feb 01, 2006

TAZTIA XT

<u>AB4</u>	ACTAVIS LABS FL INC	<u>120MG</u>	<u>A075401</u> <u>001</u>	Apr 10, 2003
<u>AB4</u>		<u>180MG</u>	<u>A075401</u> <u>002</u>	Apr 10, 2003
<u>AB4</u>		<u>240MG</u>	<u>A075401</u> <u>003</u>	Apr 10, 2003
<u>AB4</u>		<u>300MG</u>	<u>A075401</u> <u>004</u>	Apr 10, 2003
<u>AB4</u>		<u>360MG</u>	<u>A075401</u> <u>005</u>	Apr 10, 2003

TIAZAC

<u>AB4</u> +	VALEANT PHARMS NORTH	<u>120MG</u>	<u>N020401</u> <u>001</u>	Sep 11, 1995
<u>AB4</u> +		<u>180MG</u>	<u>N020401</u> <u>002</u>	Sep 11, 1995
<u>AB4</u> +		<u>240MG</u>	<u>N020401</u> <u>003</u>	Sep 11, 1995
<u>AB4</u> +		<u>300MG</u>	<u>N020401</u> <u>004</u>	Sep 11, 1995
<u>AB4</u> +		<u>360MG</u>	<u>N020401</u> <u>005</u>	Sep 11, 1995
<u>AB4</u> +!		<u>420MG</u>	<u>N020401</u> <u>006</u>	Oct 16, 1998

DILTIAZEM HYDROCHLORIDE

BC !	MYLAN	120MG	A074910	003 May 02, 1997
		60MG	A074910	001 May 02, 1997
		90MG	A074910	002 May 02, 1997

INJECTABLE; INJECTION

DILTIAZEM HYDROCHLORIDE

<u>AP</u>	AKORN INC	<u>5MG/ML</u>	<u>A075086</u> <u>001</u>	Apr 09, 1998
<u>AP</u> !	ATHENEX INC	<u>5MG/ML</u>	<u>A074617</u> <u>001</u>	Feb 28, 1996
<u>AP</u>	HIKMA FARMACEUTICA	<u>5MG/ML</u>	<u>A202651</u> <u>001</u>	Aug 09, 2012
<u>AP</u>	HOSPIRA	<u>5MG/ML</u>	<u>A074941</u> <u>001</u>	Apr 15, 1998
<u>AP</u>	INTL MEDICATION	<u>5MG/ML</u>	<u>A075749</u> <u>001</u>	Nov 21, 2001
<u>AP</u>	WEST-WARD PHARMS INT	<u>5MG/ML</u>	<u>A078538</u> <u>001</u>	Dec 17, 2008
!	HOSPIRA	100MG/VIAL	A075853	001 Dec 17, 2002

TABLET; ORAL

CARDIZEM

<u>AB</u> +	VALEANT INTL	<u>30MG</u>	<u>N018602</u> <u>001</u>	Nov 05, 1982
<u>AB</u> +		<u>60MG</u>	<u>N018602</u> <u>002</u>	Nov 05, 1982
<u>AB</u> +		<u>90MG</u>	<u>N018602</u> <u>003</u>	Dec 08, 1986
<u>AB</u> +!		<u>120MG</u>	<u>N018602</u> <u>004</u>	Dec 08, 1986

DILTIAZEM HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>30MG</u>	<u>A072838</u> <u>004</u>	Nov 05, 1992
<u>AB</u>		<u>60MG</u>	<u>A072838</u> <u>003</u>	Nov 05, 1992
<u>AB</u>		<u>90MG</u>	<u>A072838</u> <u>002</u>	Nov 05, 1992
<u>AB</u>		<u>120MG</u>	<u>A072838</u> <u>001</u>	Nov 05, 1992
<u>AB</u>	TEVA	<u>30MG</u>	<u>A074185</u> <u>001</u>	May 31, 1995
<u>AB</u>		<u>60MG</u>	<u>A074185</u> <u>002</u>	May 31, 1995
<u>AB</u>		<u>90MG</u>	<u>A074185</u> <u>003</u>	May 31, 1995
<u>AB</u>		<u>120MG</u>	<u>A074185</u> <u>004</u>	May 31, 1995

TABLET, EXTENDED RELEASE; ORAL

CARDIZEM LA

<u>AB</u> +	VALEANT INTL	<u>120MG</u>	<u>N021392</u> <u>001</u>	Feb 06, 2003
<u>AB</u> +		<u>180MG</u>	<u>N021392</u> <u>002</u>	Feb 06, 2003
<u>AB</u> +		<u>240MG</u>	<u>N021392</u> <u>003</u>	Feb 06, 2003
<u>AB</u> +		<u>300MG</u>	<u>N021392</u> <u>004</u>	Feb 06, 2003
<u>AB</u> +		<u>360MG</u>	<u>N021392</u> <u>005</u>	Feb 06, 2003
<u>AB</u> +!		<u>420MG</u>	<u>N021392</u> <u>006</u>	Feb 06, 2003

DILTIAZEM HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>120MG</u>	<u>A077686</u> <u>006</u>	Mar 15, 2010
<u>AB</u>		<u>180MG</u>	<u>A077686</u> <u>005</u>	Mar 15, 2010
<u>AB</u>		<u>240MG</u>	<u>A077686</u> <u>004</u>	Mar 15, 2010
<u>AB</u>		<u>300MG</u>	<u>A077686</u> <u>003</u>	Mar 15, 2010
<u>AB</u>		<u>360MG</u>	<u>A077686</u> <u>002</u>	Mar 15, 2010
<u>AB</u>		<u>420MG</u>	<u>A077686</u> <u>001</u>	Mar 15, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-136 (of 436)

DIMENHYDRINATE

INJECTABLE; INJECTION  
 DIMENHYDRINATE  
 ! FRESENIUS KABI USA 50MG/ML A040519 001 Jun 23, 2004

DIMERCAPOL

INJECTABLE; INJECTION  
 BAL  
 +! AKORN 10% N005939 001

DIMETHYL FUMARATE

CAPSULE, DELAYED RELEASE; ORAL  
 TECFIDERA  
 +! BIOGEN IDEC INC 120MG N204063 001 Mar 27, 2013  
 +! 240MG N204063 002 Mar 27, 2013

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL  
DIMETHYL SULFOXIDE  
AT MYLAN INSTITUTIONAL 50% A076185 001 Nov 29, 2002  
RIMSO-50  
AT +! MYLAN INSTITUTIONAL 50% N017788 001

DINOPROSTONE

GEL; ENDOCERVICAL  
 PREPIDIL  
 +! PHARMACIA AND 0.5MG/3GM N019617 001 Dec 09, 1992  
 UPJOHN  
 INSERT, EXTENDED RELEASE; VAGINAL  
 CERVIDIL  
 +! FERRING PHARMS INC 10MG N020411 001 Mar 30, 1995  
 SUPPOSITORY; VAGINAL  
 PROSTIN E2  
 +! PHARMACIA AND 20MG N017810 001  
 UPJOHN

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL  
 DIPHENHYDRAMINE HYDROCHLORIDE  
 ! BARR 50MG A080738 001  
 ELIXIR; ORAL  
 DIPHENHYDRAMINE HYDROCHLORIDE  
 ! PHARM ASSOC 12.5MG/5ML A087513 001 Feb 10, 1982  
 INJECTABLE; INJECTION  
DIPHENHYDRAMINE HYDROCHLORIDE  
AP APP PHARMS 50MG/ML A040466 001 May 28, 2002  
AP HOSPIRA 50MG/ML A040140 001 Nov 20, 1998  
AP MYLAN INSTITUTIONAL 50MG/ML A040498 001 Jul 12, 2005  
AP ! WEST-WARD PHARMS 50MG/ML A080817 002  
 INT

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

AP FRESENIUS KABI USA 50MG/ML A091526 001 Mar 26, 2013

DIPYRIDAMOLE

INJECTABLE; INJECTION  
DIPYRIDAMOLE  
AP ! ATHENEX INC 5MG/ML A074939 001 Apr 13, 1998  
AP FRESENIUS KABI USA 5MG/ML A074956 001 Sep 30, 1998  
AP WEST-WARD PHARMS 5MG/ML A074521 001 Oct 18, 1996  
 INT

TABLET; ORAL

DIPYRIDAMOLE  
AB BARR 25MG A087184 001 Oct 03, 1990  
AB 50MG A087716 001 Oct 03, 1990  
AB 75MG A087717 001 Oct 03, 1990  
AB IMPAX LABS 25MG A040782 001 Jul 18, 2007  
AB 50MG A040782 002 Jul 18, 2007  
AB 75MG A040782 003 Jul 18, 2007  
AB MURTY PHARMS 25MG A040733 001 Feb 13, 2007  
AB 50MG A040733 002 Feb 13, 2007  
AB 75MG A040733 003 Feb 13, 2007  
AB ZYDUS PHARMS USA 25MG A040874 001 Jan 28, 2008  
 INC  
AB 50MG A040874 002 Jan 28, 2008  
AB 75MG A040874 003 Jan 28, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-137 (of 436)

DIPYRIDAMOLE

TABLET;ORAL

PERSANTINE

<u>AB</u>	+	BOEHRINGER INGELHEIM	<u>25MG</u>	<u>N012836 003</u>	Dec 22, 1986
<u>AB</u>	+		<u>50MG</u>	<u>N012836 004</u>	Feb 06, 1987
<u>AB</u>	+!		<u>75MG</u>	<u>N012836 005</u>	Feb 06, 1987

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

<u>AB</u>	MAYNE PHARMA	<u>EQ 100MG BASE</u>	<u>A070173 001</u>	May 31, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A070173 002</u>	May 31, 1985
<u>AB</u>	TEVA	<u>EQ 100MG BASE</u>	<u>A070101 001</u>	Feb 22, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A070102 001</u>	Feb 22, 1985

NORPACE

<u>AB</u>	+	GD SEARLE LLC	<u>EQ 100MG BASE</u>	<u>N017447 001</u>
<u>AB</u>	+!		<u>EQ 150MG BASE</u>	<u>N017447 002</u>

CAPSULE, EXTENDED RELEASE;ORAL

NORPACE CR

+	GD SEARLE LLC	<u>EQ 100MG BASE</u>
+!		<u>EQ 150MG BASE</u>

N018655 001 Jul 20, 1982  
N018655 002 Jul 20, 1982

DISULFIRAM

TABLET;ORAL

ANTABUSE

<u>AB</u>	ODYSSEY PHARMS	<u>250MG</u>	<u>A088482 001</u>	Dec 08, 1983
<u>AB</u>	!	<u>500MG</u>	<u>A088483 001</u>	Dec 08, 1983

DISULFIRAM

<u>AB</u>	CHARTWELL MOLECULES	<u>250MG</u>	<u>A091563 001</u>	Dec 31, 2012
<u>AB</u>		<u>500MG</u>	<u>A091563 002</u>	Dec 31, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>250MG</u>	<u>A203916 001</u>	Mar 04, 2015
<u>AB</u>		<u>500MG</u>	<u>A203916 002</u>	Mar 04, 2015
<u>AB</u>	SIGMAPHARM LABS LLC	<u>250MG</u>	<u>A091619 001</u>	Mar 28, 2011
<u>AB</u>		<u>500MG</u>	<u>A091619 002</u>	Mar 28, 2011
<u>AB</u>	WEST-WARD PHARMS INT	<u>250MG</u>	<u>A202652 001</u>	Feb 05, 2014
<u>AB</u>		<u>500MG</u>	<u>A202652 002</u>	Feb 05, 2014
			<u>A091681 001</u>	Aug 08, 2013

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS;ORAL

DEPAKOTE

<u>AB</u>	++!	ABBVIE	<u>EQ 125MG VALPROIC ACID</u>	<u>N019680 001</u>	Sep 12, 1989
			<u>DIVALPROEX SODIUM</u>		
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078979 001</u>	Jan 23, 2009
<u>AB</u>		MYLAN	<u>EQ 125MG VALPROIC ACID</u>	<u>A090407 001</u>	Mar 28, 2011
<u>AB</u>		ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A078919 001</u>	Jan 27, 2009

TABLET, DELAYED RELEASE;ORAL

DEPAKOTE

<u>AB</u>	+	ABBVIE	<u>EQ 125MG VALPROIC ACID</u>	<u>N018723 003</u>	Oct 26, 1984
<u>AB</u>	+		<u>EQ 250MG VALPROIC ACID</u>	<u>N018723 001</u>	Mar 10, 1983
<u>AB</u>	+!		<u>EQ 500MG VALPROIC ACID</u>	<u>N018723 002</u>	Mar 10, 1983
			<u>DIVALPROEX SODIUM</u>		
<u>AB</u>		ACTAVIS LABS FL INC	<u>EQ 500MG VALPROIC ACID</u>	<u>A079080 001</u>	Feb 25, 2011
<u>AB</u>		ANCHEN PHARMS	<u>EQ 500MG VALPROIC ACID</u>	<u>A078411 001</u>	Nov 03, 2008
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A090554 001</u>	Apr 21, 2011
<u>AB</u>			<u>EQ 250MG VALPROIC ACID</u>	<u>A090554 002</u>	Apr 21, 2011
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A090554 003</u>	Apr 21, 2011
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A078755 001</u>	Jul 29, 2008
<u>AB</u>			<u>EQ 250MG VALPROIC ACID</u>	<u>A078755 002</u>	Jul 29, 2008
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A078755 003</u>	Jul 29, 2008
<u>AB</u>		LUPIN	<u>EQ 125MG VALPROIC ACID</u>	<u>A078790 001</u>	Jul 29, 2008
<u>AB</u>			<u>EQ 250MG VALPROIC ACID</u>	<u>A078790 002</u>	Jul 29, 2008
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A078790 003</u>	Jul 29, 2008
<u>AB</u>		MYLAN	<u>EQ 125MG VALPROIC ACID</u>	<u>A090062 001</u>	Mar 17, 2009
<u>AB</u>			<u>EQ 250MG VALPROIC ACID</u>	<u>A090062 002</u>	Mar 17, 2009
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A090062 003</u>	Mar 17, 2009
<u>AB</u>		NU PHARM	<u>EQ 125MG VALPROIC ACID</u>	<u>A077615 003</u>	Jul 29, 2008
<u>AB</u>			<u>EQ 250MG VALPROIC ACID</u>	<u>A077615 002</u>	Jul 29, 2008
<u>AB</u>			<u>EQ 500MG VALPROIC ACID</u>	<u>A077615 001</u>	Jul 29, 2008
<u>AB</u>		ORCHID HLTHCARE	<u>EQ 125MG VALPROIC ACID</u>	<u>A078853 001</u>	Nov 25, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-138 (of 436)

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DIVALPROEX SODIUM

<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078853 002</u>	Nov 25, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078853 003</u>	Nov 25, 2008
<u>AB</u>	PRINSTON INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A090210 001</u>	Nov 30, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A090210 002</u>	Nov 30, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A090210 003</u>	Nov 30, 2009
<u>AB</u>	SANDOZ	<u>EQ 125MG VALPROIC ACID</u>	<u>A078290 003</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078290 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078290 001</u>	Jul 29, 2008
<u>AB</u>	SUN PHARM INDs	<u>EQ 125MG VALPROIC ACID</u>	<u>A078597 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078597 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078597 003</u>	Jul 29, 2008
<u>AB</u>	TEVA	<u>EQ 125MG VALPROIC ACID</u>	<u>A076941 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A076941 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A076941 003</u>	Jul 29, 2008
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 125MG VALPROIC ACID</u>	<u>A079163 001</u>	Apr 05, 2011
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A079163 002</u>	Apr 05, 2011
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A079163 003</u>	Apr 05, 2011
<u>AB</u>	UPSHER-SMITH LABS	<u>EQ 125MG VALPROIC ACID</u>	<u>A078182 001</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A078182 002</u>	Jul 29, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078182 003</u>	Jul 29, 2008
<u>AB</u>	WOCKHARDT	<u>EQ 125MG VALPROIC ACID</u>	<u>A077296 001</u>	Jul 31, 2008
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077296 002</u>	Jul 31, 2008
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077296 003</u>	Jul 31, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 125MG VALPROIC ACID</u>	<u>A077100 001</u>	Mar 05, 2009
<u>AB</u>		<u>EQ 250MG VALPROIC ACID</u>	<u>A077100 002</u>	Mar 05, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077100 003</u>	Mar 05, 2009

TABLET, EXTENDED RELEASE;ORAL

DEPAKOTE ER

<u>AB</u>	+	ABBVIE	<u>EQ 250MG VALPROIC ACID</u>	<u>N021168 002</u>	May 31, 2002
<u>AB</u>	+!		<u>EQ 500MG VALPROIC ACID</u>	<u>N021168 001</u>	Aug 04, 2000

DIVALPROEX SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>EQ 250MG VALPROIC ACID</u>	<u>A203730 001</u>	May 29, 2015
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A203730 002</u>	May 29, 2015
<u>AB</u>	ANCHEN PHARMS	<u>EQ 250MG VALPROIC ACID</u>	<u>A078445 001</u>	Feb 26, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078445 002</u>	Aug 04, 2009
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A202419 001</u>	Jun 02, 2014
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A202419 002</u>	Jun 02, 2014
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 250MG VALPROIC ACID</u>	<u>A090161 001</u>	Mar 15, 2012
<u>AB</u>	IMPAK LABS	<u>EQ 250MG VALPROIC ACID</u>	<u>A078791 001</u>	May 06, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078791 002</u>	Aug 04, 2009
<u>AB</u>	MYLAN	<u>EQ 250MG VALPROIC ACID</u>	<u>A077567 001</u>	Jan 29, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A077567 002</u>	Jan 29, 2009
<u>AB</u>	REDDYS	<u>EQ 500MG VALPROIC ACID</u>	<u>A090070 001</u>	Mar 12, 2012
<u>AB</u>	WOCKHARDT	<u>EQ 250MG VALPROIC ACID</u>	<u>A078705 002</u>	Feb 10, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078705 001</u>	Aug 04, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 250MG VALPROIC ACID</u>	<u>A078239 001</u>	Feb 27, 2009
<u>AB</u>		<u>EQ 500MG VALPROIC ACID</u>	<u>A078239 002</u>	Aug 04, 2009

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>A074086 001</u>	Nov 29, 1993
<u>AP</u>	!	<u>EQ 12.5MG BASE/ML</u>	<u>A074292 001</u>	Feb 16, 1995
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 12.5MG BASE/ML</u>	<u>A074277 001</u>	Oct 31, 1994

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	++! BAXTER HLTHCARE	<u>EQ 50MG BASE/100ML</u>	<u>N020255 001</u>	Oct 19, 1993
<u>AP</u>	++!	<u>EQ 100MG BASE/100ML</u>	<u>N020255 003</u>	Oct 19, 1993
<u>AP</u>	++!	<u>EQ 200MG BASE/100ML</u>	<u>N020255 004</u>	Oct 19, 1993
<u>AP</u>	++!	<u>EQ 400MG BASE/100ML</u>	<u>N020255 005</u>	Oct 19, 1993
<u>AP</u>	++! HOSPIRA	<u>EQ 50MG BASE/100ML</u>	<u>N020201 003</u>	Oct 19, 1993
<u>AP</u>	++!	<u>EQ 100MG BASE/100ML</u>	<u>N020201 002</u>	Oct 19, 1993
<u>AP</u>	++!	<u>EQ 200MG BASE/100ML</u>	<u>N020201 001</u>	Oct 19, 1993
<u>AP</u>	++!	<u>EQ 400MG BASE/100ML</u>	<u>N020201 006</u>	Jul 07, 1994

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-139 (of 436)

DOCETAXEL

INJECTABLE; INJECTION

DOCETAXEL

<u>AP</u>	+	ACCORD HLTHCARE	<u>20MG/ML (20MG/ML)</u>	<u>N201195 003</u>	Apr 20, 2012
<u>AP</u>	+		<u>80MG/4ML (20MG/ML)</u>	<u>N201195 004</u>	Apr 20, 2012
<u>AP</u>	+!		<u>160MG/8ML (20MG/ML)</u>	<u>N201195 005</u>	Apr 20, 2012
<u>AP</u>		ACTAVIS LLC	<u>20MG/ML (20MG/ML)</u>	<u>N203551 001</u>	Apr 12, 2013
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>N203551 002</u>	Apr 12, 2013
<u>AP</u>		DFB ONCOLOGY LTD	<u>20MG/ML (20MG/ML)</u>	<u>A206177 001</u>	Jan 20, 2017
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A206177 002</u>	Jan 20, 2017
<u>AP</u>		DR REDDYS LABS LTD	<u>20MG/ML (20MG/ML)</u>	<u>A204193 001</u>	Nov 05, 2014
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A204193 002</u>	Nov 05, 2014
<u>AP</u>	+!	HOSPIRA INC	<u>20MG/2ML (10MG/ML)</u>	<u>N022234 001</u>	Mar 08, 2011
<u>AP</u>	+!		<u>80MG/8ML (10MG/ML)</u>	<u>N022234 002</u>	Mar 08, 2011
<u>AP</u>	+!		<u>160MG/16ML (10MG/ML)</u>	<u>N022234 003</u>	Mar 08, 2011
<u>AP</u>		INGENUS PHARMS LLC	<u>20MG/2ML (10MG/ML)</u>	<u>A207563 001</u>	Aug 31, 2017
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>A207563 002</u>	Aug 31, 2017
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>A207563 003</u>	Aug 31, 2017
<u>AP</u>		JIANGSU HENGRIU MED	<u>20MG/ML (20MG/ML)</u>	<u>A207252 001</u>	Aug 09, 2017
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A207252 002</u>	Aug 09, 2017
<u>AP</u>			<u>160MG/8ML (20MG/ML)</u>	<u>A207252 003</u>	Aug 09, 2017
<u>AP</u>		SANDOZ	<u>20MG/2ML (10MG/ML)</u>	<u>N201525 001</u>	Jun 29, 2011
<u>AP</u>			<u>80MG/8ML (10MG/ML)</u>	<u>N201525 002</u>	Jun 29, 2011
<u>AP</u>			<u>160MG/16ML (10MG/ML)</u>	<u>N201525 003</u>	Jun 29, 2011
<u>AP</u>		TEVA PHARMS USA	<u>20MG/ML (20MG/ML)</u>	<u>A203877 001</u>	Sep 16, 2015
<u>AP</u>			<u>80MG/4ML (20MG/ML)</u>	<u>A203877 002</u>	Sep 16, 2015
<b>TAXOTERE</b>					
<u>AP</u>	+!	SANOFI AVENTIS US	<u>20MG/ML (20MG/ML)</u>	<u>N020449 003</u>	Aug 03, 2010
<u>AP</u>	+!		<u>80MG/4ML (20MG/ML)</u>	<u>N020449 004</u>	Aug 02, 2010
<u>AP</u>	+!		<u>160MG/8ML (20MG/ML)</u>	<u>N020449 005</u>	Apr 13, 2012
<b>DOCETAXEL</b>					
	+	ACCORD HLTHCARE	20MG/0.5ML (40MG/ML)	N201195 001	Jun 08, 2011
	+		80MG/2ML (40MG/ML)	N201195 002	Jun 08, 2011
		ACTAVIS LLC	140MG/7ML (20MG/ML)	N203551 003	Apr 12, 2013
		DFB ONCOLOGY LTD	200MG/10ML (20MG/ML)	A206177 003	Jan 20, 2017
	+	HOSPIRA INC	20MG/ML (20MG/ML)	N022234 004	Jun 23, 2016
	+		80MG/4ML (20MG/ML)	N022234 005	Jun 23, 2016
	+		160MG/8ML (20MG/ML)	N022234 007	Jan 24, 2017
	!	JIANGSU HENGRIU MED	40MG/ML	A203170 001	Feb 15, 2017
SOLUTION; IV (INFUSION)					
DOCETAXEL					
		EAGLE PHARMS	20MG/ML (20MG/ML)	N205934 001	Dec 22, 2015
			80MG/4ML (20MG/ML)	N205934 002	Dec 22, 2015
			160MG/8ML (20MG/ML)	N205934 003	Dec 22, 2015

DOFETILIDE

CAPSULE; ORAL

DOFETILIDE

<u>AB</u>		MAYNE PHARMA INC	<u>0.125MG</u>	<u>A207058 001</u>	Jun 06, 2016
<u>AB</u>			<u>0.25MG</u>	<u>A207058 002</u>	Jun 06, 2016
<u>AB</u>			<u>0.5MG</u>	<u>A207058 003</u>	Jun 06, 2016
<b>TIKOSYN</b>					
<u>AB</u>	+	PFIZER	<u>0.125MG</u>	<u>N020931 001</u>	Oct 01, 1999
<u>AB</u>	+		<u>0.25MG</u>	<u>N020931 002</u>	Oct 01, 1999
<u>AB</u>	+!		<u>0.5MG</u>	<u>N020931 003</u>	Oct 01, 1999

DOLUTEGRAVIR SODIUM

TABLET; ORAL

TIVICAY

+	VIIV HLTHCARE	EQ 10MG BASE	N204790 002	Jun 09, 2016
+		EQ 25MG BASE	N204790 003	Jun 09, 2016
+		EQ 50MG BASE	N204790 001	Aug 12, 2013

DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE

TABLET; ORAL

JULUCA

+!	VIIIV HLTHCARE	EQ 50MG BASE; EQ 25MG BASE	N210192 001	Nov 21, 2017
----	----------------	----------------------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-140 (of 436)

DONEPEZIL HYDROCHLORIDE

TABLET;ORAL

ARICEPT

<u>AB</u>	+	EISAI INC	<u>5MG</u>	<u>N020690</u> <u>002</u>	Nov 25, 1996
<u>AB</u>	+!		<u>10MG</u>	<u>N020690</u> <u>001</u>	Nov 25, 1996
<u>AB</u>	+!		<u>23MG</u>	<u>N022568</u> <u>001</u>	Jul 23, 2010
<b><u>DONEPEZIL HYDROCHLORIDE</u></b>					
<u>AB</u>		ACI HEALTHCARE LTD	<u>5MG</u>	<u>A078662</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A078662</u> <u>002</u>	May 31, 2011
<u>AB</u>		ACTAVIS ELIZABETH	<u>23MG</u>	<u>A202415</u> <u>001</u>	Dec 17, 2015
<u>AB</u>		ALEMBIC PHARMS LTD	<u>5MG</u>	<u>A201724</u> <u>001</u>	Feb 25, 2013
<u>AB</u>			<u>10MG</u>	<u>A201724</u> <u>002</u>	Feb 25, 2013
<u>AB</u>		APOTEX	<u>5MG</u>	<u>A078841</u> <u>001</u>	Jun 02, 2011
<u>AB</u>			<u>10MG</u>	<u>A078841</u> <u>002</u>	Jun 02, 2011
<u>AB</u>		AUROBINDO	<u>5MG</u>	<u>A090056</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090056</u> <u>002</u>	May 31, 2011
<u>AB</u>		CADILA PHARMS LTD	<u>5MG</u>	<u>A204609</u> <u>001</u>	Sep 19, 2017
<u>AB</u>			<u>10MG</u>	<u>A204609</u> <u>002</u>	Sep 19, 2017
<u>AB</u>		CIPLA LTD	<u>5MG</u>	<u>A077518</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A077518</u> <u>002</u>	May 31, 2011
<u>AB</u>		CSPC OUYI PHARM CO	<u>5MG</u>	<u>A202114</u> <u>001</u>	Jul 05, 2013
<u>AB</u>			<u>10MG</u>	<u>A202114</u> <u>002</u>	Jul 05, 2013
<u>AB</u>		DEXCEL PHARMA	<u>23MG</u>	<u>A203713</u> <u>001</u>	Feb 19, 2016
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG</u>	<u>A201001</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A201001</u> <u>002</u>	May 31, 2011
<u>AB</u>			<u>23MG</u>	<u>A202723</u> <u>001</u>	Jul 24, 2013
<u>AB</u>		HETERO LABS LTD V	<u>5MG</u>	<u>A203034</u> <u>001</u>	Jan 30, 2015
<u>AB</u>			<u>10MG</u>	<u>A203034</u> <u>002</u>	Jan 30, 2015
<u>AB</u>		INDICUS PHARMA	<u>5MG</u>	<u>A201634</u> <u>001</u>	Jun 13, 2012
<u>AB</u>			<u>10MG</u>	<u>A201634</u> <u>002</u>	Jun 13, 2012
<u>AB</u>			<u>23MG</u>	<u>A203419</u> <u>001</u>	Apr 12, 2016
<u>AB</u>		JUBILANT GENERICS	<u>5MG</u>	<u>A090768</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090768</u> <u>002</u>	May 31, 2011
<u>AB</u>		LUPIN LTD	<u>23MG</u>	<u>A202782</u> <u>001</u>	Oct 30, 2015
<u>AB</u>		MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201146</u> <u>001</u>	Aug 17, 2012
<u>AB</u>			<u>10MG</u>	<u>A201146</u> <u>002</u>	Aug 17, 2012
<u>AB</u>			<u>23MG</u>	<u>A202631</u> <u>001</u>	Jan 22, 2014
<u>AB</u>		MYLAN PHARMS INC	<u>5MG</u>	<u>A090521</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090521</u> <u>002</u>	May 31, 2011
<u>AB</u>			<u>23MG</u>	<u>A202656</u> <u>001</u>	Oct 22, 2015
<u>AB</u>		OSMOTICA PHARM US	<u>23MG</u>	<u>A203114</u> <u>001</u>	Jan 26, 2016
<u>AB</u>		PAR PHARM	<u>23MG</u>	<u>A202542</u> <u>001</u>	Jul 24, 2013
<u>AB</u>		PLIVA HRVATSKA DOO	<u>5MG</u>	<u>A090425</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090425</u> <u>002</u>	May 31, 2011
<u>AB</u>		PRINSTON INC	<u>5MG</u>	<u>A200292</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A200292</u> <u>002</u>	May 31, 2011
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A090290</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090290</u> <u>002</u>	May 31, 2011
<u>AB</u>		SCIEGEN PHARMS INC	<u>5MG</u>	<u>A203907</u> <u>001</u>	Oct 29, 2014
<u>AB</u>			<u>10MG</u>	<u>A203907</u> <u>002</u>	Oct 29, 2014
<u>AB</u>		SUN PHARM INDS	<u>5MG</u>	<u>A090493</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090493</u> <u>002</u>	May 31, 2011
<u>AB</u>		SUN PHARM INDS LTD	<u>5MG</u>	<u>A076786</u> <u>001</u>	Nov 26, 2010
<u>AB</u>			<u>10MG</u>	<u>A076786</u> <u>002</u>	Nov 26, 2010
<u>AB</u>			<u>23MG</u>	<u>A204293</u> <u>001</u>	Jun 05, 2015
<u>AB</u>		TEVA	<u>5MG</u>	<u>A077344</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A077344</u> <u>002</u>	May 31, 2011
<u>AB</u>		TORRENT PHARMS	<u>5MG</u>	<u>A090686</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090686</u> <u>002</u>	May 31, 2011
<u>AB</u>		TWI PHARMS INC	<u>23MG</u>	<u>A203104</u> <u>001</u>	Oct 29, 2014
<u>AB</u>		UNICHEM LABS LTD	<u>5MG</u>	<u>A203656</u> <u>001</u>	Jun 23, 2016
<u>AB</u>			<u>10MG</u>	<u>A203656</u> <u>002</u>	Jun 23, 2016
<u>AB</u>		VIVIMED GLOBAL	<u>5MG</u>	<u>A090551</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A090551</u> <u>002</u>	May 31, 2011
<u>AB</u>		WOCKHARDT	<u>5MG</u>	<u>A091267</u> <u>001</u>	May 31, 2011
<u>AB</u>			<u>10MG</u>	<u>A091267</u> <u>002</u>	May 31, 2011
<u>AB</u>		ZHEJIANG HISUN PHARM	<u>23MG</u>	<u>A202410</u> <u>001</u>	Mar 24, 2017
<u>AB</u>		ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A090100</u> <u>001</u>	Oct 24, 2012
<u>AB</u>			<u>10MG</u>	<u>A090100</u> <u>002</u>	Oct 24, 2012
<u>AB</u>			<u>23MG</u>	<u>A203162</u> <u>001</u>	Aug 31, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-141 (of 436)

DONEPEZIL HYDROCHLORIDE

TABLET, ORALLY DISINTEGRATING; ORAL

ARICEPT ODT

<u>AB</u>	+	EISAI INC	<u>5MG</u>	<u>N021720 001</u>	Oct 18, 2004
<u>AB</u>	+!		<u>10MG</u>	<u>N021720 002</u>	Oct 18, 2004

DONEPEZIL HYDROCHLORIDE

<u>AB</u>	BARR	<u>5MG</u>	<u>A078388 002</u>	Nov 26, 2010
<u>AB</u>		<u>10MG</u>	<u>A078388 001</u>	Nov 26, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A201787 001</u>	Dec 14, 2012
<u>AB</u>		<u>10MG</u>	<u>A201787 002</u>	Dec 14, 2012
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A091198 001</u>	May 10, 2011
<u>AB</u>		<u>10MG</u>	<u>A091198 002</u>	May 10, 2011
<u>AB</u>	UNICHEM LABS LTD	<u>5MG</u>	<u>A204831 001</u>	Nov 10, 2016
<u>AB</u>		<u>10MG</u>	<u>A204831 002</u>	Nov 10, 2016
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A090175 001</u>	May 10, 2011
<u>AB</u>		<u>10MG</u>	<u>A090175 002</u>	May 10, 2011

DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>10MG;14MG</u>	<u>A208328 001</u>	Jan 27, 2017
<u>AB</u>		<u>10MG;28MG</u>	<u>A208328 002</u>	Jan 27, 2017

NAMZARIC

<u>AB</u>	+	FOREST LABS LLC	<u>10MG;14MG</u>	<u>N206439 001</u>	Dec 23, 2014
<u>AB</u>	+!		<u>10MG;28MG</u>	<u>N206439 002</u>	Dec 23, 2014
	+		10MG;7MG	N206439 003	Jul 18, 2016
	+		10MG;21MG	N206439 004	Jul 18, 2016

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

<u>AP</u>	++!	HOSPIRA	<u>40MG/ML</u>	<u>N018132 001</u>	
<u>AP</u>	++!		<u>80MG/100ML</u>	<u>N018132 002</u>	Feb 04, 1982
<u>AP</u>	++!		<u>80MG/ML</u>	<u>N018132 004</u>	Jul 09, 1982
<u>AP</u>	++!		<u>160MG/100ML</u>	<u>N018132 003</u>	Feb 04, 1982
<u>AP</u>	!	LUITPOLD	<u>40MG/ML</u>	<u>A070799 001</u>	Feb 11, 1987
<u>AP</u>	!		<u>80MG/ML</u>	<u>A070820 001</u>	Feb 11, 1987

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%

<u>AP</u>	++!	B BRAUN	<u>80MG/100ML</u>	<u>N019099 002</u>	Oct 15, 1986
<u>AP</u>	++!		<u>320MG/100ML</u>	<u>N019099 004</u>	Oct 15, 1986

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	++!	B BRAUN	<u>160MG/100ML</u>
-----------	-----	---------	--------------------

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	++!	BAXTER HLTHCARE	<u>80MG/100ML</u>
<u>AP</u>	++!		<u>160MG/100ML</u>
<u>AP</u>	++!		<u>320MG/100ML</u>
<u>AP</u>	++!	HOSPIRA	<u>80MG/100ML</u>
<u>AP</u>	++!		<u>160MG/100ML</u>
<u>AP</u>	++!		<u>320MG/100ML</u>

DOPAMINE HYDROCHLORIDE

! LUITPOLD 160MG/ML

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER

++! B BRAUN 40MG/100ML

DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

++! BAXTER HLTHCARE 640MG/100ML

<u>N018132 001</u>	Mar 27, 1987
<u>N018132 002</u>	Mar 27, 1987
<u>N018132 004</u>	Mar 27, 1987
<u>N018132 003</u>	Mar 27, 1987
<u>A070799 001</u>	Sep 30, 1983
<u>N018826 002</u>	Sep 30, 1983
<u>N018826 003</u>	Sep 30, 1983

A070826 001 Feb 11, 1987

N019099 001 Oct 15, 1986

N019615 004 Mar 27, 1987

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE

<u>AT</u>	Bausch and Lomb	<u>EQ 2% BASE</u>	<u>A090143 001</u>	Jun 25, 2009
<u>AT</u>	HI TECH PHARMA	<u>EQ 2% BASE</u>	<u>A077846 001</u>	Oct 28, 2008
<u>AT</u>	LUITPOLD	<u>EQ 2% BASE</u>	<u>A079186 001</u>	Nov 18, 2009
<u>AT</u>	SANDOZ INC	<u>EQ 2% BASE</u>	<u>A078748 001</u>	Nov 06, 2008
<u>AT</u>		<u>EQ 2% BASE</u>	<u>A078981 001</u>	Apr 13, 2009
<u>AT</u>	TEVA PHARMS	<u>EQ 2% BASE</u>	<u>A078756 001</u>	Dec 04, 2008
<u>AT</u>	WATSON LABS INC	<u>EQ 2% BASE</u>	<u>A202053 001</u>	Sep 11, 2014
	<b>TRUSOPT</b>		<b>N020408 001</b>	Dec 09, 1994
<u>AT</u>	+! MERCK	<u>EQ 2% BASE</u>		

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-142 (of 436)

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

**COSOPT**

**AT** +! OAK PHARMS INC **EQ 2% BASE;EQ 0.5% BASE**

**N020869 001** Apr 07, 1998

**DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE**

**AT** AKORN INC **EQ 2% BASE;EQ 0.5% BASE**

**A203058 001** Sep 22, 2014

**AT** BAUSCH AND LOMB **EQ 2% BASE;EQ 0.5% BASE**

**A090037 001** Jul 14, 2009

**AT** HI TECH PHARMA **EQ 2% BASE;EQ 0.5% BASE**

**A077847 001** Oct 28, 2008

**AT** SANDOZ **EQ 2% BASE;EQ 0.5% BASE**

**A078749 001** Nov 06, 2008

**AT** SANDOZ INC **EQ 2% BASE;EQ 0.5% BASE**

**A090604 001** Nov 18, 2009

**AT** TEVA PHARMS **EQ 2% BASE;EQ 0.5% BASE**

**A078704 001** Sep 28, 2009

**AT** WATSON LABS INC **EQ 2% BASE;EQ 0.5% BASE**

**A202054 001** Sep 03, 2014

**AT** ZAMBON SPA **EQ 2% BASE;EQ 0.5% BASE**

**A091180 001** Dec 04, 2013

COSOPT PF

+! OAK PHARMS INC **EQ 2% BASE;EQ 0.5% BASE**

**N202667 001** Feb 01, 2012

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

**DOPRAM**

**AP** +! WEST-WARD PHARMS INT **20MG/ML**

**N014879 001**

**DOXAPRAM HYDROCHLORIDE**

**AP** ATHENEX INC **20MG/ML**

**A076266 001** Jan 10, 2003

DOXAZOSIN MESYLATE

TABLET; ORAL

**CARDURA**

**AB** +! PFIZER **EQ 1MG BASE**

**N019668 001** Nov 02, 1990

**AB** + **EQ 2MG BASE**

**N019668 002** Nov 02, 1990

**AB** + **EQ 4MG BASE**

**N019668 003** Nov 02, 1990

**AB** + **EQ 8MG BASE**

**N019668 004** Nov 02, 1990

**DOXAZOSIN MESYLATE**

**AB** ACCORD HLTHCARE **EQ 1MG BASE**

**A202824 001** Jun 11, 2014

**AB** **EQ 2MG BASE**

**A202824 002** Jun 11, 2014

**AB** **EQ 4MG BASE**

**A202824 003** Jun 11, 2014

**AB** **EQ 8MG BASE**

**A202824 004** Jun 11, 2014

**AB** APOTEX **EQ 1MG BASE**

**A075580 001** Oct 18, 2000

**AB** **EQ 2MG BASE**

**A075580 002** Oct 18, 2000

**AB** **EQ 4MG BASE**

**A075580 003** Oct 18, 2000

**AB** **EQ 8MG BASE**

**A075580 004** Oct 18, 2000

**AB** DAVA PHARMS INC **EQ 1MG BASE**

**A076161 001** Jun 10, 2004

**AB** **EQ 2MG BASE**

**A076161 002** Jun 10, 2004

**AB** **EQ 4MG BASE**

**A076161 003** Jun 10, 2004

**AB** **EQ 8MG BASE**

**A076161 004** Jun 10, 2004

**AB** MYLAN **EQ 1MG BASE**

**A075509 001** Oct 19, 2000

**AB** **EQ 2MG BASE**

**A075509 002** Oct 19, 2000

**AB** **EQ 4MG BASE**

**A075509 003** Oct 19, 2000

**AB** **EQ 8MG BASE**

**A075509 004** Oct 19, 2000

**AB** PLIVA **EQ 1MG BASE**

**A075750 001** Jun 08, 2001

**AB** **EQ 2MG BASE**

**A075750 002** Jun 08, 2001

**AB** **EQ 4MG BASE**

**A075750 003** Jun 08, 2001

**AB** **EQ 8MG BASE**

**A075750 004** Jun 08, 2001

**AB** TEVA **EQ 1MG BASE**

**A075536 001** Oct 18, 2000

**AB** **EQ 2MG BASE**

**A075536 002** Oct 18, 2000

**AB** **EQ 4MG BASE**

**A075536 003** Oct 18, 2000

**AB** **EQ 8MG BASE**

**A075536 004** Oct 18, 2000

**AB** ZYDUS PHARMS USA INC **EQ 1MG BASE**

**A208719 001** Jul 07, 2017

**AB** **EQ 2MG BASE**

**A208719 002** Jul 07, 2017

**AB** **EQ 4MG BASE**

**A208719 003** Jul 07, 2017

**AB** **EQ 8MG BASE**

**A208719 004** Jul 07, 2017

TABLET, EXTENDED RELEASE; ORAL

CARDURA XL

+! PFIZER **EQ 4MG BASE**

**N021269 001** Feb 22, 2005

+! **EQ 8MG BASE**

**N021269 002** Feb 22, 2005

DOXEPIPIN HYDROCHLORIDE

CAPSULE; ORAL

**DOXEPIPIN HYDROCHLORIDE**

**AB** AMNEAL PHARMS CO **EQ 10MG BASE**

**A207482 001** Jun 28, 2017

**AB** **EQ 25MG BASE**

**A207482 002** Jun 28, 2017

**AB** **EQ 50MG BASE**

**A207482 003** Jun 28, 2017

**AB** **EQ 75MG BASE**

**A207482 004** Jun 28, 2017

**AB** **EQ 100MG BASE**

**A207482 005** Jun 28, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-143 (of 436)

DOXEPEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPEPIN HYDROCHLORIDE

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A070791 002</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A070791 003</u>	May 13, 1986
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A070791 001</u>	May 13, 1986
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A070791 004</u>	May 13, 1986
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A070791 005</u>	May 13, 1986

! PAR PHARM

CONCENTRATE; ORAL

DOXEPEPIN HYDROCHLORIDE

<u>AA</u>	SILARX	<u>EQ 10MG BASE/ML</u>	<u>A074721 001</u>	Dec 29, 1998
<u>AA</u>	!	<u>EQ 10MG BASE/ML</u>	<u>A071609 001</u>	Nov 09, 1987
<u>AA</u>	TEVA PHARMS	<u>EQ 10MG BASE/ML</u>	<u>A071918 001</u>	Jul 20, 1988
	WOCKHARDT BIO AG			
	CREAM; TOPICAL			
	ZONALON			
	+ ! MYLAN PHARMS INC	5%	N020126 001	Apr 01, 1994
	TABLET; ORAL			

DOXEPEPIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 3MG BASE</u>	<u>A201951 001</u>	Jul 26, 2013
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A201951 002</u>	Jul 26, 2013
	SILENOR			
<u>AB</u>	+	PERNIX THERAPS LLC	<u>EQ 3MG BASE</u>	N022036 001 Mar 17, 2010
<u>AB</u>	++!		<u>EQ 6MG BASE</u>	N022036 002 Mar 17, 2010

DOXERCALCIFEROL

CAPSULE; ORAL

DOXERCALCIFEROL

<u>AB</u>	RISING PHARMS INC	<u>0 .5MCG</u>	<u>A201518 001</u>	Sep 09, 2016
<u>AB</u>		<u>1MCG</u>	<u>A201518 002</u>	Sep 09, 2016
<u>AB</u>		<u>2 .5MCG</u>	<u>A201518 003</u>	Sep 09, 2016
<u>AB</u>	WEST-WARD PHARMS INT	<u>0 .5MCG</u>	<u>A091433 001</u>	Sep 23, 2011
<u>AB</u>		<u>1MCG</u>	<u>A091433 002</u>	Jan 14, 2014
<u>AB</u>		<u>2 .5MCG</u>	<u>A091433 003</u>	Jan 14, 2014
	HECTOROL			
<u>AB</u>	+	GENZYME CORP	<u>0 .5MCG</u>	N020862 002 Apr 23, 2004
<u>AB</u>	+		<u>1MCG</u>	N020862 003 Jul 13, 2009
<u>AB</u>	++!		<u>2 .5MCG</u>	N020862 001 Jun 09, 1999

INJECTABLE; INJECTION

DOXERCALCIFEROL

<u>AP</u>	AKORN INC	<u>2MCG/ML (2MCG/ML)</u>	<u>A203929 002</u>	Mar 28, 2016
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A203929 001</u>	May 07, 2015
<u>AP</u>	AMNEAL PHARMS CO	<u>2MCG/ML (2MCG/ML)</u>	<u>A208974 001</u>	May 24, 2017
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A208974 002</u>	May 24, 2017
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A208975 001</u>	May 24, 2017
<u>AP</u>	HIKMA PHARMS	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091101 001</u>	Aug 30, 2013
<u>AP</u>	SANDOZ INC	<u>4MCG/2ML (2MCG/ML)</u>	<u>A091333 001</u>	May 05, 2014
<u>AP</u>		<u>4MCG/2ML (2MCG/ML)</u>	<u>A200926 001</u>	Feb 04, 2014
	HECTOROL			
<u>AP</u>	+	GENZYME CORP	<u>2MCG/ML (2MCG/ML)</u>	N021027 002 Apr 06, 2000
<u>AP</u>	++!		<u>4MCG/2ML (2MCG/ML)</u>	N021027 001 Apr 06, 2000

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>	ACTAVIS INC	<u>2MG/ML</u>	<u>A203622 001</u>	Jun 27, 2014
<u>AP</u>		<u>200MG/100ML</u>	<u>A203622 002</u>	Jun 27, 2014
<u>AP</u>	ALVOGEN INC	<u>2MG/ML</u>	<u>A065515 001</u>	Nov 08, 2012
<u>AP</u>	AMNEAL PHARMS CO	<u>20MG/VIAL</u>	<u>A208888 001</u>	Feb 17, 2017
<u>AP</u>		<u>50MG/VIAL</u>	<u>A208888 002</u>	Feb 17, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>2MG/ML</u>	<u>A063277 001</u>	Oct 26, 1995
<u>AP</u>	GLAND PHARMA LTD	<u>2MG/ML</u>	<u>A209825 001</u>	Aug 11, 2017
<u>AP</u>	MYLAN LABS LTD	<u>2MG/ML</u>	<u>A200901 001</u>	Feb 14, 2012
<u>AP</u>		<u>10MG/VIAL</u>	<u>A200170 001</u>	Oct 28, 2011
<u>AP</u>		<u>50MG/VIAL</u>	<u>A200170 002</u>	Oct 28, 2011
<u>AP</u>	PHARMACHEMIE BV	<u>2MG/ML</u>	<u>A063336 001</u>	Feb 28, 1995
<u>AP</u>		<u>10MG/VIAL</u>	<u>A063097 001</u>	May 21, 1990
<u>AP</u>		<u>20MG/VIAL</u>	<u>A063097 002</u>	May 21, 1990
<u>AP</u>		<u>50MG/VIAL</u>	<u>A063097 003</u>	May 21, 1990
<u>AP</u>		<u>200MG/100ML</u>	<u>A063336 004</u>	Feb 28, 1995
<u>AP</u>	++!	PHARMACIA AND UPJOHN	<u>2MG/ML</u>	N050629 001 Dec 23, 1987
<u>AP</u>	++!		<u>200MG/100ML</u>	N050629 002 May 03, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-144 (of 436)

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>	SAGENT PHARMS	<u>2MG/ML</u>	<u>A091495</u> <u>001</u>	Mar 18, 2013
<u>AP</u>	SUN PHARM INDs	<u>2MG/ML</u>	<u>A091418</u> <u>001</u>	Feb 15, 2012
<u>AP</u>	TEVA PHARMS USA	<u>2MG/ML</u>	<u>A064140</u> <u>001</u>	Jul 28, 1995
<u>AP</u>		<u>200MG/100ML</u>	<u>A064140</u> <u>002</u>	Jul 28, 1995
<u>AP</u>	WEST-WARD PHARMS INT	<u>2MG/ML</u>	<u>A062975</u> <u>001</u>	Mar 17, 1989
<u>AP</u> !		<u>10MG/VIAL</u>	<u>A062921</u> <u>001</u>	Mar 17, 1989
<u>AP</u> !		<u>20MG/VIAL</u>	<u>A062921</u> <u>002</u>	Mar 17, 1989
<u>AP</u> !		<u>50MG/VIAL</u>	<u>A062921</u> <u>003</u>	Mar 17, 1989
<u>AP</u>		<u>200MG/100ML</u>	<u>A064097</u> <u>001</u>	Sep 13, 1994
+ PHARMACIA AND UPJOHN		150MG/75ML	N050629 003	Mar 28, 2011

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL (LIPOSOMAL)

<u>AB</u> +	JANSSEN RES AND DEV	<u>20MG/10ML (2MG/ML)</u>	<u>N050718</u> <u>001</u>	Nov 17, 1995
<u>AB</u> +		<u>50MG/25ML (2MG/ML)</u>	<u>N050718</u> <u>002</u>	Jun 13, 2000
<u>DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL)</u>				
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG/10ML (2MG/ML)</u>	<u>A208657</u> <u>001</u>	May 15, 2017
<u>AB</u>		<u>50MG/25ML (2MG/ML)</u>	<u>A208657</u> <u>002</u>	May 15, 2017
<u>AB</u> !	SUN PHARMA GLOBAL	<u>20MG/10ML (2MG/ML)</u>	<u>A203263</u> <u>001</u>	Feb 04, 2013
<u>AB</u> !		<u>50MG/25ML (2MG/ML)</u>	<u>A203263</u> <u>002</u>	Feb 04, 2013

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 75MG BASE</u>	<u>A209165</u> <u>001</u>	Jul 28, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209165</u> <u>002</u>	Jul 28, 2017
<u>AB</u>	G AND W LABS INC	<u>EQ 50MG BASE</u>	<u>A204446</u> <u>001</u>	May 28, 2015
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A204446</u> <u>002</u>	May 28, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204446</u> <u>003</u>	May 28, 2015
<u>AB</u>	IMPAX LABS INC	<u>EQ 150MG BASE</u>	<u>A200065</u> <u>001</u>	Feb 17, 2011
<u>AB</u>	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204234</u> <u>001</u>	Mar 05, 2014
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A204234</u> <u>002</u>	Mar 05, 2014
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A204234</u> <u>003</u>	Mar 05, 2014
<u>AB</u>	MAYNE PHARMA INC	<u>EQ 50MG BASE</u>	<u>A209396</u> <u>001</u>	Sep 29, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A209396</u> <u>002</u>	Sep 29, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209396</u> <u>003</u>	Sep 29, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 150MG BASE</u>	<u>A202778</u> <u>001</u>	Jun 08, 2012
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A065055</u> <u>001</u>	Dec 01, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065055</u> <u>002</u>	Dec 01, 2000
<u>AB</u> !		<u>EQ 150MG BASE</u>	<u>A065055</u> <u>003</u>	Jul 15, 2005
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 50MG BASE</u>	<u>A065053</u> <u>001</u>	Nov 22, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065053</u> <u>003</u>	Sep 10, 2003
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065053</u> <u>002</u>	Nov 22, 2000
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 50MG BASE</u>	<u>A205115</u> <u>001</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A205115</u> <u>002</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A205115</u> <u>003</u>	Feb 18, 2016

MONODOX

<u>AB</u> +	AQUA PHARMS	<u>EQ 50MG BASE</u>	<u>N050641</u> <u>002</u>	Feb 10, 1992
<u>AB</u> +		<u>EQ 75MG BASE</u>	<u>N050641</u> <u>003</u>	Oct 18, 2006
<u>AB</u> +!		<u>EQ 100MG BASE</u>	<u>N050641</u> <u>001</u>	Dec 29, 1989

ORACEA

+! GALDERMA LABS LP

40MG

N050805 001 May 26, 2006

FOR SUSPENSION; ORAL

DOXYCYCLINE

<u>AB</u>	CHARTWELL LIFE SCI	<u>EQ 25MG BASE/5ML</u>	<u>A065454</u> <u>001</u>	Jul 16, 2008
<u>AB</u>	LUPIN LTD	<u>EQ 25MG BASE/5ML</u>	<u>A201678</u> <u>001</u>	Mar 18, 2013

VIBRAMYCIN

<u>AB</u> +!	Pfizer	<u>EQ 25MG BASE/5ML</u>	<u>N050006</u> <u>001</u>	
--------------	--------	-------------------------	---------------------------	--

TABLET; ORAL

DOXYCYCLINE

<u>AB</u>	HERITAGE PHARMS INC	<u>EQ 50MG BASE</u>	<u>A091605</u> <u>001</u>	Dec 20, 2011
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091605</u> <u>002</u>	Dec 20, 2011
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A091605</u> <u>003</u>	Dec 20, 2011
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091605</u> <u>004</u>	Dec 20, 2011
<u>AB</u>	LANNETT	<u>EQ 50MG BASE</u>	<u>A065285</u> <u>001</u>	Dec 08, 2005
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065285</u> <u>003</u>	Jul 30, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065285</u> <u>002</u>	Dec 08, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065285</u> <u>004</u>	Jul 30, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-145 (of 436)

DOXYCYCLINE

TABLET;ORAL

DOXYCYCLINE

<u>AB</u>	MYLAN	<u>EQ 50MG BASE</u>	<u>A065377 001</u>	Nov 07, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065377 002</u>	Nov 07, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065377 003</u>	Nov 07, 2006
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A065427 001</u>	Jun 07, 2007
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A065070 001</u>	Dec 15, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065070 003</u>	Dec 30, 2002
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065070 002</u>	Dec 15, 2000
<u>AB</u>	!	<u>EQ 150MG BASE</u>	<u>A065070 004</u>	Jul 14, 2005
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 50MG BASE</u>	<u>A065356 001</u>	May 31, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065356 002</u>	May 31, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065356 003</u>	May 31, 2006
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 50MG BASE</u>	<u>A209582 001</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A209582 002</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A209582 003</u>	Sep 28, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209582 004</u>	Sep 28, 2017

DOXYCYCLINE CALCIUM

SUSPENSION;ORAL

VIBRAMYCIN

+! PFIZER

EQ 50MG BASE/5ML

N050480 001

DOXYCYCLINE HYCLATE

CAPSULE;ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 50MG BASE</u>	<u>A062031 002</u>	Oct 13, 1982
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062031 001</u>	
<u>AB</u>	AMNEAL PHARMS	<u>EQ 100MG BASE</u>	<u>A207289 001</u>	Jun 27, 2016
<u>AB</u>	CHARTWELL LIFE SCI	<u>EQ 50MG BASE</u>	<u>A062500 001</u>	Sep 11, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062500 002</u>	Sep 11, 1984
<u>AB</u>	HIKMA INTL PHARMS	<u>EQ 50MG BASE</u>	<u>A062396 002</u>	Nov 07, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062396 001</u>	May 07, 1984
<u>AB</u>	MUTUAL PHARM	<u>EQ 50MG BASE</u>	<u>A062675 001</u>	Jul 10, 1986
<u>AB</u>	MYLAN	<u>EQ 50MG BASE</u>	<u>A062337 001</u>	Mar 29, 1982
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A062337 002</u>	Mar 29, 1982
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 100MG BASE</u>	<u>A062676 001</u>	Jul 10, 1986

VIBRAMYCIN

<u>AB</u>	+! PFIZER	<u>EQ 100MG BASE</u>	<u>N050007 002</u>	
	ACTICLATE CAP			
	+! AQUA PHARMS	EQ 75MG BASE		N208253 001 Apr 26, 2016
	DOXYCYCLINE HYCLATE			
	+! HIKMA INTL PHARMS	EQ 20MG BASE		A065103 001 May 13, 2005

INJECTABLE;INJECTION

DOXY 100

<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 100MG BASE/VIAL</u>	<u>A062475 001</u>	Dec 09, 1983
<u>AP</u>	DOXY 200	<u>EQ 200MG BASE/VIAL</u>	<u>A062475 002</u>	Dec 09, 1983
<u>AP</u>	! FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>		
<u>AP</u>	<u>DOXYCYCLINE</u>			
<u>AP</u>	MYLAN LABS LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A091406 001</u>	Aug 21, 2012
<u>AP</u>	! WEST-WARD PHARMS INT	<u>EQ 100MG BASE/VIAL</u>	<u>A062569 001</u>	Mar 09, 1988
<u>AP</u>	ZYDUS PHARMS USA INC	<u>EQ 100MG BASE/VIAL</u>	<u>A207757 001</u>	Sep 28, 2017
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A207757 002</u>	Sep 28, 2017

SYSTEM, EXTENDED RELEASE;PERIODONTAL

ATRIDOX

+! TOLMAR

50MG

N050751 001 Sep 03, 1998

TABLET;ORAL

ACTICLATE

<u>AB</u>	+ AQUA PHARMS LLC	<u>EQ 75MG BASE</u>	<u>N205931 001</u>	Jul 25, 2014
<u>AB</u>	+!	<u>EQ 150MG BASE</u>	<u>N205931 002</u>	Jul 25, 2014

DOXYCYCLINE HYCLATE

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 100MG BASE</u>	<u>A062421 001</u>	Feb 02, 1983
<u>AB</u>	AMNEAL PHARMS CO	<u>EQ 75MG BASE</u>	<u>A209372 001</u>	Oct 06, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A209372 002</u>	Oct 06, 2017
<u>AB</u>	CARIBE HOLDINGS	<u>EQ 100MG BASE</u>	<u>A062269 002</u>	Nov 08, 1982
<u>AB</u>	CHARTWELL LIFE SCI	<u>EQ 100MG BASE</u>	<u>A062505 001</u>	Sep 11, 1984
<u>AB</u>	! HIKMA INTL PHARMS	<u>EQ 100MG BASE</u>	<u>A065095 001</u>	Jul 02, 2003
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A065163 001</u>	May 13, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-146 (of 436)

DOXYCYCLINE HYCLATE

TABLET;ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	!	LANNETT	<u>EQ 20MG BASE</u>	<u>A065277 001</u>	Nov 10, 2005
<u>AB</u>		LARKEN LABS	<u>EQ 20MG BASE</u>	<u>A065287 001</u>	Feb 28, 2006
<u>AB</u>		LUPIN LTD	<u>EQ 75MG BASE</u>	<u>A208818 001</u>	Sep 27, 2017
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A208818 002</u>	Sep 27, 2017
<u>AB</u>		MAYNE PHARMA INC	<u>EQ 75MG BASE</u>	<u>A208765 001</u>	Jun 14, 2017
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A208765 002</u>	Jun 14, 2017
<u>AB</u>		MYLAN	<u>EQ 100MG BASE</u>	<u>A062432 001</u>	Feb 15, 1983
<u>AB</u>		NOVEL LABS INC	<u>EQ 100MG BASE</u>	<u>A207558 001</u>	Sep 06, 2017
<u>AB</u>		SUN PHARM INDUSTRIES	<u>EQ 20MG BASE</u>	<u>A065134 001</u>	May 13, 2005
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A062677 001</u>	Jul 10, 1986
<u>AB</u>		ZYDUS PHARMS USA INC CARIBE HOLDINGS	<u>EQ 100MG BASE</u>	<u>A207773 001</u>	Oct 30, 2017
		TABLET, DELAYED RELEASE;ORAL	EQ 50MG BASE		A062269 003

DORYX

<u>AB</u>	+	MAYNE PHARMA	<u>EQ 50MG BASE</u>	<u>N050795 006</u>	Dec 19, 2014
<u>AB</u>	+		<u>EQ 75MG BASE</u>	<u>N050795 001</u>	May 06, 2005
<u>AB</u>	+		<u>EQ 80MG BASE</u>	<u>N050795 004</u>	Apr 11, 2013
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N050795 002</u>	May 06, 2005
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N050795 003</u>	Jun 20, 2008
<u>AB</u>	+!		<u>EQ 200MG BASE</u>	<u>N050795 005</u>	Apr 11, 2013

DOXYCYCLINE HYCLATE

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 75MG BASE</u>	<u>A090134 001</u>	Dec 14, 2011
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A090134 002</u>	Dec 14, 2011
<u>AB</u>		HERITAGE PHARMS INC	<u>EQ 75MG BASE</u>	<u>A200856 001</u>	Apr 30, 2013
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A200856 002</u>	Apr 30, 2013
<u>AB</u>			<u>EQ 150MG BASE</u>	<u>A200856 003</u>	Apr 30, 2013
<u>AB</u>		MYLAN	<u>EQ 50MG BASE</u>	<u>A090431 003</u>	May 23, 2016
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A090431 001</u>	Dec 28, 2010
<u>AB</u>			<u>EQ 80MG BASE</u>	<u>A090431 004</u>	Apr 29, 2016
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A090431 002</u>	Dec 28, 2010
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A090431 005</u>	May 19, 2016
<u>AB</u>		MYLAN PHARMS INC	<u>EQ 150MG BASE</u>	<u>A091052 001</u>	Feb 08, 2012
<u>AB</u>		PRINSTON INC	<u>EQ 150MG BASE</u>	<u>A207494 001</u>	Nov 15, 2016
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>A207494 002</u>	Nov 15, 2016

DORYX MPC

+! MAYNE PHARMA EQ 120MG BASE

N050795 008 May 20, 2016

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, DELAYED RELEASE;ORAL

DICLEGIS

<u>AB</u>	+!	DUCHESNAY	<u>10MG;10MG</u>	<u>N021876 001</u>	Apr 08, 2013
			<u>DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE</u>		
<u>AB</u>		ACTAVIS LABS FL INC	<u>10MG;10MG</u>	<u>A205811 001</u>	Aug 19, 2016
<u>AB</u>		PAR PHARM INC	<u>10MG;10MG</u>	<u>A208518 001</u>	Dec 06, 2017
		TABLET, EXTENDED RELEASE;ORAL			
		BONJESTA			
	+!	DUCHESNAY	20MG;20MG		

N209661 001 Nov 07, 2016

DRONABINOL

CAPSULE;ORAL

DRONABINOL

<u>AB</u>		AKORN INC	<u>2 .5MG</u>	<u>A079217 001</u>	Jun 20, 2014
<u>AB</u>			<u>5MG</u>	<u>A079217 002</u>	Jun 20, 2014
<u>AB</u>			<u>10MG</u>	<u>A079217 003</u>	Jun 20, 2014
<u>AB</u>		SVC PHARMA	<u>2 .5MG</u>	<u>A078292 001</u>	Jun 27, 2008
<u>AB</u>			<u>5MG</u>	<u>A078292 002</u>	Jun 27, 2008
<u>AB</u>			<u>10MG</u>	<u>A078292 003</u>	Jun 27, 2008

MARINOL

<u>AB</u>	+	ABBVIE	<u>2 .5MG</u>	<u>N018651 001</u>	May 31, 1985
<u>AB</u>	+!		<u>5MG</u>	<u>N018651 002</u>	May 31, 1985
<u>AB</u>	+		<u>10MG</u>	<u>N018651 003</u>	May 31, 1985
		SOLUTION;ORAL			
		SYNDROS			
	+!	INSYS DEV CO INC	5MG/ML		

N205525 001 Mar 23, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-147 (of 436)

DRONEDARONE HYDROCHLORIDE

TABLET;ORAL

MULTAQ

+! SANOFI AVENTIS US EQ 400MG BASE

N022425 001 Jul 01, 2009

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

<u>AP</u>	EUROHLTH INTL SARL	<u>2.5MG/ML</u>	<u>A208197 001</u> Dec 14, 2017
<u>AP</u>	HOSPIRA	<u>2.5MG/ML</u>	<u>A071981 001</u> Feb 29, 1988
<u>AP</u>	LUITPOLD	<u>2.5MG/ML</u>	<u>A072123 001</u> Oct 24, 1988
		<u>INAPSINE</u>	
<u>AP</u>	+! AKORN INC	<u>2.5MG/ML</u>	<u>N016796 001</u>

DROSPIRENONE; ESTRADIOL

TABLET;ORAL

ANGELIQ

+ BAYER HLTHCARE 0.25MG; 0.5MG  
+! 0.5MG; 1MG

N021355 001 Feb 29, 2012  
N021355 002 Sep 28, 2005

DROSPIRENONE; ETHINYLMESTRADIOL

TABLET;ORAL

DROSPIRENONE AND ETHINYLMESTRADIOL

<u>AB</u>	BARR	<u>3MG;0.02MG</u>	<u>A078515 001</u> Mar 30, 2009
<u>AB</u>	CYNDEA PHARMA	<u>3MG;0.02MG</u>	<u>A209423 001</u> Dec 22, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>3MG;0.02MG</u>	<u>A204296 001</u> Aug 17, 2015
<u>AB</u>	MYLAN LABS LTD	<u>3MG;0.02MG</u>	<u>A202594 001</u> Oct 22, 2015
<u>AB</u>	PII	<u>3MG;0.02MG</u>	<u>A203291 001</u> Jul 18, 2017
<u>AB</u>	WATSON LABS	<u>3MG;0.02MG</u>	<u>A078833 001</u> Nov 28, 2011
		<u>LORYNA</u>	
<u>AB</u>	LABS LEON FARMA	<u>3MG;0.02MG</u>	<u>A079221 001</u> Mar 28, 2011
		<u>MELAMISA</u>	
<u>AB</u>	NOVAST LABS LTD	<u>3MG;0.02MG</u>	<u>A202016 001</u> Jan 26, 2016
		<u>NIKKI</u>	
<u>AB</u>	LUPIN LTD	<u>3MG;0.02MG</u>	<u>A201661 001</u> May 27, 2014
		<u>YAZ</u>	
<u>AB</u>	+! BAYER HLTHCARE	<u>3MG;0.02MG</u>	<u>N021676 001</u> Mar 16, 2006

TABLET;ORAL-28

DROSPIRENONE AND ETHINYLMESTRADIOL

<u>AB</u>	ACCORD HLTHCARE	<u>3MG;0.03MG</u>	<u>A207245 001</u> Nov 22, 2016
<u>AB</u>	APOTEX INC	<u>3MG;0.03MG</u>	<u>A205876 001</u> Sep 21, 2016
<u>AB</u>	BARR	<u>3MG;0.03MG</u>	<u>A077527 001</u> May 09, 2008
<u>AB</u>	GLENMARK PHARMS LTD	<u>3MG;0.03MG</u>	<u>A204848 001</u> Mar 25, 2016
<u>AB</u>	LUPIN LTD	<u>3MG;0.03MG</u>	<u>A201663 001</u> Dec 18, 2012
<u>AB</u>	MAYNE PHARMA	<u>3MG;0.03MG</u>	<u>A090081 001</u> Sep 07, 2010
<u>AB</u>	MYLAN LABS LTD	<u>3MG;0.03MG</u>	<u>A202131 001</u> May 04, 2015
		<u>SYEDA</u>	
<u>AB</u>	LABS LEON FARMA	<u>3MG;0.03MG</u>	<u>A090114 001</u> Mar 28, 2011
		<u>YAEILA</u>	
<u>AB</u>	NOVAST LABS LTD	<u>3MG;0.03MG</u>	<u>A202015 001</u> Nov 19, 2014
		<u>YASMIN</u>	
<u>AB</u>	+! BAYER HLTHCARE	<u>3MG;0.03MG</u>	<u>N021098 001</u> May 11, 2001

DROSPIRENONE; ETHINYLMESTRADIOL; LEVOMEFOLATE CALCIUM

TABLET;ORAL

BEYAZ

<u>AB</u>	BAYER HLTHCARE	<u>3MG,N/A:0.02MG,N/A:0.451MG,0.451MG</u>	<u>N022532 001</u> Sep 24, 2010
		<u>DROSPIRENONE, ETHINYLMESTRADIOL AND LEVOMEFOLATE CALCIUM</u>	
<u>AB</u>	WATSON LABS INC	<u>3MG,N/A:0.02MG,N/A:0.451MG,0.451MG</u>	<u>A203593 001</u> Oct 11, 2016
<u>AB</u>		<u>3MG,N/A:0.03MG,N/A:0.451MG,0.451MG</u>	<u>A203594 001</u> Oct 11, 2016
		<u>SAFYRAL</u>	
<u>AB</u>	+! BAYER HLTHCARE	<u>3MG,N/A:0.03MG,N/A:0.451MG,0.451MG</u>	<u>N022574 001</u> Dec 16, 2010
		<u>TYDEMY</u>	
<u>AB</u>	LUPIN LTD	<u>3MG,N/A:0.03MG,N/A:0.451MG,0.451MG</u>	<u>A205948 001</u> Dec 12, 2017

DROXIDOPA

CAPSULE;ORAL

NORTHERA

+ LUNDBECK NA LTD 100MG  
+ 200MG  
+! 300MG

N203202 001 Feb 18, 2014  
N203202 002 Feb 18, 2014  
N203202 003 Feb 18, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-148 (of 436)

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

CYMBALTA

<u>AB</u>	+	LILLY	<u>EQ 20MG BASE</u>	<u>N021427 001</u>	Aug 03, 2004
<u>AB</u>	+		<u>EQ 30MG BASE</u>	<u>N021427 002</u>	Aug 03, 2004
<u>AB</u>	+!		<u>EQ 60MG BASE</u>	<u>N021427 004</u>	Aug 03, 2004
		<u>DULOXETINE HYDROCHLORIDE</u>			
<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 20MG BASE</u>	<u>A090776 001</u>	Dec 17, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090776 002</u>	Dec 17, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090776 003</u>	Dec 17, 2013
<u>AB</u>		AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A208706 001</u>	Jan 06, 2017
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A208706 002</u>	Jan 06, 2017
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A208706 003</u>	Jan 06, 2017
<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A202949 001</u>	Jun 09, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A202949 002</u>	Jun 09, 2014
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A202949 003</u>	Jun 09, 2014
<u>AB</u>		ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A203197 001</u>	Aug 26, 2015
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A203197 002</u>	Aug 26, 2015
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A203197 003</u>	Aug 26, 2015
<u>AB</u>		ANCHEN PHARMS	<u>EQ 20MG BASE</u>	<u>A090780 001</u>	Oct 28, 2015
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090780 002</u>	Oct 28, 2015
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090780 003</u>	Oct 28, 2015
<u>AB</u>		APOTEX INC	<u>EQ 20MG BASE</u>	<u>A202045 001</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A202045 002</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A202045 003</u>	Jun 11, 2014
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A090778 001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090778 002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090778 003</u>	Dec 11, 2013
<u>AB</u>		BRECKENRIDGE PHARM	<u>EQ 20MG BASE</u>	<u>A203088 001</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A203088 002</u>	Jun 11, 2014
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A203088 003</u>	Jun 11, 2014
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A090723 001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090723 002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090723 003</u>	Dec 11, 2013
<u>AB</u>		HETERO LABS LTD III	<u>EQ 20MG BASE</u>	<u>A204343 001</u>	Aug 03, 2016
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A204343 002</u>	Aug 03, 2016
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A204343 003</u>	Aug 03, 2016
<u>AB</u>		INVENTIA HLTHCARE	<u>EQ 20MG BASE</u>	<u>A202336 001</u>	Oct 28, 2015
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A202336 002</u>	Oct 28, 2015
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A202336 003</u>	Oct 28, 2015
<u>AB</u>		LUPIN LTD	<u>EQ 20MG BASE</u>	<u>A090694 001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090694 002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090694 004</u>	Dec 11, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A204815 001</u>	Mar 23, 2017
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A204815 002</u>	Mar 23, 2017
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A204815 003</u>	Mar 23, 2017
<u>AB</u>		PRINSTON INC	<u>EQ 20MG BASE</u>	<u>A206653 001</u>	May 18, 2017
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A206653 002</u>	May 18, 2017
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A206653 003</u>	May 18, 2017
<u>AB</u>		SUN PHARMA GLOBAL	<u>EQ 20MG BASE</u>	<u>A090745 001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090745 002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090745 003</u>	Dec 11, 2013
<u>AB</u>		TEVA PHARMS USA	<u>EQ 20MG BASE</u>	<u>A090783 001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090783 002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090783 003</u>	Dec 11, 2013
<u>AB</u>		TORRENT PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A090774 001</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090774 002</u>	Dec 11, 2013
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090774 003</u>	Dec 11, 2013
<u>AB</u>		ZYDUS HLTHCARE	<u>EQ 20MG BASE</u>	<u>A090739 001</u>	Jan 08, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090739 002</u>	Jan 08, 2014
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090739 003</u>	Jan 08, 2014
<u>AB</u>		ZYDUS PHARMS USA INC	<u>EQ 20MG BASE</u>	<u>A090728 001</u>	Jan 08, 2014
<u>AB</u>			<u>EQ 30MG BASE</u>	<u>A090728 002</u>	Jan 08, 2014
<u>AB</u>			<u>EQ 60MG BASE</u>	<u>A090728 003</u>	Jan 08, 2014
		LUPIN LTD	<u>EQ 40MG BASE</u>	A090694 003	Dec 11, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-149 (of 436)

DUTASTERIDE

CAPSULE; ORAL

AVODART

<u>AB</u>	+!	GLAXOSMITHKLINE	<u>0.5MG</u>	<u>N021319 001</u>	Nov 20, 2001
<u>DUTASTERIDE</u>					
<u>AB</u>		ACTAVIS LABS FL INC	<u>0.5MG</u>	<u>A202808 001</u>	Nov 20, 2015
<u>AB</u>		AMNEAL PHARMS	<u>0.5MG</u>	<u>A203118 001</u>	Nov 20, 2015
<u>AB</u>		APOTEX INC	<u>0.5MG</u>	<u>A204292 001</u>	Nov 24, 2015
<u>AB</u>		ASCENT PHARMS INC	<u>0.5MG</u>	<u>A206574 001</u>	Oct 21, 2016
<u>AB</u>		AUROLIFE PHARMA LLC	<u>0.5MG</u>	<u>A202660 001</u>	Nov 20, 2015
<u>AB</u>		BARR	<u>0.5MG</u>	<u>A090095 001</u>	Dec 21, 2010
<u>AB</u>		BIONPHARMA INC	<u>0.5MG</u>	<u>A200899 001</u>	Nov 20, 2015
<u>AB</u>		BRECKENRIDGE PHARM	<u>0.5MG</u>	<u>A204705 001</u>	Nov 20, 2015
<u>AB</u>		HAUTP PHARMA	<u>0.5MG</u>	<u>A207935 001</u>	Oct 13, 2017
<u>AB</u>		HUMANWELL PURACAP	<u>0.5MG</u>	<u>A209909 001</u>	Nov 21, 2017
<u>AB</u>		INTERGEL PHARMS INC	<u>0.5MG</u>	<u>A206373 001</u>	Mar 17, 2016
<u>AB</u>		MARKSANS PHARMA	<u>0.5MG</u>	<u>A204376 001</u>	Apr 07, 2017
<u>AB</u>		MYLAN PHARMS INC	<u>0.5MG</u>	<u>A203241 001</u>	Jun 14, 2016
<u>AB</u>		RISING PHARMS INC	<u>0.5MG</u>	<u>A202530 001</u>	Nov 20, 2015
<u>AB</u>		STRIDES PHARMA	<u>0.5MG</u>	<u>A204262 001</u>	Nov 20, 2015
<u>AB</u>		VINTAGE PHARMS LLC	<u>0.5MG</u>	<u>A202421 001</u>	Nov 20, 2015
<u>AB</u>		WEST-WARD PHARMS	<u>0.5MG</u>	<u>A202204 001</u>	Nov 23, 2015
		INT			
<u>AB</u>		ZYDUS PHARMS USA	<u>0.5MG</u>	<u>A204373 001</u>	Oct 04, 2017
		INC			

DUTASTERIDE; TAMSULOSIN HYDROCHLORIDE

CAPSULE; ORAL

DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>0.5MG;0.4MG</u>	<u>A202975 001</u>	Nov 20, 2015
<u>AB</u>		ANCHEN PHARMS	<u>0.5MG;0.4MG</u>	<u>A202509 001</u>	Feb 26, 2014
<u>JALYN</u>					
<u>AB</u>	+!	GLAXOSMITHKLINE	<u>0.5MG;0.4MG</u>	<u>N022460 001</u>	Jun 14, 2010

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

+! WYETH PHARMS INC 0.125%

N011963 001

ECONAZOLE NITRATE

AEROSOL, FOAM; TOPICAL

ECOZA

+! CHEMO RESEARCH SL 1%

N205175 001 Oct 24, 2013

CREAM; TOPICAL

ECONAZOLE NITRATE

<u>AB</u>		FOUGERA PHARMS	<u>1%</u>	<u>A076075 001</u>	Nov 26, 2002
<u>AB</u>	!	PERRIGO NEW YORK	<u>1%</u>	<u>A076479 001</u>	Jun 23, 2004
<u>AB</u>		TARO	<u>1%</u>	<u>A076005 001</u>	Nov 26, 2002
<u>AB</u>		TELIGENT PHARMA INC	<u>1%</u>	<u>A076574 001</u>	Dec 17, 2004

SPECTAZOLE

<u>AB</u>	+	ALVOGEN MALTA	<u>1%</u>	<u>N018751 001</u>	Dec 23, 1982
-----------	---	---------------	-----------	--------------------	--------------

EDARAVONE

SOLUTION; IV (INFUSION)

RADICAVA

+! MITSUBISHI TANABE 30MG/100ML (0.3MG/ML)

N209176 001 May 05, 2017

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

+! MEDICIS 200MG/ML

N008922 001

EDOXABAN TOSYLYLATE

TABLET; ORAL

SAVAYSIA

+ DAIICHI SANKYO INC EQ 15MG BASE  
+ EQ 30MG BASE  
+! EQ 60MG BASE

N206316 001 Jan 08, 2015  
N206316 002 Jan 08, 2015  
N206316 003 Jan 08, 2015

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON

! MYLAN INSTITUTIONAL 10MG/ML

A088873 001 Aug 06, 1985

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-150 (of 436)

EFAVIRENZ

CAPSULE;ORAL

EFAVIRENZ

<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A078064 001</u> Dec 15, 2017
-----------	----------------------	-------------	---------------------------------

<u>AB</u>		<u>200MG</u>	<u>A078064 003</u> Dec 15, 2017
-----------	--	--------------	---------------------------------

SUSTIVIA

<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>50MG</u>	<u>N020972 001</u> Sep 17, 1998
-----------	------------------------	-------------	---------------------------------

<u>AB</u>	+! EFAVIRENZ	<u>200MG</u>	<u>N020972 003</u> Sep 17, 1998
-----------	--------------	--------------	---------------------------------

EFAVIRENZ

<u>AB</u>	AUROBINDO PHARMA LTD	100MG	A078064 002 Dec 15, 2017
-----------	----------------------	-------	--------------------------

TABLET;ORAL

EFAVIRENZ

<u>AB</u>	MYLAN PHARMS INC	<u>600MG</u>	<u>A091471 001</u> Feb 17, 2016
-----------	------------------	--------------	---------------------------------

SUSTIVIA

<u>AB</u>	+! BRISTOL MYERS SQUIBB	<u>600MG</u>	<u>N021360 002</u> Feb 01, 2002
-----------	-------------------------	--------------	---------------------------------

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

ATRIPLA

+!	GILEAD	600MG;200MG;300MG	N021937 001 Jul 12, 2006
----	--------	-------------------	--------------------------

EFINACONAZOLE

SOLUTION;TOPICAL

JUBLIA

+!	DOW PHARM	10%	N203567 001 Jun 06, 2014
----	-----------	-----	--------------------------

EFLORNITHINE HYDROCHLORIDE

CREAM;TOPICAL

VANIQA

+!	SKINMEDICA	13.9%	N021145 001 Jul 27, 2000
----	------------	-------	--------------------------

ELBASVIR; GRAZOPREVIR

TABLET;ORAL

ZEPATIER

+!	MERCK SHARP DOHME	50MG;100MG	N208261 001 Jan 28, 2016
----	-------------------	------------	--------------------------

ELETRIPTAN HYDROBROMIDE

TABLET;ORAL

ELETRIPTAN HYDROBROMIDE

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A205186 001</u> Aug 29, 2017
-----------	-------------------	---------------------	---------------------------------

<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205186 002</u> Aug 29, 2017
-----------	--	---------------------	---------------------------------

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A205152 001</u> Aug 11, 2017
-----------	------------------	---------------------	---------------------------------

<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205152 002</u> Aug 11, 2017
-----------	--	---------------------	---------------------------------

<u>AB</u>	TEVA PHARMS USA	<u>EQ 20MG BASE</u>	<u>A202040 001</u> Jun 27, 2017
-----------	-----------------	---------------------	---------------------------------

<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A202040 002</u> Jun 27, 2017
-----------	--	---------------------	---------------------------------

<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 20MG BASE</u>	<u>A206409 001</u> Jun 16, 2017
-----------	----------------------	---------------------	---------------------------------

RELPAX

<u>AB</u>	+ PFIZER IRELAND	<u>EQ 20MG BASE</u>	<u>N021016 001</u> Dec 26, 2002
-----------	------------------	---------------------	---------------------------------

<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N021016 002</u> Dec 26, 2002
-----------	----	---------------------	---------------------------------

ELIGLUSTAT TARTRATE

CAPSULE;ORAL

CERDELGA

+!	GENZYME CORP	EQ 84MG BASE	N205494 001 Aug 19, 2014
----	--------------	--------------	--------------------------

ELTROMBOPAG OLAMINE

FOR SUSPENSION;ORAL

PROMACTA

+!	NOVARTIS PHARMS CORP	EQ 25MG ACID/PACKET	N207027 001 Aug 24, 2015
----	----------------------	---------------------	--------------------------

TABLET;ORAL

PROMACTA

+	NOVARTIS PHARMS CORP	EQ 12.5MG ACID	N022291 004 Oct 20, 2011
---	----------------------	----------------	--------------------------

+		EQ 25MG ACID	N022291 001 Nov 20, 2008
---	--	--------------	--------------------------

+		EQ 50MG ACID	N022291 002 Nov 20, 2008
---	--	--------------	--------------------------

+		EQ 75MG ACID	N022291 003 Sep 08, 2009
---	--	--------------	--------------------------

+		EQ 100MG ACID	N022291 005 Nov 16, 2012
---	--	---------------	--------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-151 (of 436)

ELUXADOLINE

TABLET;ORAL

VIBERZI

+ ALLERGAN HOLDINGS	75MG	N206940 001 May 27, 2015
+!	100MG	N206940 002 May 27, 2015

EMEDASTINE DIFUMARATE

SOLUTION/DROPS;OPHTHALMIC

EMADINE

+! NOVARTIS PHARMS CORP	0.05%	N020706 001 Dec 29, 1997
-------------------------	-------	--------------------------

EMPAGLIFLOZIN

TABLET;ORAL

JARDIANCE

+ BOEHRINGER INGELHEIM	10MG	N204629 001 Aug 01, 2014
+!	25MG	N204629 002 Aug 01, 2014

EMPAGLIFLOZIN; LINAGLITPTIN

TABLET;ORAL

GLYXAMBI

+ BOEHRINGER INGELHEIM	10MG;5MG	N206073 001 Jan 30, 2015
+!	25MG;5MG	N206073 002 Jan 30, 2015

EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET;ORAL

SYNJARDY

+ BOEHRINGER INGELHEIM	5MG;500MG	N206111 001 Aug 26, 2015
+	5MG;1GM	N206111 002 Aug 26, 2015
+	12.5MG;500MG	N206111 003 Aug 26, 2015
+!	12.5MG;1GM	N206111 004 Aug 26, 2015

TABLET, EXTENDED RELEASE;ORAL

SYNJARDY XR

+ BOEHRINGER INGELHEIM	5MG;1GM	N208658 001 Dec 09, 2016
+	10MG;1GM	N208658 002 Dec 09, 2016
+	12.5MG;1GM	N208658 003 Dec 09, 2016
+!	25MG;1GM	N208658 004 Dec 09, 2016

EMTRICITABINE

CAPSULE;ORAL

EMTRIVA

+! GILEAD	200MG	N021500 001 Jul 02, 2003
-----------	-------	--------------------------

SOLUTION;ORAL

EMTRIVA

+! GILEAD	10MG/ML	N021896 001 Sep 28, 2005
-----------	---------	--------------------------

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

ODEFSEY

+! GILEAD SCIENCES INC	200MG;EQ 25MG BASE;EQ 25MG BASE	N208351 001 Mar 01, 2016
------------------------	---------------------------------	--------------------------

EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

COMPLERA

+! GILEAD SCIENCES INC	200MG;EQ 25MG BASE;300MG	N202123 001 Aug 10, 2011
------------------------	--------------------------	--------------------------

EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE

TABLET;ORAL

DESCOVY

+! GILEAD SCIENCES INC	200MG;EQ 25MG BASE	N208215 001 Apr 04, 2016
------------------------	--------------------	--------------------------

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET;ORAL

EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE

<b>AB</b>	TEVA PHARMS USA	<b>200MG;300MG</b>	<b>A090894 001</b> Jun 08, 2017
<u>TRUVADA</u>			
<b>AB</b>	+! GILEAD	<b>200MG;300MG</b>	<b>N021752 001</b> Aug 02, 2004
	+	100MG;150MG	N021752 002 Mar 10, 2016
	+	133MG;200MG	N021752 003 Mar 10, 2016
	+	167MG;250MG	N021752 004 Mar 10, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-152 (of 436)

ENALAPRIL MALEATE

FOR SOLUTION;ORAL

EPANED KIT

+! SILVERGATE PHARMS 1MG/ML

N204308 001 Aug 13, 2013

SOLUTION;ORAL

EPANED

+! SILVERGATE PHARMS 1MG/ML

N208686 001 Sep 20, 2016

TABLET;ORAL

ENALAPRIL MALEATE

<u>AB</u>	APOTEX	<u>2.5MG</u>	<u>A075178 002</u>	Mar 23, 2001
<u>AB</u>		<u>5MG</u>	<u>A075178 001</u>	Mar 23, 2001
<u>AB</u>		<u>10MG</u>	<u>A075178 003</u>	Mar 23, 2001
<u>AB</u>		<u>20MG</u>	<u>A075178 004</u>	Mar 23, 2001
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A075480 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075480 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075480 003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075480 004</u>	Aug 22, 2000
<u>AB</u>	SANDOZ INC	<u>2.5MG</u>	<u>A075496 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075496 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075459 001</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075459 002</u>	Aug 22, 2000
<u>AB</u>	TARO	<u>2.5MG</u>	<u>A075657 001</u>	Jan 23, 2001
<u>AB</u>		<u>5MG</u>	<u>A075657 002</u>	Jan 23, 2001
<u>AB</u>		<u>10MG</u>	<u>A075657 003</u>	Jan 23, 2001
<u>AB</u>		<u>20MG</u>	<u>A075657 004</u>	Jan 23, 2001
<u>AB</u>	TEVA	<u>2.5MG</u>	<u>A075479 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075479 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075479 003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075479 004</u>	Aug 22, 2000
<u>AB</u>	WOCKHARDT LTD	<u>2.5MG</u>	<u>A075483 001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>A075483 002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>A075483 003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>A075483 004</u>	Aug 22, 2000
<u>VASOTEC</u>				
<u>AB</u>	+ VALEANT PHARMS NORTH	<u>2.5MG</u>	<u>N018998 005</u>	Jul 26, 1988
<u>AB</u>	+ !	<u>5MG</u>	<u>N018998 001</u>	Dec 24, 1985
<u>AB</u>	+ !	<u>10MG</u>	<u>N018998 002</u>	Dec 24, 1985
<u>AB</u>	+ !	<u>20MG</u>	<u>N018998 003</u>	Dec 24, 1985

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX INC	<u>5MG;12.5MG</u>	<u>A076486 001</u>	Oct 27, 2004
<u>AB</u>		<u>10MG;25MG</u>	<u>A076486 002</u>	Oct 27, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG;12.5MG</u>	<u>A075909 001</u>	Oct 15, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075909 002</u>	Oct 15, 2001
<u>AB</u>	G AND W LABS INC	<u>5MG;12.5MG</u>	<u>A075727 001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075727 002</u>	Sep 18, 2001
<u>AB</u>	MYLAN	<u>5MG;12.5MG</u>	<u>A075624 001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075624 002</u>	Sep 18, 2001
<u>AB</u>	TARO PHARM IND	<u>5MG;12.5MG</u>	<u>A075788 001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG;25MG</u>	<u>A075788 002</u>	Sep 18, 2001
<u>VASERETIC</u>				
<u>AB</u>	+ VALEANT INTL	<u>5MG;12.5MG</u>	<u>N019221 003</u>	Jul 12, 1995
<u>AB</u>	+ !	<u>10MG;25MG</u>	<u>N019221 001</u>	Oct 31, 1986

ENALAPRILAT

INJECTABLE;INJECTION

ENALAPRILAT

<u>AP</u>	! ATHENEX INC	<u>1.25MG/ML</u>	<u>A075634 001</u>	Aug 22, 2000
<u>AP</u>	HIKMA FARMACEUTICA	<u>1.25MG/ML</u>	<u>A078687 001</u>	Dec 23, 2008
<u>AP</u>	! HOSPIRA	<u>1.25MG/ML</u>	<u>A075458 001</u>	Aug 22, 2000
<u>AP</u>	TEVA PHARMS USA	<u>1.25MG/ML</u>	<u>A075578 001</u>	Aug 22, 2000

ENASIDENIB MESYLATE

TABLET;ORAL

IDHIFA

+! CELGENE CORP EQ 50MG BASE  
+! EQ 100MG BASE

N209606 001 Aug 01, 2017  
N209606 002 Aug 01, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-153 (of 436)

ENFUVIRTIDE

INJECTABLE; SUBCUTANEOUS

FUZEON

+! ROCHE

90MG/VIAL

N021481 001 Mar 13, 2003

ENOXAPARIN SODIUM

INJECTABLE; INTRAVENOUS, SUBCUTANEOUS

ENOXAPARIN SODIUM

AB SANDOZ INC 300MG/3ML (100MG/ML)

A078660 001 Nov 28, 2011

LOVENOX

AB + SANOFI AVENTIS US

300MG/3ML (100MG/ML)

N020164 009 Jan 23, 2003

INJECTABLE; SUBCUTANEOUS

ENOXAPARIN SODIUM (PRESERVATIVE FREE)

AP AMPHASTAR PHARM 30MG/0.3ML (100MG/ML)

A076684 001 Sep 19, 2011

AP 40MG/0.4ML (100MG/ML)

A076684 002 Sep 19, 2011

AP 60MG/0.6ML (100MG/ML)

A076684 003 Sep 19, 2011

AP 80MG/0.8ML (100MG/ML)

A076684 004 Sep 19, 2011

AP 100MG/ML (100MG/ML)

A076684 005 Sep 19, 2011

AP 120MG/0.8ML (150MG/ML)

A076684 006 Sep 19, 2011

AP 150MG/ML (150MG/ML)

A076684 007 Sep 19, 2011

AP SANDOZ 30MG/0.3ML (100MG/ML)

A077857 002 Jul 23, 2010

AP 40MG/0.4ML (100MG/ML)

A077857 003 Jul 23, 2010

AP 60MG/0.6ML (100MG/ML)

A077857 004 Jul 23, 2010

AP 80MG/0.8ML (100MG/ML)

A077857 005 Jul 23, 2010

AP 100MG/ML (100MG/ML)

A077857 006 Jul 23, 2010

AP 120MG/0.8ML (150MG/ML)

A077857 007 Jul 23, 2010

AP 150MG/ML (150MG/ML)

A076726 001 Jun 23, 2014

AP TEVA 30MG/0.3ML (100MG/ML)

A076726 002 Jun 23, 2014

AP 40MG/0.4ML (100MG/ML)

A076726 003 Jun 23, 2014

AP 60MG/0.6ML (100MG/ML)

A076726 004 Jun 23, 2014

AP 80MG/0.8ML (100MG/ML)

A076726 005 Jun 23, 2014

AP 100MG/ML (100MG/ML)

A076726 006 Jun 23, 2014

AP 120MG/0.8ML (150MG/ML)

A076726 007 Jun 23, 2014

AP 150MG/ML (150MG/ML)

A076726 008 Jun 23, 2014

LOVENOX (PRESERVATIVE FREE)

AP + SANOFI AVENTIS US 30MG/0.3ML (100MG/ML)

N020164 001 Mar 29, 1993

AP + 40MG/0.4ML (100MG/ML)

N020164 002 Jan 30, 1998

AP + 60MG/0.6ML (100MG/ML)

N020164 003 Mar 27, 1998

AP + 80MG/0.8ML (100MG/ML)

N020164 004 Mar 27, 1998

AP +! 100MG/ML (100MG/ML)

N020164 005 Mar 27, 1998

AP + 120MG/0.8ML (150MG/ML)

N020164 007 Jun 02, 2000

AP + 150MG/ML (150MG/ML)

N020164 008 Jun 02, 2000

ENTACAPONE

TABLET; ORAL

COMTAN

AB +! ORION PHARMA 200MG

N020796 001 Oct 19, 1999

ENTACAPONE

AB AJANTA PHARMA LTD 200MG

A205792 001 Aug 31, 2017

AB AUROBINDO PHARMA LTD 200MG

A203437 001 Jun 19, 2015

AB MACLEODS PHARMS LTD 200MG

A207210 001 Jun 05, 2017

AB SUN PHARMA GLOBAL 200MG

A090690 001 Jul 16, 2012

AB WOCKHARDT LTD 200MG

A078941 001 Aug 16, 2012

ENTECAVIR

SOLUTION; ORAL

BARACLUDE

+! BRISTOL MYERS SQUIBB

0.05MG/ML

N021798 001 Mar 29, 2005

TABLET; ORAL

BARACLUDE

+! BRISTOL MYERS SQUIBB

0.5MG

N021797 001 Mar 29, 2005

+!

1MG

N021797 002 Mar 29, 2005

ENTECAVIR

AB ACCORD HLTHCARE 0.5MG

A205824 001 Aug 25, 2017

AB AMNEAL PHARMS 1MG

A205824 002 Aug 25, 2017

AB AUROBINDO PHARMA LTD 0.5MG

A206652 001 Nov 12, 2015

AB AUROBINDO PHARMA LTD 1MG

A206652 002 Nov 12, 2015

AB CIPLA LTD 0.5MG

A206217 001 Aug 26, 2015

AB CIPLA LTD 1MG

A206217 002 Aug 26, 2015

AB CIPLA LTD 0.5MG

A206872 001 Dec 06, 2016

AB CIPLA LTD 1MG

A206872 002 Dec 06, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-154 (of 436)

ENTECAVIR

TABLET;ORAL

ENTECAVIR

<u>AB</u>	HETERO LABS LTD V	<u>0.5MG</u>	<u>A205740 001</u>	Aug 21, 2015
<u>AB</u>		<u>1MG</u>	<u>A205740 002</u>	Aug 21, 2015
<u>AB</u>	PAR PHARM INC	<u>0.5MG</u>	<u>A206294 001</u>	Nov 23, 2016
<u>AB</u>		<u>1MG</u>	<u>A206294 002</u>	Nov 23, 2016
<u>AB</u>	PRINSTON INC	<u>0.5MG</u>	<u>A208782 001</u>	Oct 10, 2017
<u>AB</u>		<u>1MG</u>	<u>A208782 002</u>	Oct 10, 2017
<u>AB</u>	SANDOZ INC	<u>0.5MG</u>	<u>A206672 001</u>	May 11, 2017
<u>AB</u>		<u>1MG</u>	<u>A206672 002</u>	May 11, 2017
<u>AB</u>	TEVA PHARMS USA	<u>0.5MG</u>	<u>A202122 001</u>	Aug 26, 2014
<u>AB</u>		<u>1MG</u>	<u>A202122 002</u>	Aug 26, 2014
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.5MG</u>	<u>A206745 001</u>	Jun 23, 2017
<u>AB</u>		<u>1MG</u>	<u>A206745 002</u>	Jun 23, 2017

ENZALUTAMIDE

CAPSULE;ORAL

XTANDI

+! ASTELLAS

40MG

N203415 001 Aug 31, 2012

EPHEDRINE SULFATE

SOLUTION;INTRAVENOUS

AKOVAZ

<u>AP</u>	+! FLAMEL IRELAND LTD	<u>50MG/ML (50MG/ML)</u>	<u>N208289 001</u>	Apr 29, 2016
<u>AP</u>	<u>EPHEDRINE SULFATE</u>			
<u>AP</u>	AKORN INC	<u>50MG/ML (50MG/ML)</u>	<u>N208609 001</u>	Mar 01, 2017
<u>AP</u>	SANDOZ INC	<u>50MG/ML (50MG/ML)</u>	<u>A209784 001</u>	Aug 23, 2017
	SOLUTION;IV (INFUSION) CORPHEDRA			
	PAR STERILE PRODUCTS	50MG/ML (50MG/ML)	N208943 001	Jan 27, 2017

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ELESTAT

<u>AT</u>	+! ALLERGAN	<u>0.05%</u>	<u>N0211565 001</u>	Oct 16, 2003
	<u>EPINASTINE HYDROCHLORIDE</u>			
<u>AT</u>	AKORN	<u>0.05%</u>	<u>A204055 001</u>	May 05, 2017
<u>AT</u>	APOTEX	<u>0.05%</u>	<u>A090919 001</u>	Oct 31, 2011
<u>AT</u>	BRECKENRIDGE PHARM	<u>0.05%</u>	<u>A090870 001</u>	Mar 14, 2011
<u>AT</u>	SANDOZ INC	<u>0.05%</u>	<u>A203384 001</u>	Dec 07, 2016
<u>AT</u>	SOMERSET THERAPS LLC	<u>0.05%</u>	<u>A090951 001</u>	Oct 31, 2011
<u>AT</u>	SUN PHARM IND	<u>0.05%</u>	<u>A0911626 001</u>	Oct 31, 2011

EPINEPHRINE

INJECTABLE;INTRAMUSCULAR, SUBCUTANEOUS

ADRENACCLICK

BX	+! IMPAX LABS INC	EQ 0.15MG/DELIVERY	N020800 003	Nov 25, 2009
BX	+!	EQ 0.3MG/DELIVERY	N020800 004	Nov 25, 2009
	EPIPEN			
BX	+! MYLAN SPECIALITY LP	0.3MG/DELIVERY	N019430 001	Dec 22, 1987
	EPIPEN JR.			
BX	+! MYLAN SPECIALITY LP	0.15MG/DELIVERY	N019430 002	Dec 22, 1987
	SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS			
	AUVI-Q			
BX	+! KALEO INC	EQ 0.15MG/DELIVERY	N201739 002	Aug 10, 2012
BX	+	EQ 0.3MG/DELIVERY	N201739 001	Aug 10, 2012
	ADRENALIN			
	+! PAR STERILE PRODUCTS	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N204200 001	Dec 07, 2012
	+	EQ 30MG BASE/30ML (EQ 1MG BASE/ML)	N204640 001	Dec 18, 2013
	AUVI-Q			
	+ KALEO INC	EQ 0.1MG/DELIVERY	N201739 003	Nov 17, 2017
	SYMJEPI			
	+! ADAMIS PHARMS CORP	0.3MG/0.3ML (0.3MG/0.3ML)	N207534 001	Jun 15, 2017
	SOLUTION;IV (INFUSION), INTRAOCULAR, INTRAMUSCULAR, SUBCUTANEOUS			
	EPINEPHRINE			
	BELCHER PHARMS LLC	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N205029 001	Jul 29, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-155 (of 436)

EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIGNOSPAN FORTE

! DEPROCO

EQ 0.02MG BASE/ML; 2%

A088389 001 Jan 22, 1985

LIGNOSPAN STANDARD

! DEPROCO

EQ 0.01MG BASE/ML; 2%

A088390 001 Jan 22, 1985

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

**CITANESE FORTE DENTAL**

**AP +! DENTSPLY PHARM 0.005MG/ML; 4%**

**N021383 001**

**PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE**

**AP SEPTODONT INC 0.005MG/ML; 4%**

**A078959 001 Aug 30, 2011**

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

**LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE**

**AP EASTMAN KODAK 0.01MG/ML; 2%**

**A040057 002 Feb 26, 1993**

**AP 0.02MG/ML; 2%**

**A040057 001 Feb 26, 1993**

**AP HOSPIRA 0.005MG/ML; 0.5%**

**A089635 001 Jun 21, 1988**

**AP 0.005MG/ML; 1.5%**

**A088571 001 Sep 13, 1985**

**AP 0.005MG/ML; 1.5%**

**A089645 001 Jun 21, 1988**

**AP 0.005MG/ML; 2%**

**A089651 001 Jun 21, 1988**

**AP 0.01MG/ML; 1%**

**A089644 001 Jun 21, 1988**

**AP 0.01MG/ML; 2%**

**A078772 001 May 12, 2008**

**AP 0.01MG/ML; 2%**

**A089646 001 Jun 21, 1988**

**AP 0.02MG/ML; 2%**

**A078772 002 May 12, 2008**

**OCTOCOCAINE**

**AP ! SEPTODONT 0.01MG/ML; 2%**

**A084048 001**

**AP ! 0.02MG/ML; 2%**

**A084048 002**

**XYLOCAINE W/ EPINEPHRINE**

**AP +! FRESENIUS KABI USA 0.005MG/ML; 0.5%**

**N006488 012**

**AP +! 0.005MG/ML; 1.5%**

**N006488 017**

**AP +! 0.005MG/ML; 2%**

**N006488 019 Nov 13, 1986**

**AP +! 0.01MG/ML; 1%**

**N006488 004**

**AP +! 0.02MG/ML; 2%**

**N006488 005**

+!

0.005MG/ML; 1% N006488 018 Nov 13, 1986

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

**ELLENCE**

**AP +! PFIZER INC 200MG/100ML (2MG/ML)**

**N050778 001 Sep 15, 1999**

**AP + 50MG/25ML (2MG/ML)**

**N050778 002 Sep 15, 1999**

**EPIRUBICIN HYDROCHLORIDE**

**AP ACTAVIS TOTOWA 10MG/5ML (2MG/ML)**

**A065445 001 Sep 18, 2008**

**AP 50MG/25ML (2MG/ML)**

**A065445 002 Sep 18, 2008**

**AP 200MG/100ML (2MG/ML)**

**A065445 003 Sep 18, 2008**

**AP AKORN INC 50MG/25ML (2MG/ML)**

**A090163 001 Jun 24, 2009**

**AP CIPLA LTD 50MG/25ML (2MG/ML)**

**A065361 001 Oct 22, 2007**

**AP FRESENIUS KABI 200MG/100ML (2MG/ML)**

**A065361 002 Oct 22, 2007**

**AP ONCOL 200MG/100ML (2MG/ML)**

**A065411 001 Aug 20, 2007**

**AP 50MG/25ML (2MG/ML)**

**A065411 002 Aug 20, 2007**

**AP FRESENIUS KABI USA 10MG/5ML (2MG/ML)**

**A065408 001 Oct 15, 2007**

**AP 50MG/25ML (2MG/ML)**

**A065408 002 Oct 15, 2007**

**AP 150MG/75ML (2MG/ML)**

**A065408 003 Oct 15, 2007**

**AP 200MG/100ML (2MG/ML)**

**A065408 004 Oct 15, 2007**

**AP HISUN PHARM 50MG/25ML (2MG/ML)**

**A090075 001 Mar 25, 2010**

**AP HANGZHOU 200MG/100ML (2MG/ML)**

**A090075 002 Mar 25, 2010**

**AP HOSPIRA 10MG/5ML (2MG/ML)**

**A065343 001 Apr 19, 2007**

**AP 150MG/75ML (2MG/ML)**

**A065343 003 Apr 19, 2007**

**AP 200MG/100ML (2MG/ML)**

**A065343 004 Apr 19, 2007**

**AP IMPAX LABS INC 50MG/25ML (2MG/ML)**

**A065331 001 Aug 09, 2007**

**AP 200MG/100ML (2MG/ML)**

**A065331 002 Aug 09, 2007**

**AP MYLAN LABS LTD 50MG/25ML (2MG/ML)**

**A091599 001 Mar 12, 2012**

**AP 200MG/100ML (2MG/ML)**

**A091599 002 Mar 12, 2012**

**AP WEST-WARD PHARMS INT 50MG/25ML (2MG/ML)**

**A065289 001 Jun 27, 2007**

**AP 200MG/100ML (2MG/ML)**

**A065289 002 Jun 27, 2007**

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-156 (of 436)

**EPLERENONE**

TABLET;ORAL

**EPLERENONE**

<b>AB</b>	ACCORD HLTHCARE	<u>25MG</u>	<b>A206922 001</b>	Jul 13, 2017
<b>AB</b>		<u>50MG</u>	<b>A206922 002</b>	Jul 13, 2017
<b>AB</b>	APOTEX	<u>25MG</u>	<b>A078482 001</b>	Jul 30, 2008
<b>AB</b>		<u>50MG</u>	<b>A078482 002</b>	Jul 30, 2008
<b>AB</b>	MYLAN PHARMS INC	<u>25MG</u>	<b>A203896 001</b>	Feb 02, 2017
<b>AB</b>		<u>50MG</u>	<b>A203896 002</b>	Feb 02, 2017
<b>AB</b>	SANDOZ	<u>25MG</u>	<b>A078510 001</b>	Aug 01, 2008
<b>AB</b>		<u>50MG</u>	<b>A078510 002</b>	Aug 01, 2008

**INSPRA**

<b>AB</b>	+	GD SEARLE LLC	<u>25MG</u>	<b>N021437 001</b>	Sep 27, 2002
<b>AB</b>	+	!	<u>50MG</u>	<b>N021437 002</b>	Sep 27, 2002

**EPOPROSTENOL SODIUM**

INJECTABLE; INJECTION

**EPOPROSTENOL SODIUM**

<b>AP</b>	TEVA PHARMS USA	<u>EQ 0.5MG BASE/VIAL</u>	<b>A078396 001</b>	Apr 23, 2008
<b>AP</b>		<u>EQ 1.5MG BASE/VIAL</u>	<b>A078396 002</b>	Apr 23, 2008
<b>AP</b>	+	!	GLAXOSMITHKLINE LLC	<u>EQ 0.5MG BASE/VIAL</u>
<b>AP</b>	+	!		<u>EQ 1.5MG BASE/VIAL</u>
	VELETRI			
	+ ACTELION PHARMS LTD	EQ 0.5MG BASE/VIAL	N022260 002	Jun 28, 2012
	+	EQ 1.5MG BASE/VIAL	N022260 001	Jun 27, 2008

**EPROSARTAN MESYLATE**

TABLET;ORAL

**EPROSARTAN MESYLATE**

<b>AB</b>	MYLAN PHARMS INC	<u>EQ 400MG BASE</u>	<b>A202012 001</b>	Nov 16, 2011
<b>AB</b>		<u>EQ 600MG BASE</u>	<b>A202012 002</b>	Nov 16, 2011
<b>AB</b>	TEVETEN			
<b>AB</b>	+	ABBVIE	<u>EQ 400MG BASE</u>	<b>N020738 005</b>
<b>AB</b>	+	!	<u>EQ 600MG BASE</u>	<b>N020738 006</b>

**EPTIFIBATIDE**

INJECTABLE; INJECTION

**EPTIFIBATIDE**

<b>AP</b>	ACCORD HLTHCARE	<u>2MG/ML</u>	<b>A205557 001</b>	Nov 06, 2017
<b>AP</b>		<u>75MG/100ML</u>	<b>A205557 002</b>	Nov 06, 2017
<b>AP</b>	AKORN	<u>2MG/ML</u>	<b>A204589 001</b>	Apr 18, 2017
<b>AP</b>		<u>75MG/100ML</u>	<b>A204589 002</b>	Apr 18, 2017
<b>AP</b>	AMNEAL PHARMS	<u>2MG/ML</u>	<b>A205581 001</b>	Dec 08, 2016
<b>AP</b>		<u>75MG/100ML</u>	<b>A205581 002</b>	Dec 08, 2016
<b>AP</b>	AUROBINDO PHARMA LTD	<u>2MG/ML</u>	<b>A206127 001</b>	Dec 08, 2015
<b>AP</b>		<u>75MG/100ML</u>	<b>A206127 002</b>	Dec 08, 2015
<b>AP</b>	TEVA PHARMS USA	<u>2MG/ML</u>	<b>A090854 001</b>	Jun 12, 2015
<b>AP</b>	INTEGRILIN			
<b>AP</b>	+	!	SCHERING	<u>2MG/ML</u>
<b>AP</b>	+	!		<u>75MG/100ML</u>

**ERGOCALCIFEROL**

CAPSULE;ORAL

**DRISDOL**

<b>AA</b>	+	!	US PHARM HOLDINGS	<u>50,000 IU</u>	<b>N003444 001</b>
<b>AA</b>	ERGOCALCIFEROL				
<b>AA</b>	ORIT LABS LLC		<u>50,000 IU</u>	<b>A040833 001</b>	May 20, 2009
<b>AA</b>	SIGMAPHARM LABS LLC		<u>50,000 IU</u>	<b>A091004 001</b>	Jul 14, 2010
<b>AA</b>	STRIDES PHARMA		<u>50,000 IU</u>	<b>A090455 001</b>	Aug 03, 2010
<b>AA</b>	SUN PHARM INDs INC		<u>50,000 IU</u>	<b>A040865 001</b>	Dec 29, 2009

**VITAMIN D**

<b>AA</b>	BIONPHARMA INC	<u>50,000 IU</u>	<b>A080704 001</b>
-----------	----------------	------------------	--------------------

**ERGOLOID MESYLATES**

TABLET;ORAL

ERGOLOID MESYLATES

!	SUN PHARM INDUSTRIES	1MG	A081113 001	Oct 31, 1991
---	----------------------	-----	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-157 (of 436)

ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

! TERSERA THERAPS LLC 2MG

A087693 001 Feb 24, 1983

ERIBULIN MESYLATE

SOLUTION; INTRAVENOUS

HALAVEN

+! EISAI INC

1MG/2ML (0.5MG/ML)

N201532 001 Nov 15, 2010

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

TARCEVA

+! OSI PHARMS

EQ 25MG BASE

N021743 001 Nov 18, 2004

+

EQ 100MG BASE

N021743 002 Nov 18, 2004

! +!

EQ 150MG BASE

N021743 003 Nov 18, 2004

ERTAPENEM SODIUM

INJECTABLE; INTRAMUSCULAR, IV (INFUSION)

INVANZ

+! MERCK SHARP DOHME

EQ 1GM BASE/VIAL

N021337 001 Nov 21, 2001

ERTUGLIFLOZIN

TABLET; ORAL

STEGLATRO

+! MERCK SHARP DOHME

5MG

N209803 001 Dec 19, 2017

! +!

15MG

N209803 002 Dec 19, 2017

ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE

TABLET; ORAL

SEGLUROMET

+! MERCK SHARP DOHME

2.5MG; 500MG

N209806 001 Dec 19, 2017

+

2.5MG; 1GM

N209806 002 Dec 19, 2017

+

7.5MG; 500MG

N209806 003 Dec 19, 2017

! +!

7.5MG; 1GM

N209806 004 Dec 19, 2017

ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

STELUJAN

+! MERCK SHARP DOHME

5MG; EQ 100MG BASE

N209805 001 Dec 19, 2017

! +!

15MG; EQ 100MG BASE

N209805 002 Dec 19, 2017

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYC

AB +! MAYNE PHARMA 250MG

N050536 001

ERYTHROMYCIN

AB ARBOR PHARMS LLC 250MG

A062746 001 Dec 22, 1986

GEL; TOPICAL

ERYGEL

AT +! MYLAN PHARMS INC 2%

N050617 001 Oct 21, 1987

ERYTHROMYCIN

AT FOUGERA PHARMS 2%

A064184 001 Sep 30, 1997

AT PERRIGO CO 2%

A063211 001 Jan 29, 1993

AT TELIGENT PHARMA INC 2%

A208154 001 Jul 19, 2017

OINTMENT; OPHTHALMIC

ERYTHROMYCIN

AT AKORN 0.5%

A064030 001 Jul 18, 1996

AT BAUSCH AND LOMB 0.5%

A064067 001 Jul 29, 1994

AT ! PERRIGO CO 0.5%

A062447 001 Sep 26, 1983

TENNESSEE

SOLUTION; TOPICAL

ERYTHROMYCIN

AT ! PERRIGO NEW YORK 2%

A063038 001 Jan 11, 1991

AT TELIGENT PHARMA INC 2%

A208100 001 Nov 20, 2017

AT WOCKHARDT BIO AG 2%

A062825 001 Oct 23, 1987

SWAB; TOPICAL

ERYTHROMYCIN

AT AKORN 2%

A090215 001 May 12, 2010

AT ! PERRIGO CO 2%

A064126 001 Jul 03, 1996

TABLET; ORAL

ERYTHROMYCIN

ARBOR PHARMS LLC 250MG

A061621 001

!

500MG

A061621 002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-158 (of 436)

**ERYTHROMYCIN**

TABLET, COATED PARTICLES;ORAL

PCE			
+ ARBOR PHARMS LLC	333MG	N050611 001	Sep 09, 1986
+!	500MG	N050611 002	Aug 22, 1990

TABLET, DELAYED RELEASE;ORAL

ERY-TAB			
ARBOR PHARMS LLC	250MG	A062298 001	
	333MG	A062298 003	Mar 29, 1982
!	500MG	A062298 002	

**ERYTHROMYCIN ETHYLSUCCINATE**

GRANULE;ORAL

E.E.S.			
+ ARBOR PHARMS LLC	EQ 200MG BASE/5ML	N050207 001	
ERYPED			
+ ARBOR PHARMS LLC	EQ 200MG BASE/5ML	N050207 003	Mar 30, 1987
+!	EQ 400MG BASE/5ML	N050207 002	

TABLET;ORAL

E.E.S. 400

BX ! ARBOR PHARMS LLC	EQ 400MG BASE	A061905 002	Aug 12, 1982
ERYTHROMYCIN ETHYLSUCCINATE			
BX ! ARBOR PHARMS LLC	EQ 400MG BASE	A061904 001	

**ERYTHROMYCIN LACTOBIONATE**

INJECTABLE;INJECTION

**ERYTHROCIN**

<b>AP</b> HOSPIRA	<b>EQ 500MG BASE/VIAL</b>	<b>A062638 001</b>	Oct 31, 1986
<b>AP</b> +!	<b>EQ 500MG BASE/VIAL</b>	<b>N050609 001</b>	Sep 24, 1986
!	EQ 1GM BASE/VIAL	A062638 002	Oct 31, 1986

**ERYTHROMYCIN STEARATE**

TABLET;ORAL

ERYTHROCIN STEARATE

! ARBOR PHARMS LLC	EQ 250MG BASE	A060359 001	
--------------------	---------------	-------------	--

**ESCITALOPRAM OXALATE**

SOLUTION;ORAL

**ESCITALOPRAM OXALATE**

<b>AA</b> AMNEAL PHARMS	<b>EQ 5MG BASE/5ML</b>	<b>A202227 001</b>	Mar 14, 2012
<b>AA</b> ANTRIM PHARMS LLC	<b>EQ 5MG BASE/5ML</b>	<b>A203967 001</b>	May 26, 2015
<b>AA</b> AUROBINDO PHARMA LTD	<b>EQ 5MG BASE/5ML</b>	<b>A079062 001</b>	Apr 02, 2012
<b>AA</b> HETERO LABS LTD III	<b>EQ 5MG BASE/5ML</b>	<b>A202221 001</b>	Jun 12, 2012
<b>AA</b> MACLEODS PHARMS LTD	<b>EQ 5MG BASE/5ML</b>	<b>A202754 001</b>	Mar 31, 2016
<b>AA</b> SILARX PHARMS INC	<b>EQ 5MG BASE/5ML</b>	<b>A090477 001</b>	Jun 12, 2013
<b>AA</b> TARO	<b>EQ 5MG BASE/5ML</b>	<b>A079121 001</b>	May 03, 2012

**LEXAPRO**

<b>AA</b> +! FOREST LABS	<b>EQ 5MG BASE/5ML</b>	<b>N021365 001</b>	Nov 27, 2002
--------------------------	------------------------	--------------------	--------------

TABLET;ORAL

**ESCITALOPRAM OXALATE**

<b>AB</b> ACCORD HLTHCARE	<b>EQ 5MG BASE</b>	<b>A202389 001</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A202389 002</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A202389 003</b>	Sep 11, 2012
<b>AB</b> AMNEAL PHARMS	<b>EQ 5MG BASE</b>	<b>A205619 001</b>	May 17, 2017
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A205619 002</b>	May 17, 2017
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A205619 003</b>	May 17, 2017
<b>AB</b> APOTEX INC	<b>EQ 5MG BASE</b>	<b>A078777 001</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A078777 002</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A078777 003</b>	Sep 11, 2012
<b>AB</b> AUROBINDO PHARMA LTD	<b>EQ 5MG BASE</b>	<b>A090432 001</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A090432 002</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A090432 003</b>	Sep 11, 2012
<b>AB</b> HIKMA PHARMS	<b>EQ 5MG BASE</b>	<b>A078766 001</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A078766 002</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A078766 003</b>	Sep 11, 2012
<b>AB</b> INVAGEN PHARMS	<b>EQ 5MG BASE</b>	<b>A078604 001</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A078604 002</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A078604 003</b>	Sep 11, 2012
<b>AB</b> JUBILANT GENERICS	<b>EQ 5MG BASE</b>	<b>A202280 001</b>	Sep 12, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A202280 002</b>	Sep 12, 2012
<b>AB</b>	<b>EQ 20MG BASE</b>	<b>A202280 003</b>	Sep 12, 2012
<b>AB</b> LUPIN LTD	<b>EQ 5MG BASE</b>	<b>A078169 001</b>	Sep 11, 2012
<b>AB</b>	<b>EQ 10MG BASE</b>	<b>A078169 002</b>	Sep 11, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-159 (of 436)

ESCITALOPRAM OXALATE

TABLET;ORAL

ESCITALOPRAM OXALATE

<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078169 003</u>	Sep 11, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A202210 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202210 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A202210 003</u>	Sep 11, 2012
<u>AB</u>	PRINSTON INC	<u>EQ 5MG BASE</u>	<u>A078032 001</u>	Aug 28, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078032 002</u>	Aug 28, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078032 003</u>	Aug 28, 2015
<u>AB</u>	STI PHARMA LLC	<u>EQ 5MG BASE</u>	<u>A077512 001</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077512 002</u>	Sep 12, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077512 003</u>	Sep 12, 2012
<u>AB</u>	TEVA PHARMS USA	<u>EQ 5MG BASE</u>	<u>A076765 001</u>	Mar 14, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076765 002</u>	Mar 14, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076765 003</u>	Mar 14, 2012
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A090939 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A090939 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A090939 003</u>	Sep 11, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 5MG BASE</u>	<u>A077734 001</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077734 002</u>	Sep 11, 2012
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077734 003</u>	Sep 11, 2012
<u>LEXAPRO</u>				
<u>AB</u>	+ FOREST LABS	<u>EQ 5MG BASE</u>	<u>N021323 001</u>	Aug 14, 2002
<u>AB</u>	+	<u>EQ 10MG BASE</u>	<u>N021323 002</u>	Aug 14, 2002
<u>AB</u>	+!	<u>EQ 20MG BASE</u>	<u>N021323 003</u>	Aug 14, 2002

ESLICARBAZEPINE ACETATE

TABLET;ORAL

APTOIM

+	SUNOVION PHARMS INC	200MG	N022416 001	Nov 08, 2013
+		400MG	N022416 002	Nov 08, 2013
+		600MG	N022416 003	Nov 08, 2013
+		800MG	N022416 004	Nov 08, 2013

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+! BAXTER HLTHCARE	<u>10MG/ML</u>	<u>N019386 006</u>	Feb 25, 2003
<u>ESMOLOL HYDROCHLORIDE</u>				
<u>AP</u>	AUROBINDO PHARMA LTD	<u>10MG/ML</u>	<u>A205520 001</u>	Jul 23, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>A076573 001</u>	May 02, 2005
<u>AP</u>	LUITPOLD PHARMS INC	<u>10MG/ML</u>	<u>A201126 001</u>	Feb 20, 2015
<u>AP</u>	MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>A076474 001</u>	May 02, 2005
<u>AP</u>	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A076323 001</u>	Aug 10, 2004

BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER

++! BAXTER HLTHCARE	2GM/100ML	N019386 005	Jan 27, 2003
---------------------	-----------	-------------	--------------

BREVIBLOC IN PLASTIC CONTAINER

++! BAXTER HLTHCARE	1GM/100ML	N019386 004	Feb 16, 2001
---------------------	-----------	-------------	--------------

SOLUTION; INTRAVENOUS

ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER

++! HQ SPCLT PHARMA	2GM/100ML (20MG/ML)	N205703 002	Apr 07, 2016
---------------------	---------------------	-------------	--------------

ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER

++! HQ SPCLT PHARMA	2.5GM/250ML (10MG/ML)	N205703 001	Apr 07, 2016
---------------------	-----------------------	-------------	--------------

ESOMEPRAZOLE MAGNESEIUM

CAPSULE, DELAYED REL PELLETS;ORAL

ESOMEPRAZOLE MAGNESEIUM

<u>AB</u>	ALKEM LABS LTD	<u>EQ 20MG BASE</u>	<u>A208333 001</u>	Oct 20, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A208333 002</u>	Oct 20, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A205606 001</u>	Apr 21, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205606 002</u>	Apr 21, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A078279 001</u>	Sep 25, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078279 002</u>	Sep 25, 2015
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078003 001</u>	Jan 26, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078003 002</u>	Jan 26, 2015
<u>AB</u>	KREMERS URBAN PHARMS	<u>EQ 20MG BASE</u>	<u>A205563 001</u>	Sep 01, 2017
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205563 002</u>	Sep 01, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A078936 001</u>	Aug 02, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-160 (of 436)

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS;ORAL

ESOMEPRAZOLE MAGNESIUM

<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078936 002</u>	Aug 03, 2015
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A203636 001</u>	Oct 19, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203636 002</u>	Oct 19, 2015
				<b>NEXIUM</b>
<u>AB</u>	+ ASTRAZENECA PHARMS	<u>EQ 20MG BASE</u>	<u>N021153 001</u>	Feb 20, 2001
<u>AB</u>	+!	<u>EQ 40MG BASE</u>	<u>N021153 002</u>	Feb 20, 2001
	ESOMEPRAZOLE MAGNESIUM			
BX	HETERO LABS LTD III	EQ 20MG BASE	A202784 001	Sep 21, 2015
BX		EQ 40MG BASE	A202784 002	Sep 21, 2015
	FOR SUSPENSION, DELAYED RELEASE;ORAL			
	NEXIUM			
	+ ASTRAZENECA PHARMS	EQ 2.5MG BASE/PACKET	N021957 003	Dec 15, 2011
	+	EQ 5MG BASE/PACKET	N021957 004	Dec 15, 2011
	+	EQ 10MG BASE/PACKET	N022101 001	Feb 27, 2008
	+	EQ 20MG BASE/PACKET	N021957 001	Oct 20, 2006
	+!	EQ 40MG BASE/PACKET	N021957 002	Oct 20, 2006

ESOMEPRAZOLE MAGNESIUM; NAPROXEN

TABLET, DELAYED RELEASE;ORAL

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 20MG BASE;375MG</u>	<u>A202461 001</u>	Sep 27, 2013
<u>AB</u>		<u>EQ 20MG BASE;500MG</u>	<u>A202461 002</u>	Sep 27, 2013
				<b>VIMOVO</b>
<u>AB</u>	+ HORIZON PHARMA USA	<u>EQ 20MG BASE;375MG</u>	<u>N022511 002</u>	Apr 30, 2010
<u>AB</u>	+!	<u>EQ 20MG BASE;500MG</u>	<u>N022511 001</u>	Apr 30, 2010

ESOMEPRAZOLE SODIUM

INJECTABLE;INTRAVENOUS

ESOMEPRAZOLE SODIUM

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 40MG BASE/VIAL</u>	<u>A205379 001</u>	Sep 25, 2015
<u>AP</u>	! AUROBINDO PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A204657 002</u>	Aug 10, 2016
<u>AP</u>	DEVA HOLDING AS	<u>EQ 40MG BASE/VIAL</u>	<u>A207181 001</u>	Mar 06, 2017
<u>AP</u>	MYLAN LABS LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A202686 002</u>	May 17, 2017
<u>AP</u>	SUN PHARMA GLOBAL	<u>EQ 40MG BASE/VIAL</u>	<u>A200882 002</u>	Mar 18, 2013
				<b>NEXIUM IV</b>
<u>AP</u>	+! ASTRAZENECA PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>N021689 002</u>	Mar 31, 2005

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE STRONTIUM

+!	R2 PHARMA LLC	49.3MG	N202342 002	Aug 06, 2013
----	---------------	--------	-------------	--------------

ESTAZOLAM

TABLET;ORAL

ESTAZOLAM

<u>AB</u>	MAYNE PHARMA	<u>1MG</u>	<u>A074921 001</u>	Jul 10, 1997
<u>AB</u>	!	<u>2MG</u>	<u>A074921 002</u>	Jul 10, 1997
<u>AB</u>	PAR PHARM	<u>1MG</u>	<u>A074826 001</u>	Jul 03, 1997
<u>AB</u>		<u>2MG</u>	<u>A074826 002</u>	Jul 03, 1997
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>A074818 001</u>	Aug 19, 1997
<u>AB</u>		<u>2MG</u>	<u>A074818 002</u>	Aug 19, 1997

ESTRADIOL

CREAM;VAGINAL

ESTRACE

<u>AB</u>	! ALLERGAN SALES LLC	<u>0.01%</u>	<u>A086069 001</u>	Jan 31, 1984
				<b>ESTRADIOL</b>

<u>AB</u>	MYLAN PHARMS INC	<u>0.01%</u>	<u>A208788 001</u>	Dec 29, 2017
	FILM, EXTENDED RELEASE;TRANSDERMAL			

CLIMARA

<u>AB</u>	+ BAYER HLTHCARE	<u>0.0375MG/24HR</u>	<u>N020375 005</u>	May 27, 2003
<u>AB</u>	+	<u>0.06MG/24HR</u>	<u>N020375 006</u>	May 27, 2003

ESTRADIOL

<u>AB</u>	MYLAN TECHNOLOGIES	<u>0.0375MG/24HR</u>	<u>A075182 004</u>	Jul 20, 2006
<u>AB</u>		<u>0.06MG/24HR</u>	<u>A075182 005</u>	Jul 20, 2006
<u>AB1</u>		<u>0.025MG/24HR</u>	<u>A201675 001</u>	Dec 19, 2014
<u>AB1</u>		<u>0.0375MG/24HR</u>	<u>A201675 002</u>	Dec 19, 2014
<u>AB1</u>		<u>0.05MG/24HR</u>	<u>A201675 003</u>	Dec 19, 2014
<u>AB1</u>		<u>0.075MG/24HR</u>	<u>A201675 004</u>	Dec 19, 2014
<u>AB1</u>		<u>0.1MG/24HR</u>	<u>A201675 005</u>	Dec 19, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-161 (of 436)

**ESTRADIOL**

FILM, EXTENDED RELEASE;TRANSDERMAL

**VIVELLE-DOT**

<b>AB1</b>	+	NOVARTIS	<b><u>0.025MG/24HR</u></b>
<b>AB1</b>	+		<b><u>0.0375MG/24HR</u></b>
<b>AB1</b>	+		<b><u>0.05MG/24HR</u></b>
<b>AB1</b>	+		<b><u>0.075MG/24HR</u></b>
<b>AB1</b>	+!		<b><u>0.1MG/24HR</u></b>

<b>N020538</b>	<b>009</b>	May 03, 2002
<b>N020538</b>	<b>005</b>	Jan 08, 1999
<b>N020538</b>	<b>006</b>	Jan 08, 1999
<b>N020538</b>	<b>007</b>	Jan 08, 1999
<b>N020538</b>	<b>008</b>	Jan 08, 1999

**CLIMARA**

<b>AB2</b>	+	BAYER HLTHCARE	<b><u>0.025MG/24HR</u></b>
<b>AB2</b>	+		<b><u>0.05MG/24HR</u></b>
<b>AB2</b>	+		<b><u>0.075MG/24HR</u></b>
<b>AB2</b>	+!		<b><u>0.1MG/24HR</u></b>

<b>N020375</b>	<b>004</b>	Mar 05, 1999
<b>N020375</b>	<b>001</b>	Dec 22, 1994
<b>N020375</b>	<b>003</b>	Mar 23, 1998
<b>N020375</b>	<b>002</b>	Dec 22, 1994

**ESTRADIOL**

<b>AB2</b>	MYLAN TECHNOLOGIES	<b><u>0.025MG/24HR</u></b>
<b>AB2</b>		<b><u>0.05MG/24HR</u></b>
<b>AB2</b>		<b><u>0.075MG/24HR</u></b>
<b>AB2</b>		<b><u>0.1MG/24HR</u></b>

<b>A075182</b>	<b>003</b>	Jan 26, 2005
<b>A075182</b>	<b>006</b>	Feb 24, 2000
<b>A075182</b>	<b>002</b>	Jan 26, 2005
<b>A075182</b>	<b>001</b>	Feb 24, 2000

ALORA

BX	ALLERGAN SALES LLC	0.025MG/24HR
BX		0.05MG/24HR
BX		0.075MG/24HR
BX		0.1MG/24HR

N020655	004	Apr 05, 2002
N020655	001	Dec 20, 1996
N020655	002	Dec 20, 1996
N020655	003	Dec 20, 1996

MENOSTAR

+!	BAYER HLTHCARE	0.014MG/24HR
----	----------------	--------------

N021674	001	Jun 08, 2004
---------	-----	--------------

MINIVELLE

+	NOVEN	0.025MG/24HR
+		0.0375MG/24HR
+		0.05MG/24HR
+		0.075MG/24HR
+!		0.1MG/24HR

N203752	005	Sep 23, 2014
N203752	001	Oct 29, 2012
N203752	003	Oct 29, 2012
N203752	002	Oct 29, 2012
N203752	004	Oct 29, 2012

GEL;TRANSDERMAL

DIVIGEL

+	VERTICAL PHARMS LLC	0.1% (0.25GM/PACKET)
+		0.1% (0.5GM/PACKET)
+!		0.1% (1GM/PACKET)

N022038	001	Jun 04, 2007
N022038	002	Jun 04, 2007
N022038	003	Jun 04, 2007

GEL, METERED;TRANSDERMAL

ELESTRIN

+!	MYLAN SPECIALITY LP	0.06% (0.87GM/ACTIVATION)
----	---------------------	---------------------------

N021813	001	Dec 15, 2006
---------	-----	--------------

ESTROGEL

+!	ASCEND THERAPS US	0.06% (1.25GM/ACTIVATION)
----	-------------------	---------------------------

N021166	002	Feb 09, 2004
---------	-----	--------------

INSERT, EXTENDED RELEASE;VAGINAL

ESTRING

+!	PHARMACIA AND UPJOHN	0.0075MG/24HR
----	----------------------	---------------

N020472	001	Apr 26, 1996
---------	-----	--------------

SPRAY;TRANSDERMAL

EVAMIST

+!	PERRIGO PHARMA INTL	1.53MG/SPRAY
----	---------------------	--------------

N022014	001	Jul 27, 2007
---------	-----	--------------

TABLET;ORAL

**ESTRADIOL**

<b>AB</b>	BARR LABS INC	<b><u>0.5MG</u></b>
<b>AB</b>		<b><u>1MG</u></b>
<b>AB</b>	!	<b><u>2MG</u></b>
<b>AB</b>	EPIC PHARMA INC	<b><u>0.5MG</u></b>
<b>AB</b>		<b><u>1MG</u></b>
<b>AB</b>		<b><u>2MG</u></b>
<b>AB</b>	MAYNE PHARMA	<b><u>0.5MG</u></b>
<b>AB</b>		<b><u>1MG</u></b>
<b>AB</b>		<b><u>2MG</u></b>
<b>AB</b>	MYLAN	<b><u>0.5MG</u></b>
<b>AB</b>		<b><u>1MG</u></b>
<b>AB</b>		<b><u>2MG</u></b>

<b>A040197</b>	<b>001</b>	Oct 22, 1997
<b>A040197</b>	<b>002</b>	Oct 22, 1997
<b>A040197</b>	<b>003</b>	Oct 22, 1997
<b>A040275</b>	<b>001</b>	Dec 29, 1998
<b>A040275</b>	<b>002</b>	Dec 29, 1998
<b>A040275</b>	<b>003</b>	Dec 29, 1998
<b>A040114</b>	<b>003</b>	Mar 14, 1996
<b>A040114</b>	<b>001</b>	Mar 14, 1996
<b>A040114</b>	<b>002</b>	Mar 14, 1996
<b>A040326</b>	<b>001</b>	Apr 21, 1999
<b>A040326</b>	<b>002</b>	Apr 21, 1999
<b>A040326</b>	<b>003</b>	Apr 21, 1999

TABLET;VAGINAL

**ESTRADIOL**

<b>AB</b>	AMNEAL PHARMS	<b><u>10MCG</u></b>
<b>AB</b>	TEVA PHARMS USA	<b><u>10MCG</u></b>
<b>AB</b>	<b><u>VAGIFEM</u></b>	

<b>A205256</b>	<b>001</b>	May 29, 2015
<b>A206388</b>	<b>001</b>	Jul 21, 2017
<b>N020908</b>	<b>002</b>	Nov 25, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-162 (of 436)

ESTRADIOL ACETATE

INSERT, EXTENDED RELEASE;VAGINAL

FEMRING

+ APIL  
+!

EQ 0.05MG BASE/24HR  
EQ 0.1MG BASE/24HR

N021367 001 Mar 20, 2003  
N021367 002 Mar 20, 2003

ESTRADIOL CYPIONATE

INJECTABLE;INJECTION

DEPO-ESTRADIOL

! PHARMACIA AND  
UPJOHN

5MG/ML

A085470 003

ESTRADIOL HEMIHYDRATE

EMULSION;TOPICAL

ESTRASORB

+! EXELTIS USA INC

0.25%

N021371 001 Oct 09, 2003

ESTRADIOL VALERATE

INJECTABLE;INJECTION

DELESTROGEN

AO +! PAR STERILE  
PRODUCTS

20MG/ML

N009402 004

AO +!

40MG/ML

N009402 003

ESTRADIOL VALERATE

AO LUITPOLD

20MG/ML

A090920 001 Jan 19, 2010

AO

40MG/ML

A090920 002 Jan 19, 2010

DELESTROGEN

+! PAR STERILE  
PRODUCTS

10MG/ML

N009402 002

ESTRADIOL; LEVONORGESTREL

FILM, EXTENDED RELEASE;TRANSDERMAL

CLIMARA PRO

+! BAYER HLTHCARE

0.045MG/24HR;0.015MG/24HR

N021258 001 Nov 21, 2003

ESTRADIOL; NORETHINDRONE ACETATE

FILM, EXTENDED RELEASE;TRANSDERMAL

COBIPATCH

+ NOVEN PHARMS INC

0.05MG/24HR;0.14MG/24HR

N020870 001 Aug 07, 1998

+!

0.05MG/24HR;0.25MG/24HR

N020870 002 Aug 07, 1998

TABLET;ORAL

ACTIVELLA

AB + AMNEAL PHARMS LLC

0.5MG;0.1MG

N020907 002 Dec 28, 2006

AB +!

1MG;0.5MG

N020907 001 Nov 18, 1998

AMABELZ

AB LUPIN LTD

0.5MG;0.1MG

A203339 001 Jun 20, 2016

AB

1MG;0.5MG

A203339 002 Jun 20, 2016

ESTRADIOL AND NORETHINDRONE ACETATE

AB BARR

1MG;0.5MG

A079193 001 May 11, 2010

AB BRECKENRIDGE PHARM

0.5MG;0.1MG

A078324 002 Jun 09, 2011

AB

1MG;0.5MG

A078324 001 Apr 17, 2008

AB MYLAN LABS LTD

0.5MG;0.1MG

A207261 001 Feb 10, 2017

AB

1MG;0.5MG

A207261 002 Feb 10, 2017

AB TEVA PHARMS USA

0.5MG;0.1MG

A200747 001 Mar 08, 2012

ESTRADIOL; NORGESTIMATE

TABLET;ORAL

ESTRADIOL AND NORGESTIMATE

!

BARR

1MG,1MG;N/A,0.09MG

A076812 001 Apr 29, 2005

ESTRAMUSTINE PHOSPHATE SODIUM

CAPSULE;ORAL

EMCYT

+! PHARMACIA AND  
UPJOHN

EQ 140MG PHOSPHATE

N018045 001

ESTROGENS, CONJUGATED

CREAM;TOPICAL, VAGINAL

PREMARIN

+! WYETH PHARMS INC

0.625MG/GM

N020216 001

INJECTABLE;INJECTION

PREMARIN

+! WYETH PHARMS INC

25MG/VIAL

N010402 001

TABLET;ORAL

PREMARIN

+ WYETH PHARMS INC

0.3MG

N004782 003

+

0.45MG

N004782 006 Jul 16, 2003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-163 (of 436)

ESTROGENS, CONJUGATED

TABLET;ORAL

PREMARIN

+!	0.625MG	N004782 004
+!	0.9MG	N004782 005 Jan 26, 1984
+!	1.25MG	N004782 001

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET;ORAL-28

PREMPHASE 14/14

+! WYETH PHARMS INC	0.625MG, 0.625MG;N/A, 5MG	N020527 002 Nov 17, 1995
PREMPRO		
+! WYETH PHARMS INC	0.3MG;1.5MG	N020527 005 Jun 04, 2003
+!	0.45MG;1.5MG	N020527 004 Mar 12, 2003
+!	0.625MG;2.5MG	N020527 001 Nov 17, 1995
+!	0.625MG;5MG	N020527 003 Jan 09, 1998

ESTROGENS, ESTERIFIED

TABLET;ORAL

MENEST

MONARCH PHARMS	0.3MG	A084951 001
	0.625MG	A084948 001
	1.25MG	A084950 001
!	2.5MG	A084949 001

ESTROPIPATE

TABLET;ORAL

ESTROPIPATE

MYLAN	0.75MG	A040359 001 Aug 26, 1999
	1.5MG	A040359 002 Aug 26, 1999
OGEN 5		
PHARMACIA AND UPJOHN	6MG	A083220 004

ESZOPICLONE

TABLET;ORAL

**ESZOPICLONE**

<b>AB</b>	AUROBINDO PHARMA LTD	<b>1MG</b>	<b>A208451 001</b> Sep 15, 2016
<b>AB</b>		<b>2MG</b>	<b>A208451 002</b> Sep 15, 2016
<b>AB</b>		<b>3MG</b>	<b>A208451 003</b> Sep 15, 2016
<b>AB</b>	DR REDDYS LABS LTD	<b>1MG</b>	<b>A091024 001</b> Apr 15, 2014
<b>AB</b>		<b>2MG</b>	<b>A091024 002</b> Apr 15, 2014
<b>AB</b>		<b>3MG</b>	<b>A091024 003</b> Apr 15, 2014
<b>AB</b>	GLENMARK GENERICS	<b>1MG</b>	<b>A091166 001</b> Apr 15, 2014
<b>AB</b>		<b>2MG</b>	<b>A091166 002</b> Apr 15, 2014
<b>AB</b>		<b>3MG</b>	<b>A091166 003</b> Apr 15, 2014
<b>AB</b>	LUPIN LTD	<b>1MG</b>	<b>A091124 001</b> Sep 13, 2011
<b>AB</b>		<b>2MG</b>	<b>A091124 002</b> Sep 13, 2011
<b>AB</b>		<b>3MG</b>	<b>A091124 003</b> Sep 13, 2011
<b>AB</b>	MACLEODS PHARMS LTD	<b>1MG</b>	<b>A202929 001</b> Jan 30, 2015
<b>AB</b>		<b>2MG</b>	<b>A202929 002</b> Jan 30, 2015
<b>AB</b>		<b>3MG</b>	<b>A202929 003</b> Jan 30, 2015
<b>AB</b>	MYLAN PHARMS INC	<b>1MG</b>	<b>A091151 001</b> Mar 26, 2013
<b>AB</b>		<b>2MG</b>	<b>A091151 002</b> Mar 26, 2013
<b>AB</b>		<b>3MG</b>	<b>A091151 003</b> Mar 26, 2013
<b>AB</b>	ORCHID HLTHCARE	<b>1MG</b>	<b>A091113 001</b> Jun 10, 2014
<b>AB</b>		<b>2MG</b>	<b>A091113 002</b> Jun 10, 2014
<b>AB</b>		<b>3MG</b>	<b>A091113 003</b> Jun 10, 2014
<b>AB</b>	SUN PHARMA GLOBAL	<b>1MG</b>	<b>A091103 001</b> Apr 03, 2013
<b>AB</b>		<b>2MG</b>	<b>A091103 002</b> Apr 03, 2013
<b>AB</b>		<b>3MG</b>	<b>A091103 003</b> Apr 03, 2013
<b>AB</b>	TEVA	<b>1MG</b>	<b>A091169 001</b> May 23, 2011
<b>AB</b>		<b>2MG</b>	<b>A091169 002</b> May 23, 2011
<b>AB</b>		<b>3MG</b>	<b>A091169 003</b> May 23, 2011
<b>AB</b>	WEST-WARD PHARMS INT	<b>1MG</b>	<b>A091153 001</b> Apr 15, 2014
<b>AB</b>		<b>2MG</b>	<b>A091153 002</b> Apr 15, 2014
<b>AB</b>		<b>3MG</b>	<b>A091153 003</b> Apr 15, 2014
<b>LUNESTA</b>			
<b>AB</b>	+ SUNOVION PHARMS INC	<b>1MG</b>	<b>N021476 001</b> Dec 15, 2004
<b>AB</b>	+	<b>2MG</b>	<b>N021476 002</b> Dec 15, 2004
<b>AB</b>	+!	<b>3MG</b>	<b>N021476 003</b> Dec 15, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-164 (of 436)

ETELCALCETIDE

SOLUTION; INTRAVENOUS

PARSABIV

+!	KAI PHARMS INC	2.5MG/0.5ML (2.5MG/0.5ML)	N208325 001 Feb 07, 2017
+!		5MG/ML (5MG/ML)	N208325 002 Feb 07, 2017
+!		10MG/2ML (5MG/ML)	N208325 003 Feb 07, 2017

ETEPLIRSEN

SOLUTION; IV (INFUSION)

EXONDYS 51

+!	SAREPTA THERAPS INC	100MG/2ML (50MG/ML)	N206488 001 Sep 19, 2016
+!		500MG/10ML (50MG/ML)	N206488 002 Sep 19, 2016

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECRRIN

<u>AP</u>	+!	ATON	<u>EQ 50MG BASE/VIAL</u>	<u>N016093 001</u>
<u>AP</u>		<u>ETHACRYNATE SODIUM</u>		
<u>AP</u>		MYLAN INSTITUTIONAL	<u>EQ 50MG BASE/VIAL</u>	<u>A204634 001</u> Aug 23, 2016
<u>AP</u>		PAR STERILE PRODUCTS	<u>EQ 50MG BASE/VIAL</u>	<u>A205473 001</u> Jul 29, 2015

ETHACRYNIC ACID

TABLET; ORAL

EDECRRIN

<u>AB</u>	+!	ATON	<u>25MG</u>	<u>N016092 001</u>
		<u>ETHACRYNIC ACID</u>		
<u>AB</u>		EDENBRIDGE PHARMS	<u>25MG</u>	<u>A205609 001</u> Jun 30, 2016
<u>AB</u>		PAR PHARM INC	<u>25MG</u>	<u>A208501 001</u> Jul 21, 2017

AB

WEST-WARD PHARMS INT

<u>AB</u>		AKORN	<u>100MG</u>	<u>A075095 001</u> Nov 30, 1999
<u>AB</u>			<u>400MG</u>	<u>A075095 002</u> Nov 30, 1999
<u>AB</u>		BARR	<u>400MG</u>	<u>A076057 001</u> Nov 26, 2001
<u>AB</u>		LUPIN	<u>100MG</u>	<u>A078939 001</u> Jun 17, 2009
<u>AB</u>			<u>400MG</u>	<u>A078939 002</u> Jun 17, 2009
		<u>MYAMBUTOL</u>		
<u>AB</u>	+	STI PHARMA LLC	<u>100MG</u>	<u>N016320 001</u>
<u>AB</u>	+		<u>400MG</u>	<u>N016320 003</u>

ETHANOLAMINE OLEATE

INJECTABLE; INJECTION

ETHAMOLIN

+!	QOL MEDCL	50MG/ML	N019357 001 Dec 22, 1988
----	-----------	---------	--------------------------

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-28

ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL

<u>AB</u>		MYLAN LABS LTD	<u>0.035MG;1MG</u>	<u>A204703 001</u> Jul 28, 2016
<u>AB</u>			<u>0.05MG;1MG</u>	<u>A204704 001</u> Feb 09, 2016
		<u>KELNOR</u>		
<u>AB</u>		BARR	<u>0.035MG;1MG</u>	<u>A076785 001</u> May 23, 2005
		<u>ZOVIA 1/35E-28</u>		
<u>AB</u>		MAYNE PHARMA	<u>0.035MG;1MG</u>	<u>A072721 001</u> Dec 30, 1991
		<u>ZOVIA 1/50E-28</u>		
<u>AB</u>	!	WATSON LABS	<u>0.05MG;1MG</u>	<u>A072723 001</u> Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL

NUVARING

+!	ORGANON SUB MERCK	0.015MG/24HR;0.12MG/24HR	N021187 001 Oct 03, 2001
----	-------------------	--------------------------	--------------------------

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

ASHLYNA

<u>AB</u>		GLENMARK GENERICS	<u>0.03MG, 0.01MG;0.15MG, N/A</u>	<u>A203163 001</u> Feb 23, 2015
		<u>DAYSEE</u>		
<u>AB</u>		LUPIN LTD	<u>0.03MG, 0.01MG;0.15MG, N/A</u>	<u>A091467 001</u> Apr 10, 2013
		<u>FAYOSIM</u>		
<u>AB</u>		LUPIN LTD	<u>0.02MG, 0.15MG;0.025MG, 0.15MG;0.03MG, 0.15MG, N/A</u>	<u>A205943 001</u> Mar 29, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-165 (of 436)

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL

INTROVALE

AB LABS LEON FARMA 0.03MG;0.15MG

A079064 001 Sep 27, 2010

JAIMIESS

AB LABS LEON FARMA 0.03MG,0.01MG;0.15MG,N/A

A203770 001 Dec 27, 2017

LEVONORGESTREL AND ETHINYL ESTRADIOOL

AB AMNEAL PHARMS 0.03MG;0.15MG

A203871 001 Nov 13, 2015

AB 0.03MG,0.01MG;0.15MG,N/A

A203872 001 Dec 22, 2015

AB GLENMARK GENERICS 0.02MG;0.09MG

A202791 001 Apr 09, 2015

AB GLENMARK PHARMS LTD 0.03MG;0.15MG

A203164 001 Jun 12, 2015

AB LUPIN LTD 0.02MG,0.01MG;0.1MG,N/A

A091674 001 Oct 26, 2011

AB 0.03MG;0.15MG

A091440 001 Oct 23, 2012

AB MAYNE PHARMA 0.02MG,0.01MG;0.1MG,N/A

A200407 001 Oct 25, 2011

AB MYLAN LABS LTD 0.03MG;0.15MG

A200490 001 Apr 21, 2015

AB ! WATSON LABS 0.02MG;0.09MG

A079218 001 Jun 06, 2011

LEVONORGESTREL AND ETHINYL ESTRADIOOL AND ETHINYL ESTRADIOL

AB LABS LEON FARMA 0.02MG,0.01MG;0.1MG,N/A

A205131 001 Dec 14, 2017

AB MAYNE PHARMA 0.03MG,0.01MG;0.15MG,N/A

A078834 001 May 31, 2011

AB MYLAN LABS LTD 0.02MG,0.15MG;

A206053 001 Oct 02, 2017

AB 0.025MG,0.15MG;0.03MG,0.15MG;

A200493 001 Jun 17, 2015

AB 0.02MG,0.01MG;0.1MG,N/A

A200492 001 May 27, 2015

LO SIMPESSE

AB AUROBINDO PHARMA LTD 0.02MG,0.01MG;0.01MG,N/A

A206852 001 Apr 28, 2017

LOSEASONIQUE

AB TEVA BRANDED PHARM 0.02MG,0.01MG;0.1MG,N/A

N022262 001 Oct 24, 2008

QUARTETTE

AB +! TEVA BRANDED PHARM 0.02MG,0.15MG;0.025MG,0.15MG;0.03MG,0.15MG;0.01MG,N/A

N204061 001 Mar 28, 2013

QUASENSE

AB WATSON LABS 0.03MG;0.15MG

A077101 001 Sep 06, 2006

SEASONALE

AB +! TEVA BRANDED PHARM 0.03MG;0.15MG

N021544 001 Sep 05, 2003

SEASONIQUE

AB +! TEVA BRANDED PHARM 0.03MG,0.01MG;0.15MG,N/A

N021840 001 May 25, 2006

SETLAKIN

AB NOVAST LABS LTD 0.03MG;0.15MG

A090716 001 Sep 15, 2014

SIMPESSE

AB AUROBINDO PHARMA LTD 0.03MG,0.01MG;0.15MG,N/A

A206851 001 Apr 07, 2017

TABLET;ORAL-28

ALTAVERA

AB LABS LEON FARMA 0.03MG;0.15MG

A079102 001 Aug 03, 2010

AYUNA

AB AUROBINDO PHARMA LTD 0.03MG;0.15MG

A206866 001 Sep 23, 2016

ELIFEMME

AB LABS LEON FARMA 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG

A202507 001 Dec 04, 2015

ENPRESSE-28

AB DURAMED PHARMS BARR 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG

A075809 002 Jul 16, 2001

KURVELO

AB LUPIN LTD 0.03MG;0.15MG

A091408 001 Oct 17, 2012

LEVONEST

AB NOVAST LABS LTD 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG

A090719 001 Dec 29, 2010

LEVONORGESTREL AND ETHINYL ESTRADIOOL

AB AMNEAL PHARMS 0.03MG;0.15MG

A201095 001 Dec 08, 2014

AB LUPIN LTD 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG

A200248 001 Nov 19, 2015

AB MYLAN LABS LTD 0.03MG;0.15MG

A091663 001 Dec 21, 2012

LEVORA 0.15/30-28

AB ! MAYNE PHARMA 0.03MG;0.15MG

A073594 001 Dec 13, 1993

MARLISSA

AB GLENMARK GENERICS 0.03MG;0.15MG

A091452 001 Feb 29, 2012

MYZILRA

AB VINTAGE PHARMS LLC 0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG

A077502 001 Nov 23, 2011

PORTIA-28

AB BARR 0.03MG;0.15MG

A075866 002 May 23, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-166 (of 436)

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL-28

TRIVORA-28

<u>AB</u>	! MAYNE PHARMA	<u>0.03MG, 0.04MG, 0.03MG, 0.05MG, 0.075MG, 0.1 25MG</u>	<u>A074538 002</u>	Dec 18, 1997
-----------	----------------	--	--------------------	--------------

AFIRMELLE

<u>AB1</u>	AUROBINDO PHARMA LTD	<u>0.02MG; 0.1MG</u>	<u>A206886 001</u>	Nov 14, 2016
------------	----------------------	----------------------	--------------------	--------------

AVIANE-28

<u>AB1</u>	DURAMED PHARMS BARR	<u>0.02MG; 0.1MG</u>	<u>A075796 001</u>	Apr 30, 2001
------------	---------------------	----------------------	--------------------	--------------

FALMINA

<u>AB1</u>	NOVAST LABS LTD	<u>0.02MG; 0.1MG</u>	<u>A090721 001</u>	Mar 28, 2012
------------	-----------------	----------------------	--------------------	--------------

LEVONORGESTREL AND ETHINYL ESTRADIOL

<u>AB1</u>	AMNEAL PHARMS	<u>0.02MG; 0.1MG</u>	<u>A201108 001</u>	Feb 05, 2014
------------	---------------	----------------------	--------------------	--------------

<u>AB1</u>	LUPIN LTD	<u>0.02MG; 0.1MG</u>	<u>A091425 001</u>	Jan 18, 2013
------------	-----------	----------------------	--------------------	--------------

<u>AB1</u>	! MAYNE PHARMA	<u>0.02MG; 0.1MG</u>	<u>A076625 001</u>	Nov 18, 2004
------------	----------------	----------------------	--------------------	--------------

<u>AB1</u>	MYLAN LABS LTD	<u>0.02MG; 0.1MG</u>	<u>A200245 001</u>	Oct 09, 2013
------------	----------------	----------------------	--------------------	--------------

ORSYTHIA

<u>AB1</u>	VINTAGE PHARMS LLC	<u>0.02MG; 0.1MG</u>	<u>A077099 001</u>	May 11, 2011
------------	--------------------	----------------------	--------------------	--------------

VIENVA

<u>AB1</u>	LABS LEON FARMA	<u>0.02MG; 0.1MG</u>	<u>A201088 001</u>	May 21, 2015
------------	-----------------	----------------------	--------------------	--------------

LESSINA-28

<u>AB2</u>	BARR	<u>0.02MG; 0.1MG</u>	<u>A075803 002</u>	Mar 20, 2002
------------	------	----------------------	--------------------	--------------

LEVONORGESTREL AND ETHINYL ESTRADIOL

<u>AB2</u>	! MAYNE PHARMA	<u>0.02MG; 0.1MG</u>	<u>A077681 001</u>	May 31, 2006
------------	----------------	----------------------	--------------------	--------------

<u>AB2</u>	MYLAN LABS LTD	<u>0.02MG; 0.1MG</u>	<u>A202247 001</u>	Dec 08, 2014
------------	----------------	----------------------	--------------------	--------------

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE;TRANSDERMAL

XULANE

!	MYLAN TECHNOLOGIES	0.035MG/24HR; 0.15MG/24HR	A200910 001	Apr 16, 2014
---	--------------------	---------------------------	-------------	--------------

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET;ORAL-21

NORINYL 1+35 21-DAY

<u>AB</u>	ALLERGAN SALES LLC	<u>0.035MG; 1MG</u>	<u>N017565 001</u>	
-----------	--------------------	---------------------	--------------------	--

NORTREL 1/35-21

<u>AB</u>	BARR	<u>0.035MG; 1MG</u>	<u>A072693 001</u>	Feb 28, 1992
	NORTREL 7/7/7			
	BARR	0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG	A075478 001	Aug 30, 2002
	G			

TABLET;ORAL-28

ALYACEN 1/35

<u>AB</u>	GLENMARK GENERICS	<u>0.035MG; 1MG</u>	<u>A091634 001</u>	Jan 19, 2012
-----------	-------------------	---------------------	--------------------	--------------

ALYACEN 7/7/7

<u>AB</u>	GLENMARK GENERICS	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG</u>	<u>A091636 001</u>	Jan 19, 2012
	G			

ARANELLE

<u>AB</u>	BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MG</u>	<u>A076783 001</u>	Sep 29, 2004
-----------	------	---	--------------------	--------------

BALZIVA-28

<u>AB</u>	BARR	<u>0.035MG; 0.4MG</u>	<u>A076238 001</u>	Apr 22, 2004
-----------	------	-----------------------	--------------------	--------------

BREVICON 28-DAY

<u>AB</u>	ALLERGAN SALES LLC	<u>0.035MG; 0.5MG</u>	<u>N017743 001</u>	
-----------	--------------------	-----------------------	--------------------	--

BRIELLYN

<u>AB</u>	GLENMARK GENERICS	<u>0.035MG; 0.4MG</u>	<u>A090538 001</u>	Mar 22, 2011
-----------	-------------------	-----------------------	--------------------	--------------

CYCLAFEM 0.5/35

<u>AB</u>	VINTAGE PHARMS	<u>0.035MG; 0.5MG</u>	<u>A203413 001</u>	Dec 16, 2015
-----------	----------------	-----------------------	--------------------	--------------

CYCLAFEM 1/35

<u>AB</u>	VINTAGE PHARMS LLC	<u>0.035MG; 1MG</u>	<u>A076337 001</u>	Nov 12, 2010
-----------	--------------------	---------------------	--------------------	--------------

CYCLAFEM 7/7/7

<u>AB</u>	VINTAGE PHARMS LLC	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG</u>	<u>A076338 001</u>	Nov 16, 2010
	G			

CYONANZ

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG; 0.5MG</u>	<u>A207055 001</u>	Oct 21, 2016
-----------	----------------------	-----------------------	--------------------	--------------

DASETTA 1/35

<u>AB</u>	NOVAST LABS LTD	<u>0.035MG; 1MG</u>	<u>A090948 001</u>	Dec 22, 2011
-----------	-----------------	---------------------	--------------------	--------------

DASETTA 7/7/7

<u>AB</u>	NOVAST LABS LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG</u>	<u>A090946 001</u>	Dec 22, 2011
	G			

GILDAGIA

<u>AB</u>	VINTAGE PHARMS	<u>0.035MG; 0.4MG</u>	<u>A078376 001</u>	Nov 06, 2012
-----------	----------------	-----------------------	--------------------	--------------

MODICON 28

<u>AB</u>	+! JANSSEN PHARMS	<u>0.035MG; 0.5MG</u>	<u>N017735 001</u>	
-----------	-------------------	-----------------------	--------------------	--

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-167 (of 436)

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET;ORAL-28

NORETHINDRONE AND ETHINYL ESTRADIOL

<u>AB</u>	ACCORD HLTHCARE	<u>0.035MG;1MG</u>	<u>A206864 001</u>	Apr 28, 2017
<u>AB</u>	MAYNE PHARMA	<u>0.035MG;0.5MG</u>	<u>A070686 001</u>	Jan 29, 1987
<u>AB</u>	! MYLAN LABS LTD	<u>0.035MG;0.4MG</u>	<u>A200897 001</u>	May 11, 2015
<u>AB</u>	WATSON LABS	<u>0.035MG;0.4MG</u>	<u>A078323 001</u>	Feb 04, 2010
<u>AB</u>	WATSON LABS TEVA	<u>0.035MG;1MG</u>	<u>A070687 001</u>	Jan 29, 1987
<b><u>NORINYL 1+35 28-DAY</u></b>				
<u>AB</u>	ALLERGAN SALES LLC	<u>0.035MG;1MG</u>	<u>N017565 002</u>	
<b><u>NORTREL 0.5/35-28</u></b>				
<u>AB</u>	BARR	<u>0.035MG;0.5MG</u>	<u>A072695 001</u>	Feb 28, 1992
<b><u>NORTREL 1/35-28</u></b>				
<u>AB</u>	BARR	<u>0.035MG;1MG</u>	<u>A072696 001</u>	Feb 28, 1992
<b><u>NORTREL 7/7/7</u></b>				
<u>AB</u>	BARR	<u>0.035MG, 0.035MG, 0.035MG;0.5MG, 0.75MG, 1M</u>	<u>A075478 002</u>	Aug 30, 2002
<b><u>G</u></b>				
<b><u>NYLIA 1/35</u></b>				
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG;1MG</u>	<u>A207056 001</u>	Oct 21, 2016
<b><u>NYLIA 7/7/7</u></b>				
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG, 0.035MG, 0.035MG;0.5MG, 0.75MG, 1M</u>	<u>A207054 001</u>	Oct 21, 2016
<b><u>G</u></b>				
<b><u>ORTHO-NOVUM 1/35-28</u></b>				
<u>AB</u>	+! JANSSEN PHARMS	<u>0.035MG;1MG</u>	<u>N017919 002</u>	
<b><u>ORTHO-NOVUM 7/7/7-28</u></b>				
<u>AB</u>	+! JANSSEN PHARMS	<u>0.035MG, 0.035MG, 0.035MG;0.5MG, 0.75MG, 1M</u>	<u>N018985 002</u>	Apr 04, 1984
<b><u>G</u></b>				
<b><u>PHILITH</u></b>				
<u>AB</u>	NOVAST LABS LTD	<u>0.035MG;0.4MG</u>	<u>A090947 001</u>	Dec 22, 2011
<b><u>PIRMELLA 1/35</u></b>				
<u>AB</u>	LUPIN LTD	<u>0.035MG;1MG</u>	<u>A201512 001</u>	Apr 24, 2013
<b><u>PIRMELLA 7/7/7</u></b>				
<u>AB</u>	LUPIN LTD	<u>0.035MG, 0.035MG, 0.035MG;0.5MG, 0.75MG, 1M</u>	<u>A201510 001</u>	Apr 24, 2013
<b><u>G</u></b>				
<b><u>TRI-NORINYL 28-DAY</u></b>				
<u>AB</u>	+! MAYNE PHARMA	<u>0.035MG, 0.035MG, 0.035MG;0.5MG, 1MG, 0.5MG</u>	<u>N018977 002</u>	Apr 13, 1984
<b><u>VYFEMLA</u></b>				
<u>AB</u>	LUPIN LTD	<u>0.035MG;0.4MG</u>	<u>A201886 001</u>	Sep 26, 2013
<b><u>WERA</u></b>				
<u>AB</u>	NOVAST LABS LTD	<u>0.035MG;0.5MG</u>	<u>A091204 001</u>	Mar 27, 2012
NORETHINDRONE AND ETHINYL ESTRADIOL				
! MYLAN LABS LTD 0.05MG;1MG				
NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)				
WATSON LABS TEVA 0.035MG, 0.035MG;0.5MG, 1MG				
TABLET, CHEWABLE;ORAL				
<b><u>FEMCON FE</u></b>				
<u>AB</u>	+! APIL	<u>0.035MG;0.4MG</u>	<u>N021490 001</u>	Nov 14, 2003
<b><u>KAITLIB FE</u></b>				
<u>AB</u>	LUPIN LTD	<u>0.025MG;0.8MG</u>	<u>A203448 001</u>	Dec 17, 2015
<b><u>NEXESTA FE</u></b>				
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG;0.4MG</u>	<u>A207535 001</u>	Feb 02, 2017
<b><u>NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u></b>				
<u>AB</u>	ACCORD HLTHCARE	<u>0.035MG;0.4MG</u>	<u>A207066 001</u>	Mar 29, 2017
<u>AB</u>	AMNEAL PHARMS	<u>0.035MG;0.4MG</u>	<u>A078892 001</u>	Sep 26, 2011
<u>AB</u>	+! APIL	<u>0.025MG;0.8MG</u>	<u>N022573 001</u>	Dec 22, 2010
<u>AB</u>	BARR	<u>0.035MG;0.4MG</u>	<u>A078965 001</u>	Aug 05, 2010
<u>AB</u>	LUPIN LTD	<u>0.035MG;0.4MG</u>	<u>A091332 001</u>	Mar 23, 2016
<u>AB</u>	MYLAN LABS LTD	<u>0.025MG;0.8MG</u>	<u>A203371 001</u>	Apr 23, 2014
<u>AB</u>		<u>0.035MG;0.4MG</u>	<u>A202086 001</u>	Apr 01, 2015
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE</u></b>				
CAPSULE;ORAL				
TAYTULLA				
+! APIL 0.02MG;1MG				
TABLET;ORAL				
<b><u>AUROVELA 24 FE</u></b>				
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207504 001</u>	Jun 15, 2017
<b><u>BLISOVI 24 FE</u></b>				
<u>AB</u>	LUPIN LTD	<u>0.02MG;1MG</u>	<u>A091398 001</u>	Oct 28, 2015
<b><u>FEMHRT</u></b>				
<u>AB</u>	APIL	<u>0.0025MG;0.5MG</u>	<u>N021065 001</u>	Jan 14, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-168 (of 436)

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL

FYAVOLV

<u>AB</u>	LUPIN LTD	<u>0.005MG;1MG</u>	<u>A204213 002</u>	Dec 10, 2015
<u>AB</u>		<u>0.0025MG;0.5MG</u>	<u>A204213 001</u>	Dec 10, 2015

GILDESS 24 FE

<u>AB</u>	VINTAGE PHARMS	<u>0.02MG;1MG</u>	<u>A090293 001</u>	Dec 01, 2014
-----------	----------------	-------------------	--------------------	--------------

LARIN 24 FE

<u>AB</u>	NOVAST LABS LTD	<u>0.02MG;1MG</u>	<u>A202994 001</u>	Feb 18, 2015
-----------	-----------------	-------------------	--------------------	--------------

LERIBANE

<u>AB</u>	NOVAST LABS LTD	<u>0.0025MG;0.5MG</u>	<u>A203435 002</u>	Jun 03, 2016
<u>AB</u>		<u>0.005MG;1MG</u>	<u>A203435 001</u>	Jun 03, 2016
<u>AB</u>		<u>0.005MG;1MG</u>	<u>A203435 001</u>	Jun 03, 2016

LO LOESTRIN FE

<u>AB</u>	+!	<u>APIL</u>	<u>0.01MG,0.01MG;1MG,N/A</u>	<u>N022501 001</u>	Oct 21, 2010
-----------	----	-------------	------------------------------	--------------------	--------------

LOESTRIN 24 FE

<u>AB</u>	+	<u>APIL</u>	<u>0.02MG;1MG</u>	<u>N021871 001</u>	Feb 17, 2006
-----------	---	-------------	-------------------	--------------------	--------------

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

<u>AB</u>	!	<u>BARR LABS INC</u>	<u>0.005MG;1MG</u>	<u>A076221 001</u>	Nov 06, 2009
<u>AB</u>		<u>GLENMARK GENERICS</u>	<u>0.0025MG;0.5MG</u>	<u>A203038 001</u>	Apr 02, 2015
<u>AB</u>		<u>GLENMARK GENERICS</u>	<u>0.0025MG;0.5MG</u>	<u>A203038 001</u>	Apr 02, 2015
<u>AB</u>		<u>MYLAN LABS LTD</u>	<u>0.005MG;1MG</u>	<u>A207260 001</u>	Feb 02, 2017
<u>AB</u>		<u>MYLAN LABS LTD</u>	<u>0.005MG;1MG</u>	<u>A207259 001</u>	Dec 27, 2016

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

<u>AB</u>	MYLAN LABS LTD	<u>0.01MG,0.01MG;1MG,N/A</u>	<u>A205049 001</u>	May 31, 2016
-----------	----------------	------------------------------	--------------------	--------------

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

<u>AB</u>	!	<u>AMNEAL PHARMS</u>	<u>0.02MG;1MG</u>	<u>A078267 001</u>	Sep 01, 2009
<u>AB</u>		<u>BARR LABS INC</u>	<u>0.02MG;1MG</u>	<u>A090938 001</u>	Dec 01, 2014
<u>AB</u>		<u>BARR LABS INC</u>	<u>0.02MG;1MG</u>	<u>A090938 001</u>	Dec 01, 2014
<u>AB</u>		<u>GLENMARK PHARMS LTD</u>	<u>0.02MG;1MG</u>	<u>A204847 001</u>	Nov 17, 2017
<u>AB</u>		<u>GLENMARK PHARMS LTD</u>	<u>0.02MG;1MG</u>	<u>A204847 001</u>	Nov 17, 2017
<u>AB</u>		<u>MYLAN LABS LTD</u>	<u>0.02MG;1MG</u>	<u>A202742 001</u>	Oct 30, 2014
<u>AB</u>		<u>MYLAN LABS LTD</u>	<u>0.02MG;1MG</u>	<u>A202742 001</u>	Oct 30, 2014

TABLET;ORAL-21

AUROVELA 1.5/30

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.03MG;1.5MG</u>	<u>A207581 001</u>	Jun 26, 2017
-----------	----------------------	---------------------	--------------------	--------------

AUROVELA 1/20

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207506 001</u>	Jun 16, 2017
-----------	----------------------	-------------------	--------------------	--------------

GILDESS 1.5/30

<u>AB</u>	VINTAGE PHARMS LLC	<u>0.03MG;1.5MG</u>	<u>A077075 002</u>	Jul 24, 2012
<u>AB</u>	VINTAGE PHARMS LLC	<u>0.02MG;1MG</u>	<u>A077077 002</u>	Jul 24, 2012
<u>AB</u>	VINTAGE PHARMS LLC	<u>0.02MG;1MG</u>	<u>A077077 002</u>	Jul 24, 2012

JUNEL 1.5/30

<u>AB</u>	BARR	<u>0.03MG;1.5MG</u>	<u>A076381 001</u>	May 30, 2003
<u>AB</u>	JUNEL 1/20	<u>0.02MG;1MG</u>	<u>A076380 001</u>	May 30, 2003
<u>AB</u>	JUNEL 1/20	<u>0.02MG;1MG</u>	<u>A076380 001</u>	May 30, 2003

LARIN 1.5/30

<u>AB</u>	NOVAST LABS LTD	<u>0.03MG;1.5MG</u>	<u>A202996 001</u>	Mar 20, 2014
<u>AB</u>	LARIN 1/20	<u>0.02MG;1MG</u>	<u>A202995 001</u>	Dec 04, 2013
<u>AB</u>	LARIN 1/20	<u>0.02MG;1MG</u>	<u>A202995 001</u>	Dec 04, 2013
<u>AB</u>	LOESTRIN 21 1.5/30	<u>0.03MG;1.5MG</u>	<u>N017875 001</u>	
<u>AB</u>	LOESTRIN 21 1.5/30	<u>0.03MG;1.5MG</u>	<u>N017875 001</u>	
<u>AB</u>	APIL	<u>0.02MG;1MG</u>	<u>N017876 001</u>	
<u>AB</u>	APIL	<u>0.02MG;1MG</u>	<u>N017876 001</u>	

MICROGESTIN 1.5/30

<u>AB</u>	MAYNE PHARMA	<u>0.03MG;1.5MG</u>	<u>A075548 002</u>	Jul 30, 2003
<u>AB</u>	MICROGESTIN 1/20	<u>0.02MG;1MG</u>	<u>A075647 002</u>	Jul 30, 2003
<u>AB</u>	MICROGESTIN 1/20	<u>0.02MG;1MG</u>	<u>A075647 002</u>	Jul 30, 2003
<u>AB</u>	MAYNE PHARMA	<u>0.02MG;1MG</u>	<u>A075647 001</u>	Jul 30, 2003
<u>AB</u>	MAYNE PHARMA	<u>0.02MG;1MG</u>	<u>A075647 001</u>	Jul 30, 2003

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL

<u>AB</u>	GLENMARK PHARMS LTD	<u>0.02MG;1MG</u>	<u>A206969 001</u>	Jan 20, 2016
<u>AB</u>	MYLAN LABS LTD	<u>0.02MG;1MG</u>	<u>A202771 001</u>	Nov 06, 2013
<u>AB</u>	MYLAN LABS LTD	<u>0.02MG;1MG</u>	<u>A202771 001</u>	Nov 06, 2013
<u>AB</u>		<u>0.03MG;1.5MG</u>	<u>A202770 001</u>	Feb 19, 2015
<u>AB</u>		<u>0.03MG;1.5MG</u>	<u>A202770 001</u>	Feb 19, 2015

TRI-LEGEST 21

BARR

0.02MG, 0.03MG, 0.035MG;1MG, 1MG, 1MG

A076405 001 Oct 26, 2007

TABLET;ORAL-28

AUROVELA FE 1.5/30

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.03MG;1.5MG</u>	<u>A207580 001</u>	Jun 15, 2017
-----------	----------------------	---------------------	--------------------	--------------

AUROVELA FE 1/20

<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.02MG;1MG</u>	<u>A207505 001</u>	Jun 16, 2017
-----------	----------------------	-------------------	--------------------	--------------

BLISOVI FE 1.5/30

<u>AB</u>	LUPIN LTD	<u>0.03MG;1.5MG</u>	<u>A201585 001</u>	Nov 18, 2015
<u>AB</u>	BLISOVI FE 1/20	<u>0.02MG;1MG</u>	<u>A201584 001</u>	Nov 18, 2015
<u>AB</u>	BLISOVI FE 1/20	<u>0.02MG;1MG</u>	<u>A201584 001</u>	Nov 18, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-169 (of 436)

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL-28

ESTROSTEP FE

<u>AB</u>	+!	APIL	<u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u>	<u>N020130 002</u>	Oct 09, 1996
<u>AB</u>		<u>GILDESS FE 1.5/30</u>	<u>0.03MG; 1.5MG</u>	<u>A077075 001</u>	Apr 28, 2005
<u>AB</u>		VINTAGE PHARMS LLC	<u>0.02MG; 1MG</u>	<u>A077077 001</u>	May 20, 2005
<u>AB</u>		VINTAGE PHARMS LLC	<u>0.02MG; 1.5MG</u>	<u>A206597 001</u>	Nov 21, 2017
<u>AB</u>		HAILEY FE 1/20	<u>0.02MG; 1MG</u>	<u>A076064 001</u>	Sep 18, 2003
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.02MG; 1.5MG</u>	<u>A076081 001</u>	Sep 18, 2003
<u>AB</u>		JUNEL FE 1.5/30	<u>0.03MG; 1.5MG</u>	<u>A091453 001</u>	Aug 23, 2013
<u>AB</u>		BARR	<u>0.02MG; 1MG</u>	<u>A091454 001</u>	Aug 26, 2013
<u>AB</u>		JUNEL FE 1/20	<u>0.02MG; 1.5MG</u>	<u>N017355 001</u>	
<u>AB</u>		LARIN FE 1.5/30	<u>0.03MG; 1.5MG</u>	<u>N017354 001</u>	
<u>AB</u>		NOVAST LABS LTD	<u>0.02MG; 1.5MG</u>	<u>A075548 001</u>	Feb 05, 2001
<u>AB</u>		LARIN FE 1/20	<u>0.02MG; 1MG</u>	<u>A075647 001</u>	Feb 05, 2001
<u>AB</u>		NOVAST LABS LTD	<u>0.02MG; 1.5MG</u>	<u>A076629 001</u>	Mar 18, 2010
<u>AB</u>		LOESTRIN FE 1.5/30	<u>0.03MG; 1.5MG</u>	<u>A202772 001</u>	Nov 14, 2013
<u>AB</u>	+!	APIL	<u>0.02MG; 1MG</u>	<u>A202741 001</u>	Feb 20, 2015
<u>AB</u>		LOESTRIN FE 1/20	<u>0.02MG; 1.5MG</u>	<u>A076105 001</u>	Oct 26, 2007
<u>AB</u>		MICROGESTIN FE 1.5/30	<u>0.02MG; 1MG</u>	<u>A206287 001</u>	May 24, 2016
<u>AB</u>		MAYNE PHARMA	<u>0.03MG; 1.5MG</u>	<u>N203667 001</u>	May 08, 2013
<u>AB</u>		MICROGESTIN FE 1/20	<u>0.02MG; 1.5MG</u>	<u>A207514 001</u>	Sep 11, 2017
<u>AB</u>		MAYNE PHARMA	<u>0.02MG; 1MG</u>	<u>A210369 001</u>	Dec 26, 2017
		<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL</u>		<u>A206120 001</u>	Sep 12, 2017
<u>AB</u>		MAYNE PHARMA	<u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u>		
<u>AB</u>		MYLAN LABS LTD	<u>0.02MG; 1MG</u>		
<u>AB</u>			<u>0.03MG; 1.5MG</u>		
		<u>TRI-LEGEST FE</u>			
<u>AB</u>		BARR	<u>0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG</u>		
		TABLET, CHEWABLE;ORAL			
		<u>MIBELAS 24 FE</u>			
<u>AB</u>		LUPIN ATLANTIS	<u>0.02MG; 1MG</u>		
		<u>MINASTRIN 24 FE</u>			
<u>AB</u>	+!	APIL	<u>0.02MG; 1MG</u>		
		<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>			
<u>AB</u>		AMNEAL PHARMS	<u>0.02MG; 1MG</u>		
<u>AB</u>		GLENMARK PHARMS LTD	<u>0.02MG; 1MG</u>		
<u>AB</u>		MYLAN LABS LTD	<u>0.02MG; 1MG</u>		
		TABLET, CHEWABLE, TABLET;ORAL			
		LO MINASTRIN FE			
	+!	APIL	<u>0.01MG, 0.01MG, N/A, 1MG, N/A, N/A</u>		
				<u>N204654 001</u>	Jul 24, 2013

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET;ORAL

NORGESTIMATE AND ETHINYL ESTRADIOL

<u>AB</u>		GLENMARK GENERICS	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200494 001</u>	Jun 17, 2011
		TABLET;ORAL-28			
		<u>ESTARYLLA</u>			
<u>AB</u>		LABS LEON FARMA	<u>0.035MG; 0.25MG</u>	<u>A090794 001</u>	Jan 30, 2013
		<u>MILI</u>			
<u>AB</u>		AUROBINDO PHARMA LTD	<u>0.035MG; 0.25MG</u>	<u>A205449 001</u>	Jul 07, 2016
		<u>MONO-LINYAH</u>			
<u>AB</u>		NOVAST LABS LTD	<u>0.035MG; 0.25MG</u>	<u>A090523 001</u>	May 23, 2012
		<u>NORGESTIMATE AND ETHINYL ESTRADIOL</u>			
<u>AB</u>		AMNEAL PHARMS	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A203870 001</u>	Nov 12, 2015
			<u>0.035MG; 0.25MG</u>	<u>A203865 001</u>	Oct 27, 2015
<u>AB</u>			<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A203873 001</u>	May 12, 2016
<u>AB</u>		GLENMARK GENERICS	<u>0.035MG; 0.25MG</u>	<u>A200538 001</u>	Apr 05, 2012
<u>AB</u>		GLENMARK PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A204057 001</u>	Feb 23, 2016
<u>AB</u>		LUPIN LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205588 001</u>	Apr 26, 2016
<u>AB</u>			<u>0.035MG; 0.25MG</u>	<u>A205630 001</u>	Oct 27, 2016
<u>AB</u>		LUPIN PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A200541 001</u>	Jun 25, 2012
<u>AB</u>		MYLAN LABS LTD	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A202132 001</u>	Sep 09, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-170 (of 436)

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET;ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

<u>AB</u>		<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A201897 001</u>	Jan 27, 2016	
<u>AB</u>		<u>0.035MG; 0.25MG</u>	<u>A201896 001</u>	Jan 27, 2016	
<u>AB</u>	OC PHARMA	<u>0.035MG; 0.035MG; 0.035MG; 0.18MG; 0.215MG; 0.25MG</u>	<u>A200383 001</u>	Apr 07, 2015	
<u>AB</u>		<u>0.035MG; 0.25MG</u>	<u>A200384 001</u>	Apr 07, 2015	
		<u><b>ORTHO CYCLEN-28</b></u>			
<u>AB</u>	+!	JANSSEN PHARMS	<u>0.035MG; 0.25MG</u>	<u>N019653 002</u>	Dec 29, 1989
		<u><b>ORTHO TRI-CYCLEN</b></u>			
<u>AB</u>	+!	JANSSEN PHARMS	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>N019697 001</u>	Jul 03, 1992
		<u><b>ORTHO TRI-CYCLEN LO</b></u>			
<u>AB</u>	+!	JANSSEN PHARMS	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>N021241 001</u>	Aug 22, 2002
		<u><b>PREVIFEM</b></u>			
<u>AB</u>	VINTAGE PHARMS LLC	<u>0.035MG; 0.25MG</u>	<u>A076334 001</u>	Jan 09, 2004	
		<u><b>SPRINTEC</b></u>			
<u>AB</u>	BARR	<u>0.035MG; 0.25MG</u>	<u>A075804 001</u>	Sep 25, 2002	
		<u><b>TRI LO SPRINTEC</b></u>			
<u>AB</u>	BARR LABS INC	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076784 001</u>	Jun 29, 2009	
		<u><b>TRI-ESTARYLLA</b></u>			
<u>AB</u>	LABS LEON FARMA	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090793 001</u>	Jan 30, 2013	
		<u><b>TRI-LINYAH</b></u>			
<u>AB</u>	NOVAST LABS LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A090524 001</u>	May 30, 2012	
		<u><b>TRI-LO-ESTARYLLA</b></u>			
<u>AB</u>	LABS LEON FARMA	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A091232 001</u>	Jun 29, 2015	
		<u><b>TRI-LO-MILI</b></u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.025MG, 0.025MG, 0.025MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205762 001</u>	Nov 04, 2016	
		<u><b>TRI-MILI</b></u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A205441 001</u>	Jul 06, 2016	
		<u><b>TRI-PREVIFEM</b></u>			
<u>AB</u>	VINTAGE PHARMS LLC	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A076335 001</u>	Mar 26, 2004	
		<u><b>TRI-SPRINTEC</b></u>			
<u>AB</u>	BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>A075808 001</u>	Dec 29, 2003	

ETHINYL ESTRADIOL; NORGESTREL

TABLET;ORAL-21

CRYSELLE

<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 001</u>	Nov 30, 2001
		<u>TABLET;ORAL-28</u>		
<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>A075840 002</u>	Nov 30, 2001
		<u><b>ELINEST</b></u>		
<u>AB</u>	NOVAST LABS LTD	<u>0.03MG; 0.3MG</u>	<u>A091105 001</u>	Mar 28, 2012
		<u><b>LOW-OGESTREL-28</b></u>		
<u>AB</u>	MAYNE PHARMA OGESTREL 0.5/50-28 ! WATSON LABS	<u>0.03MG; 0.3MG</u>	<u>A075288 002</u>	Jul 28, 1999
		0.05MG; 0.5MG		A075406 002 Dec 15, 1999

ETHIODIZED OIL

OIL;INTRALYMPHATIC, INTRAUTERINE

LIPIODOL

+! GUERBET

EQ 4.8GM IODINE/10ML (EQ 480MG IODINE/ML)

N009190 001

ETHIONAMIDE

TABLET;ORAL

TRECATOR

+! WYETH PHARMS INC

250MG

N013026 002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-171 (of 436)

**ETHOSUXIMIDE**

CAPSULE; ORAL

**ETHOSUXIMIDE**

<b>AB</b>	AKORN	<b>250MG</b>	<b>A040686 001</b>	May 28, 2008
<b>AB</b>	BIONPHARMA INC	<b>250MG</b>	<b>A040430 001</b>	Oct 28, 2002
<b>AB</b>	HERITAGE PHARMS INC	<b>250MG</b>	<b>A200892 001</b>	Sep 25, 2012

**ZARONTIN**

<b>AB</b>	+!	PARKER DAVIS	<b>250MG</b>	<b>N012380 001</b>
-----------	----	--------------	--------------	--------------------

SYRUP; ORAL

**ETHOSUXIMIDE**

<b>AA</b>	MIKART	<b>250MG/5ML</b>	<b>A040506 001</b>	Dec 22, 2003
<b>AA</b>	PHARM ASSOC	<b>250MG/5ML</b>	<b>A040253 001</b>	Nov 22, 2000
<b>AA</b>	TEVA PHARMS	<b>250MG/5ML</b>	<b>A081306 001</b>	Jul 30, 1993

**ZARONTIN**

<b>AA</b>	!	PARKER-DAVIS	<b>250MG/5ML</b>	<b>A080258 001</b>
-----------	---	--------------	------------------	--------------------

**ETHOTOIN**

TABLET; ORAL

PEGANONE

+!	RECORDATI RARE	250MG
----	----------------	-------

N010841 001

**ETIDRONATE DISODIUM**

TABLET; ORAL

ETIDRONATE DISODIUM

MYLAN	200MG	A075800 001	Jan 24, 2003
!	400MG	A075800 002	Jan 24, 2003

**ETODOLAC**

CAPSULE; ORAL

**ETODOLAC**

<b>AB</b>	ANI PHARMS INC	<b>200MG</b>	<b>A075126 001</b>	Sep 16, 1999
<b>AB</b>		<b>300MG</b>	<b>A075126 002</b>	Sep 16, 1999
<b>AB</b>	APOTEX	<b>200MG</b>	<b>A075419 001</b>	Jul 28, 2000
<b>AB</b>		<b>300MG</b>	<b>A075419 002</b>	Jul 28, 2000
<b>AB</b>	TARO	<b>200MG</b>	<b>A075078 001</b>	Apr 30, 1998
<b>AB</b>	!	<b>300MG</b>	<b>A075078 002</b>	Apr 30, 1998

TABLET; ORAL

**ETODOLAC**

<b>AB</b>	APOTEX INC	<b>400MG</b>	<b>A076004 001</b>	Dec 03, 2002
<b>AB</b>		<b>500MG</b>	<b>A076004 002</b>	Dec 03, 2002
<b>AB</b>	SANDOZ	<b>400MG</b>	<b>A074903 001</b>	Apr 11, 1997
<b>AB</b>		<b>500MG</b>	<b>A074903 002</b>	Apr 19, 1999
<b>AB</b>	TARO PHARM INDS	<b>400MG</b>	<b>A075074 001</b>	Mar 11, 1998
<b>AB</b>	!	<b>500MG</b>	<b>A075074 002</b>	Apr 25, 2000
<b>AB</b>	TEVA	<b>400MG</b>	<b>A075009 001</b>	Nov 26, 1997
<b>AB</b>		<b>500MG</b>	<b>A075009 002</b>	Dec 28, 1999

TABLET, EXTENDED RELEASE; ORAL

**ETODOLAC**

<b>AB</b>	TARO	<b>400MG</b>	<b>A076174 001</b>	Mar 13, 2003
<b>AB</b>		<b>500MG</b>	<b>A076174 002</b>	Mar 13, 2003
<b>AB</b>		<b>600MG</b>	<b>A076174 003</b>	Mar 13, 2003
<b>AB</b>	TEVA	<b>400MG</b>	<b>A075665 003</b>	Feb 05, 2001
<b>AB</b>		<b>500MG</b>	<b>A075665 002</b>	Jul 31, 2000
<b>AB</b>	!	<b>600MG</b>	<b>A075665 001</b>	Jul 31, 2000
<b>AB</b>	ZYDUS PHARMS USA INC	<b>400MG</b>	<b>A091134 001</b>	Jan 23, 2014
<b>AB</b>		<b>500MG</b>	<b>A091134 002</b>	Jan 23, 2014
<b>AB</b>		<b>600MG</b>	<b>A091134 003</b>	Jan 23, 2014

**ETOMIDATE**

INJECTABLE; INJECTION

**AMIDATE**

<b>AP</b>	+!	HOSPIRA	<b>2MG/ML</b>	<b>N018227 001</b>	Sep 07, 1982
-----------	----	---------	---------------	--------------------	--------------

**ETOMIDATE**

<b>AP</b>	AUROBINDO PHARMA LTD	<b>2MG/ML</b>	<b>A206126 001</b>	Feb 24, 2017
<b>AP</b>	EMCURE PHARMS LTD	<b>2MG/ML</b>	<b>A204618 001</b>	Aug 13, 2014
<b>AP</b>	GLAND PHARMA LTD	<b>2MG/ML</b>	<b>A209058 001</b>	Apr 18, 2017
<b>AP</b>	HIKMA FARMACEUTICA	<b>2MG/ML</b>	<b>A202354 001</b>	Feb 25, 2016
<b>AP</b>	LUITPOLD	<b>2MG/ML</b>	<b>A078867 001</b>	Dec 22, 2009
<b>AP</b>	MYLAN LABS LTD	<b>2MG/ML</b>	<b>A078289 001</b>	Jan 02, 2009
<b>AP</b>		<b>2MG/ML</b>	<b>A201044 001</b>	Feb 07, 2017
<b>AP</b>	PAR STERILE PRODUCTS	<b>2MG/ML</b>	<b>A091297 001</b>	Jun 20, 2012
<b>AP</b>	WEST-WARD PHARMS	<b>2MG/ML</b>	<b>A074593 001</b>	Nov 04, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-172 (of 436)

ETOMIDATE

INJECTABLE; INJECTION

ETOMIDATE

INT

AP ZYDUS PHARMS USA INC

2MG/ML

A202360 001 Jul 18, 2014

ETONOGESTREL

IMPLANT; IMPLANTATION

NEXPLANON

+! ORGANON USA INC

68MG/IMPLANT

N021529 002 May 31, 2011

ETOPOSIDE

CAPSULE; ORAL

ETOPOSIDE

! MYLAN

50MG

A075635 001 Sep 19, 2001

INJECTABLE; INJECTION

ETOPOSIDE

AP ACCORD HLTHCARE

20MG/ML

A074513 001 Mar 14, 1996

AP ! FRESENIUS KABI USA

20MG/ML

A074983 001 Sep 30, 1998

AP MYLAN LABS LTD

20MG/ML

A203507 001 Nov 20, 2017

AP

20MG/ML

A204927 001 Oct 31, 2017

AP TEVA PHARMS USA

20MG/ML

A074529 001 Jul 24, 1996

AP WEST-WARD PHARMS

20MG/ML

A074290 001 Jul 17, 1995

INT

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

+! BRISTOL MYERS SQUIBB

EQ 100MG BASE/VIAL

N020457 001 May 17, 1996

ETRAVIRINE

TABLET; ORAL

INTELENCE

+

25MG

N022187 003 Mar 26, 2012

+

100MG

N022187 001 Jan 18, 2008

+!

200MG

N022187 002 Dec 22, 2010

EVEROLIMUS

TABLET; ORAL

AFINITOR

+

2.5MG

N022334 003 Jul 09, 2010

+

5MG

N022334 001 Mar 30, 2009

+

7.5MG

N022334 004 Mar 30, 2012

+!

10MG

N022334 002 Mar 30, 2009

ZORTRESS

+

0.25MG

N021560 001 Apr 20, 2010

+

0.5MG

N021560 002 Apr 20, 2010

+!

0.75MG

N021560 003 Apr 20, 2010

TABLET, FOR SUSPENSION; ORAL

AFINITOR DISPERZ

+

2MG

N203985 001 Aug 29, 2012

+

3MG

N203985 002 Aug 29, 2012

+!

5MG

N203985 003 Aug 29, 2012

EXEMESTANE

TABLET; ORAL

AROMASIN

AB +! PHARMACIA AND UPJOHN

25MG

N020753 001 Oct 21, 1999

EXEMESTANE

AB ALVOGEN MALTA

25MG

A200898 001 Jul 28, 2014

AB MYLAN PHARMS INC

25MG

A203315 001 Mar 10, 2017

AB UPSHER-SMITH LABS

25MG

A209208 001 Jul 26, 2017

AB WEST-WARD PHARMS

25MG

A077431 001 Apr 01, 2011

INT

EXENATIDE

SUSPENSION, EXTENDED RELEASE; SUBCUTANEOUS

BYDUREON BCISE

+! ASTRazeneca AB

2MG/0.85ML (2MG/0.85ML)

N209210 001 Oct 20, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-173 (of 436)

**EXENATIDE SYNTHETIC**

FOR SUSPENSION, EXTENDED RELEASE;SUBCUTANEOUS

BYDUREON

+! ASTRAZENECA AB 2MG/VIAL

N022200 001 Jan 27, 2012

BYDUREON PEN

+! ASTRAZENECA AB 2MG

N022200 002 Feb 28, 2014

INJECTABLE;SUBCUTANEOUS

BYETTA

+! ASTRAZENECA AB 300MCG/1.2ML (250MCG/ML)

N021773 001 Apr 28, 2005

+! ASTRAZENECA AB 600MCG/2.4ML (250MCG/ML)

N021773 002 Apr 28, 2005

**EZETIMIBE**

TABLET;ORAL

**EZETIMIBE**

<b><u>AB</u></b>	ALKEM LABS LTD	<b><u>10MG</u></b>	<b><u>A209234 001</u></b>	Dec 21, 2017
<b><u>AB</u></b>	AMNEAL PHARMS CO	<b><u>10MG</u></b>	<b><u>A208803 001</u></b>	Jun 12, 2017
<b><u>AB</u></b>	APOTEX INC	<b><u>10MG</u></b>	<b><u>A208332 001</u></b>	Jun 12, 2017
<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>10MG</u></b>	<b><u>A209838 001</u></b>	Aug 25, 2017
<b><u>AB</u></b>	GLENMARK PHARMS LTD	<b><u>10MG</u></b>	<b><u>A078560 001</u></b>	Jun 26, 2015
<b><u>AB</u></b>	OHM LABS INC	<b><u>10MG</u></b>	<b><u>A207311 001</u></b>	Jun 12, 2017
<b><u>AB</u></b>	SANDOZ INC	<b><u>10MG</u></b>	<b><u>A203931 001</u></b>	Jun 12, 2017
<b><u>AB</u></b>	TEVA PHARMS USA	<b><u>10MG</u></b>	<b><u>A078724 001</u></b>	Jun 12, 2017
<b><u>AB</u></b>	WATSON LABS INC	<b><u>10MG</u></b>	<b><u>A200831 001</u></b>	Jun 12, 2017
<b><u>AB</u></b>	ZYDUS PHARMS USA INC	<b><u>10MG</u></b>	<b><u>A204331 001</u></b>	Jun 12, 2017

**ZETIA**

<b><u>AB</u></b>	+! MSD INTL GMBH	<b><u>10MG</u></b>	<b><u>N021445 001</u></b>	Oct 25, 2002
------------------	------------------	--------------------	---------------------------	--------------

**EZETIMIBE; SIMVASTATIN**

TABLET;ORAL

**EZETIMIBE AND SIMVASTATIN**

<b><u>AB</u></b>	ALKEM LABS LTD	<b><u>10MG;10MG</u></b>	<b><u>A209222 001</u></b>	Dec 22, 2017
<b><u>AB</u></b>		<b><u>10MG;20MG</u></b>	<b><u>A209222 002</u></b>	Dec 22, 2017
<b><u>AB</u></b>		<b><u>10MG;40MG</u></b>	<b><u>A209222 003</u></b>	Dec 22, 2017
<b><u>AB</u></b>		<b><u>10MG;80MG</u></b>	<b><u>A209222 004</u></b>	Dec 22, 2017
<b><u>AB</u></b>	AMNEAL PHARMS CO	<b><u>10MG;10MG</u></b>	<b><u>A208831 001</u></b>	Nov 21, 2017
<b><u>AB</u></b>		<b><u>10MG;20MG</u></b>	<b><u>A208831 002</u></b>	Nov 21, 2017
<b><u>AB</u></b>		<b><u>10MG;40MG</u></b>	<b><u>A208831 003</u></b>	Nov 21, 2017
<b><u>AB</u></b>		<b><u>10MG;80MG</u></b>	<b><u>A208831 004</u></b>	Nov 21, 2017
<b><u>AB</u></b>	DR REDDYS LABS SA	<b><u>10MG;10MG</u></b>	<b><u>A200909 001</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;20MG</u></b>	<b><u>A200909 002</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;40MG</u></b>	<b><u>A200909 003</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;80MG</u></b>	<b><u>A200909 004</u></b>	Apr 26, 2017
<b><u>AB</u></b>	IMPAX LABS INC	<b><u>10MG;10MG</u></b>	<b><u>A201890 001</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;20MG</u></b>	<b><u>A201890 002</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;40MG</u></b>	<b><u>A201890 003</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;80MG</u></b>	<b><u>A201890 004</u></b>	Apr 26, 2017
<b><u>AB</u></b>	WATSON LABS INC	<b><u>10MG;10MG</u></b>	<b><u>A202968 001</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;20MG</u></b>	<b><u>A202968 002</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;40MG</u></b>	<b><u>A202968 003</u></b>	Apr 26, 2017
<b><u>AB</u></b>		<b><u>10MG;80MG</u></b>	<b><u>A202968 004</u></b>	Apr 26, 2017

**VYTORIN**

<b><u>AB</u></b>	+ MSD INTL	<b><u>10MG;10MG</u></b>	<b><u>N021687 001</u></b>	Jul 23, 2004
<b><u>AB</u></b>	+	<b><u>10MG;20MG</u></b>	<b><u>N021687 002</u></b>	Jul 23, 2004
<b><u>AB</u></b>	+	<b><u>10MG;40MG</u></b>	<b><u>N021687 003</u></b>	Jul 23, 2004
<b><u>AB</u></b>	!+	<b><u>10MG;80MG</u></b>	<b><u>N021687 004</u></b>	Jul 23, 2004

**FAMCICLOVIR**

TABLET;ORAL

**FAMCICLOVIR**

<b><u>AB</u></b>	APOTEX	<b><u>125MG</u></b>	<b><u>A091480 001</u></b>	Jul 22, 2011
<b><u>AB</u></b>		<b><u>250MG</u></b>	<b><u>A091480 002</u></b>	Jul 22, 2011
<b><u>AB</u></b>		<b><u>500MG</u></b>	<b><u>A091480 003</u></b>	Jul 22, 2011
<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>125MG</u></b>	<b><u>A091114 001</u></b>	Mar 21, 2011
<b><u>AB</u></b>		<b><u>250MG</u></b>	<b><u>A091114 002</u></b>	Mar 21, 2011
<b><u>AB</u></b>		<b><u>500MG</u></b>	<b><u>A091114 003</u></b>	Mar 21, 2011
<b><u>AB</u></b>	CIPLA LTD	<b><u>125MG</u></b>	<b><u>A078278 001</u></b>	Mar 21, 2011
<b><u>AB</u></b>		<b><u>250MG</u></b>	<b><u>A078278 002</u></b>	Mar 21, 2011
<b><u>AB</u></b>		<b><u>500MG</u></b>	<b><u>A078278 003</u></b>	Mar 21, 2011
<b><u>AB</u></b>	HETERO LABS LTD V	<b><u>125MG</u></b>	<b><u>A202438 001</u></b>	Sep 10, 2014
<b><u>AB</u></b>		<b><u>250MG</u></b>	<b><u>A202438 002</u></b>	Sep 10, 2014
<b><u>AB</u></b>		<b><u>500MG</u></b>	<b><u>A202438 003</u></b>	Sep 10, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-174 (of 436)

**FAMCICLOVIR**

TABLET;ORAL

**FAMCICLOVIR**

<b>AB</b>	MACLEODS PHARMS LTD	<u>125MG</u>	<b>A201022 001</b>	Jan 12, 2012
<b>AB</b>		<u>250MG</u>	<b>A201022 002</b>	Jan 12, 2012
<b>AB</b>		<u>500MG</u>	<b>A201022 003</b>	Jan 12, 2012
<b>AB</b>	MYLAN	<u>125MG</u>	<b>A201333 001</b>	Mar 24, 2011
<b>AB</b>		<u>250MG</u>	<b>A201333 002</b>	Mar 24, 2011
<b>AB</b>		<u>500MG</u>	<b>A201333 003</b>	Mar 24, 2011
<b>AB</b>	TEVA PHARMS	<u>125MG</u>	<b>A077487 001</b>	Aug 24, 2007
<b>AB</b>		<u>250MG</u>	<b>A077487 002</b>	Aug 24, 2007
<b>AB</b>	!	<u>500MG</u>	<b>A077487 003</b>	Aug 24, 2007
<b>AB</b>	WEST-WARD PHARMS INT	<u>125MG</u>	<b>A090128 001</b>	Mar 21, 2011
<b>AB</b>		<u>250MG</u>	<b>A090128 002</b>	Mar 21, 2011
<b>AB</b>		<u>500MG</u>	<b>A090128 003</b>	Mar 21, 2011

**FAMOTIDINE**

FOR SUSPENSION;ORAL

**FAMOTIDINE**

<b>AB</b>	HI-TECH PHARMA CO	<u>40MG/5ML</u>	<b>A201995 001</b>	May 30, 2014
<b>AB</b>	LUPIN LTD	<u>40MG/5ML</u>	<b>A090440 001</b>	Jun 29, 2010
<b>AB</b>	NAVINTA LLC	<u>40MG/5ML</u>	<b>A091020 001</b>	May 27, 2010
<b>AB</b>	NOVEL LABS INC	<u>40MG/5ML</u>	<b>A201695 001</b>	Dec 17, 2012

**PEPCID**

<b>AB</b>	+!	SALIX PHARMS	<u>40MG/5ML</u>	<b>N019527 001</b>	Feb 02, 1987
-----------	----	--------------	-----------------	--------------------	--------------

INJECTABLE;INJECTION

**FAMOTIDINE**

<b>AP</b>	ATHENEX INC	<u>10MG/ML</u>	<b>A075651 001</b>	Apr 16, 2001	
<b>AP</b>		<u>10MG/ML</u>	<b>A075684 001</b>	Apr 16, 2001	
<b>AP</b>	FRESENIUS KABI USA	<u>10MG/ML</u>	<b>A075709 001</b>	Apr 16, 2001	
<b>AP</b>	MYLAN LABS LTD	<u>10MG/ML</u>	<b>A078641 001</b>	Jun 25, 2008	
<b>AP</b>	!	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<b>A075488 001</b>	Apr 16, 2001

**FAMOTIDINE PRESERVATIVE FREE**

<b>AP</b>	ATHENEX INC	<u>10MG/ML</u>	<b>A075622 001</b>	Apr 16, 2001	
<b>AP</b>		<u>10MG/ML</u>	<b>A075825 001</b>	Apr 17, 2001	
<b>AP</b>	FRESENIUS KABI USA	<u>10MG/ML</u>	<b>A075813 001</b>	Apr 16, 2001	
<b>AP</b>	MYLAN LABS LTD	<u>10MG/ML</u>	<b>A078642 001</b>	Jun 25, 2008	
<b>AP</b>	!	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<b>A075486 001</b>	Apr 16, 2001

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

! BAXTER HLTHCARE 0.4MG/ML

A075591 001 May 10, 2001

TABLET;ORAL

**FAMOTIDINE**

<b>AB</b>	ALEMBIC PHARMS LTD	<u>20MG</u>	<b>A078916 001</b>	May 22, 2009
<b>AB</b>		<u>40MG</u>	<b>A078916 002</b>	May 22, 2009
<b>AB</b>	APOTEX	<u>20MG</u>	<b>A075611 001</b>	Jul 23, 2001
<b>AB</b>		<u>40MG</u>	<b>A075611 002</b>	Jul 23, 2001
<b>AB</b>	AUROBINDO PHARMA LTD	<u>20MG</u>	<b>A206530 001</b>	Dec 22, 2015
<b>AB</b>		<u>40MG</u>	<b>A206530 002</b>	Dec 22, 2015
<b>AB</b>	CARLSBAD	<u>20MG</u>	<b>A075805 001</b>	Apr 16, 2001
<b>AB</b>		<u>40MG</u>	<b>A075805 002</b>	Apr 16, 2001
<b>AB</b>	DR REDDYS LABS LTD	<u>20MG</u>	<b>A075718 001</b>	Apr 16, 2001
<b>AB</b>		<u>40MG</u>	<b>A075718 002</b>	Apr 16, 2001
<b>AB</b>	IVAX SUB TEVA PHARMS	<u>20MG</u>	<b>A075511 001</b>	Apr 16, 2001
<b>AB</b>		<u>40MG</u>	<b>A075511 002</b>	Apr 16, 2001
<b>AB</b>	MYLAN	<u>20MG</u>	<b>A075704 001</b>	Apr 16, 2001
<b>AB</b>		<u>40MG</u>	<b>A075704 002</b>	Apr 16, 2001
<b>AB</b>	PERRIGO R AND D	<u>20MG</u>	<b>A077352 002</b>	Jul 27, 2005
<b>AB</b>		<u>40MG</u>	<b>A077352 001</b>	Jul 27, 2005
<b>AB</b>	TEVA	<u>20MG</u>	<b>A075311 001</b>	Apr 16, 2001
<b>AB</b>		<u>40MG</u>	<b>A075311 002</b>	Apr 16, 2001
<b>AB</b>	WOCKHARDT LTD	<u>20MG</u>	<b>A075786 001</b>	Apr 16, 2001
<b>AB</b>		<u>40MG</u>	<b>A075786 002</b>	Apr 16, 2001

**PEPCID**

<b>AB</b>	+	VALEANT PHARMS NORTH	<u>20MG</u>	<b>N019462 001</b>	Oct 15, 1986
<b>AB</b>	+		<u>40MG</u>	<b>N019462 002</b>	Oct 15, 1986

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-175 (of 436)

FAMOTIDINE; IBUPROFEN

TABLET;ORAL

DUEXIS

+! HORIZON PHARMA 26.6MG; 800MG

N022519 001 Apr 23, 2011

FEBUXOSTAT

TABLET;ORAL

ULORIC

+ TAKEDA PHARMS USA 40MG  
+! 80MG

N021856 001 Feb 13, 2009  
N021856 002 Feb 13, 2009

FELBAMATE

SUSPENSION;ORAL

**FELBAMATE**

AB AMNEAL PHARMS 600MG/5ML

AB TARO PHARM 600MG/5ML

**FELBATOL**

AB +! MYLAN SPECIALITY LP 600MG/5ML

TABLET;ORAL

**FELBAMATE**

AB ALVOGEN MALTA 400MG

AB 600MG

AB AMNEAL PHARMS 400MG

AB 600MG

AB IMPAX LABS INC 400MG

AB 600MG

AB TARO PHARM 400MG

AB 600MG

AB ZYDUS PHARMS USA 400MG

INC

AB 600MG

A202385 001 Dec 16, 2011

A206314 001 Jun 16, 2017

**FELBATOL**

AB + MYLAN SPECIALITY LP 400MG

AB +! 600MG

N020189 003 Jul 29, 1993

**FELDIPINE**

TABLET, EXTENDED RELEASE;ORAL

**FELODIPINE**

AB AUROBINDO PHARMA 2.5MG

LTD

AB 5MG

AB 10MG

AB GLENMARK GENERICS 2.5MG

AB 5MG

AB 10MG

AB HERITAGE PHARMS INC 2.5MG

AB 5MG

AB 10MG

AB JUBILANT GENERICS 2.5MG

AB 5MG

AB 10MG

AB MYLAN 2.5MG

AB 5MG

AB ! 10MG

AB ORCHID HLTHCARE 2.5MG

AB 5MG

AB 10MG

AB SUN PHARM INDs LTD 2.5MG

AB 5MG

AB 10MG

AB SUN PHARM INDUSTRIES 2.5MG

AB 5MG

AB 10MG

AB TORRENT PHARMS LTD 2.5MG

AB 5MG

AB 10MG

AB VINTAGE PHARMS LLC 2.5MG

AB 5MG

AB 10MG

A203417 001 Jan 17, 2013

A203417 002 Jan 17, 2013

A203417 003 Jan 17, 2013

A090365 001 Dec 17, 2010

A090365 002 Dec 17, 2010

A090365 003 Dec 17, 2010

A201964 001 Nov 08, 2013

A201964 002 Nov 08, 2013

A201964 003 Nov 08, 2013

A203983 001 Aug 19, 2016

A203983 002 Aug 19, 2016

A203983 003 Aug 19, 2016

A078855 001 Apr 17, 2008

A078855 002 Apr 17, 2008

A078855 003 Apr 17, 2008

A203032 001 May 21, 2015

A203032 002 May 21, 2015

A203032 003 May 21, 2015

A091200 001 Dec 13, 2013

A091200 002 Dec 13, 2013

A091200 003 Dec 13, 2013

A075896 001 Nov 02, 2004

A075896 002 Nov 02, 2004

A075896 003 Nov 02, 2004

A202170 001 Nov 28, 2011

A202170 002 Nov 28, 2011

A202170 003 Nov 28, 2011

A200815 001 Oct 28, 2011

A200815 002 Oct 28, 2011

A200815 003 Oct 28, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-176 (of 436)

**FENOFIBRATE**

CAPSULE; ORAL

**ANTARA (MICRONIZED)**

<b>AB</b>	+	LUPIN ATLANTIS	<b>43MG</b>	<b>N021695 001</b>	Nov 30, 2004
<b>AB</b>	!+		<b>130MG</b>	<b>N021695 003</b>	Nov 30, 2004

**FENOFIBRATE**

<b>AB</b>		SUN PHARM INDs LTD	<b>43MG</b>	<b>A201748 001</b>	Oct 31, 2014
<b>AB</b>			<b>130MG</b>	<b>A201748 002</b>	Oct 31, 2014

**FENOFIBRATE (MICRONIZED)**

<b>AB</b>		APOTEX INC	<b>43MG</b>	<b>A202252 001</b>	Jul 26, 2013
<b>AB</b>			<b>130MG</b>	<b>A202252 002</b>	Jul 26, 2013
<b>AB</b>		CNTY LINE PHARMS	<b>67MG</b>	<b>A207805 001</b>	Nov 16, 2017
<b>AB</b>			<b>134MG</b>	<b>A207805 002</b>	Nov 16, 2017
<b>AB</b>			<b>200MG</b>	<b>A207805 003</b>	Nov 16, 2017
<b>AB</b>		DR REDDYS LABS SA	<b>43MG</b>	<b>A090859 001</b>	Mar 01, 2012
<b>AB</b>			<b>130MG</b>	<b>A090859 002</b>	Mar 01, 2012
<b>AB</b>		GLENMARK PHARMS LTD	<b>67MG</b>	<b>A205566 001</b>	Apr 07, 2017
<b>AB</b>			<b>134MG</b>	<b>A205566 002</b>	Apr 07, 2017
<b>AB</b>			<b>200MG</b>	<b>A205566 003</b>	Apr 07, 2017
<b>AB</b>		IMPAK LABS	<b>67MG</b>	<b>A075868 001</b>	Oct 27, 2003
<b>AB</b>			<b>134MG</b>	<b>A075868 002</b>	Oct 27, 2003
<b>AB</b>	!		<b>200MG</b>	<b>A075868 003</b>	Oct 27, 2003
<b>AB</b>		INVAGEN PHARMS	<b>67MG</b>	<b>A207378 001</b>	Mar 28, 2017
<b>AB</b>			<b>134MG</b>	<b>A207378 002</b>	Mar 28, 2017
<b>AB</b>			<b>200MG</b>	<b>A207378 003</b>	Mar 28, 2017
<b>AB</b>		MYLAN PHARMS INC	<b>43MG</b>	<b>A202579 001</b>	Jan 10, 2013
<b>AB</b>			<b>67MG</b>	<b>A202676 001</b>	Oct 23, 2012
<b>AB</b>			<b>130MG</b>	<b>A202579 002</b>	Jan 10, 2013
<b>AB</b>			<b>134MG</b>	<b>A202676 002</b>	Oct 23, 2012
<b>AB</b>			<b>200MG</b>	<b>A202676 003</b>	Oct 23, 2012
<b>AB</b>		RHODES PHARMS	<b>67MG</b>	<b>A075753 001</b>	Sep 03, 2002
<b>AB</b>			<b>134MG</b>	<b>A075753 002</b>	Apr 09, 2002
<b>AB</b>			<b>200MG</b>	<b>A075753 003</b>	Apr 09, 2002

ANTARA (MICRONIZED)

+	LUPIN ATLANTIS	30MG	N021695 004	Oct 18, 2013
+		90MG	N021695 005	Oct 18, 2013

LIPOFEN

+	CIPHER PHARMS INC	50MG	N021612 001	Jan 11, 2006
!+		150MG	N021612 003	Jan 11, 2006

TABLET; ORAL

**FENOFIBRATE**

<b>AB</b>		AUROBINDO PHARMA LTD	<b>48MG</b>	<b>A205118 001</b>	May 05, 2016
<b>AB</b>			<b>145MG</b>	<b>A205118 002</b>	May 05, 2016
<b>AB</b>		CIPLA LTD	<b>48MG</b>	<b>A208709 001</b>	Dec 15, 2016
<b>AB</b>			<b>145MG</b>	<b>A208709 002</b>	Dec 15, 2016
<b>AB</b>		CNTY LINE PHARMS	<b>54MG</b>	<b>A207803 001</b>	Dec 19, 2017
<b>AB</b>			<b>160MG</b>	<b>A207803 002</b>	Dec 19, 2017
<b>AB</b>		HETERO LABS LTD III	<b>48MG</b>	<b>A204598 001</b>	Jul 12, 2016
<b>AB</b>			<b>145MG</b>	<b>A204598 002</b>	Jul 12, 2016
<b>AB</b>		IMPAX LABS	<b>54MG</b>	<b>A076509 001</b>	Mar 26, 2008
<b>AB</b>	!		<b>160MG</b>	<b>A076509 002</b>	Mar 26, 2008
<b>AB</b>		LUPIN LTD	<b>48MG</b>	<b>A090856 001</b>	Dec 23, 2011
<b>AB</b>			<b>54MG</b>	<b>A204019 001</b>	Aug 17, 2015
<b>AB</b>			<b>145MG</b>	<b>A090856 002</b>	Dec 23, 2011
<b>AB</b>			<b>160MG</b>	<b>A204019 002</b>	Aug 17, 2015
<b>AB</b>		MYLAN	<b>40MG</b>	<b>A204475 001</b>	Jun 23, 2016
<b>AB</b>			<b>54MG</b>	<b>A076520 001</b>	Oct 25, 2007
<b>AB</b>			<b>120MG</b>	<b>A204475 002</b>	Jun 23, 2016
<b>AB</b>			<b>160MG</b>	<b>A076520 003</b>	Oct 25, 2007
<b>AB</b>		MYLAN PHARMS INC	<b>48MG</b>	<b>A202856 001</b>	Dec 07, 2012
<b>AB</b>			<b>145MG</b>	<b>A202856 002</b>	Dec 07, 2012
<b>AB</b>		RHODES PHARMS	<b>54MG</b>	<b>A076433 001</b>	May 13, 2005
<b>AB</b>			<b>160MG</b>	<b>A076433 002</b>	May 13, 2005
<b>AB</b>		SUN PHARM INDs LTD	<b>48MG</b>	<b>A200884 001</b>	Sep 07, 2017
<b>AB</b>			<b>54MG</b>	<b>A076635 001</b>	Oct 31, 2005
<b>AB</b>			<b>145MG</b>	<b>A200884 002</b>	Sep 07, 2017
<b>AB</b>			<b>160MG</b>	<b>A076635 003</b>	Oct 31, 2005
<b>AB</b>		VALEANT PHARMS NORTH	<b>48MG</b>	<b>A090715 001</b>	Apr 05, 2012
<b>AB</b>			<b>145MG</b>	<b>A090715 002</b>	Apr 05, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-177 (of 436)

FENOFIBRATE

TABLET;ORAL

FENOGLIDE

<u>AB</u>	+	SANTARUS INC	<u>40MG</u>	<u>N022118 001</u>	Aug 10, 2007
<u>AB</u>	+!		<u>120MG</u>	<u>N022118 002</u>	Aug 10, 2007

TRICOR

<u>AB</u>	+	ABBVIE	<u>48MG</u>	<u>N021656 001</u>	Nov 05, 2004
<u>AB</u>	+!		<u>145MG</u>	<u>N021656 002</u>	Nov 05, 2004

TRIGLIDE

BX	+!	SKYEPHARMA AG	160MG	N021350 002	May 07, 2005
		FENOFIBRATE		A076635 002	Oct 31, 2005
		SUN PHARM INDs LTD	107MG		

FENOFIBRIC ACID

TABLET;ORAL

FIBRICOR

	+	ARALEZ PHARMS INC	35MG	N022418 001	Aug 14, 2009
	+!		105MG	N022418 002	Aug 14, 2009

FENOLDOPAM MESYLATE

INJECTABLE;INJECTION

CORLOPAM

<u>AP</u>	+!	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>N019922 001</u>	Sep 23, 1997
-----------	----	---------	------------------------	--------------------	--------------

FENOLDOPAM MESYLATE

<u>AP</u>		SANDOZ INC	<u>EQ 10MG BASE/ML</u>	<u>A077155 001</u>	Feb 15, 2005
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 10MG BASE/ML</u>	<u>A076582 001</u>	Oct 12, 2004

FENOPROFEN CALCIUM

CAPSULE;ORAL

NALFON

	+	XSPIRE PHARMA	EQ 200MG BASE	N017604 003	
	+		EQ 400MG BASE	N017604 004	Jul 21, 2009

TABLET;ORAL

FENOPROFEN CALCIUM

	!	XSPIRE PHARMA	EQ 600MG BASE	A072267 001	Aug 17, 1988
--	---	---------------	---------------	-------------	--------------

FENTANYL

FILM, EXTENDED RELEASE;TRANSDERMAL

DURAGESIC-100

<u>AB</u>	+	JANSSEN PHARMS	<u>100MCG/HR</u>	<u>N019813 001</u>	Aug 07, 1990
-----------	---	----------------	------------------	--------------------	--------------

DURAGESIC-12

<u>AB</u>	+	JANSSEN PHARMS	<u>12.5MCG/HR</u>	<u>N019813 005</u>	Feb 04, 2005
-----------	---	----------------	-------------------	--------------------	--------------

DURAGESIC-25

<u>AB</u>	+!	JANSSEN PHARMS	<u>25MCG/HR</u>	<u>N019813 004</u>	Aug 07, 1990
-----------	----	----------------	-----------------	--------------------	--------------

DURAGESIC-50

<u>AB</u>	+	JANSSEN PHARMS	<u>50MCG/HR</u>	<u>N019813 003</u>	Aug 07, 1990
-----------	---	----------------	-----------------	--------------------	--------------

DURAGESIC-75

<u>AB</u>	+	JANSSEN PHARMS	<u>75MCG/HR</u>	<u>N019813 002</u>	Aug 07, 1990
-----------	---	----------------	-----------------	--------------------	--------------

FENTANYL-100

<u>AB</u>		3M DRUG DELIVERY	<u>100MCG/HR</u>	<u>A202097 005</u>	Nov 04, 2016
-----------	--	------------------	------------------	--------------------	--------------

ACTAVIS LABS UT INC

<u>AB</u>		ACTAVIS LABS UT INC	<u>100MCG/HR</u>	<u>A076709 004</u>	Aug 20, 2007
-----------	--	---------------------	------------------	--------------------	--------------

AVEVA

<u>AB</u>		AVEVA	<u>100MCG/HR</u>	<u>A077449 004</u>	Oct 20, 2008
-----------	--	-------	------------------	--------------------	--------------

LAVIPHARM LABS

<u>AB</u>		LAVIPHARM LABS	<u>100MCG/HR</u>	<u>A077051 004</u>	Aug 04, 2006
-----------	--	----------------	------------------	--------------------	--------------

MAYNE PHARMA

<u>AB</u>		MAYNE PHARMA	<u>100MCG/HR</u>	<u>A077062 004</u>	Aug 20, 2007
-----------	--	--------------	------------------	--------------------	--------------

MYLAN TECHNOLOGIES

<u>AB</u>		MYLAN TECHNOLOGIES	<u>100MCG/HR</u>	<u>A076258 004</u>	Jan 28, 2005
-----------	--	--------------------	------------------	--------------------	--------------

SPECGX LLC

<u>AB</u>		SPECGX LLC	<u>100MCG/HR</u>	<u>A077154 004</u>	Feb 09, 2011
-----------	--	------------	------------------	--------------------	--------------

FENTANYL-12

<u>AB</u>		3M DRUG DELIVERY	<u>12.5MCG/HR</u>	<u>A202097 001</u>	Nov 04, 2016
-----------	--	------------------	-------------------	--------------------	--------------

AVEVA

<u>AB</u>		AVEVA	<u>12.5MCG/HR</u>	<u>A077449 005</u>	Sep 11, 2015
-----------	--	-------	-------------------	--------------------	--------------

MYLAN TECHNOLOGIES

<u>AB</u>		MYLAN TECHNOLOGIES	<u>12.5MCG/HR</u>	<u>A076258 005</u>	Jan 23, 2007
-----------	--	--------------------	-------------------	--------------------	--------------

SPECGX LLC

<u>AB</u>		SPECGX LLC	<u>12.5MCG/HR</u>	<u>A077154 005</u>	Jun 11, 2015
-----------	--	------------	-------------------	--------------------	--------------

FENTANYL-25

<u>AB</u>		3M DRUG DELIVERY	<u>25MCG/HR</u>	<u>A202097 002</u>	Nov 04, 2016
-----------	--	------------------	-----------------	--------------------	--------------

ACTAVIS LABS UT INC

<u>AB</u>		ACTAVIS LABS UT INC	<u>25MCG/HR</u>	<u>A076709 001</u>	Aug 20, 2007
-----------	--	---------------------	-----------------	--------------------	--------------

AVEVA

<u>AB</u>		AVEVA	<u>25MCG/HR</u>	<u>A077449 001</u>	Oct 20, 2008
-----------	--	-------	-----------------	--------------------	--------------

LAVIPHARM LABS

<u>AB</u>		LAVIPHARM LABS	<u>25MCG/HR</u>	<u>A077051 001</u>	Aug 04, 2006
-----------	--	----------------	-----------------	--------------------	--------------

MAYNE PHARMA

<u>AB</u>		MAYNE PHARMA	<u>25MCG/HR</u>	<u>A077062 001</u>	Aug 20, 2007
-----------	--	--------------	-----------------	--------------------	--------------

MYLAN TECHNOLOGIES

<u>AB</u>		MYLAN TECHNOLOGIES	<u>25MCG/HR</u>	<u>A076258 001</u>	Jan 28, 2005
-----------	--	--------------------	-----------------	--------------------	--------------

SPECGX LLC

<u>AB</u>		SPECGX LLC	<u>25MCG/HR</u>	<u>A077154 001</u>	Feb 09, 2011
-----------	--	------------	-----------------	--------------------	--------------

FENTANYL-37

<u>AB</u>		AVEVA	<u>37.5MCG/HR</u>	<u>A077449 006</u>	Dec 06, 2017
-----------	--	-------	-------------------	--------------------	--------------

FENTANYL-50

<u>AB</u>		3M DRUG DELIVERY	<u>50MCG/HR</u>	<u>A202097 003</u>	Nov 04, 2016
-----------	--	------------------	-----------------	--------------------	--------------

<u>AB</u>		ACTAVIS LABS UT INC	<u>50MCG/HR</u>	<u>A076709 002</u>	Aug 20, 2007
-----------	--	---------------------	-----------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-178 (of 436)

**FENTANYL**

FILM, EXTENDED RELEASE;TRANSDERMAL

**FENTANYL-50**

<b>AB</b>	AVEVA	<b>50MCG/HR</b>	<b>A077449 002</b>	Oct 20, 2008
<b>AB</b>	LAVIPHARM LABS	<b>50MCG/HR</b>	<b>A077051 002</b>	Aug 04, 2006
<b>AB</b>	MAYNE PHARMA	<b>50MCG/HR</b>	<b>A077062 002</b>	Aug 20, 2007
<b>AB</b>	MYLAN TECHNOLOGIES	<b>50MCG/HR</b>	<b>A076258 002</b>	Jan 28, 2005
<b>AB</b>	SPECGX LLC	<b>50MCG/HR</b>	<b>A077154 002</b>	Feb 09, 2011

**FENTANYL-62**

<b>AB</b>	AVEVA	<b>62.5MCG/HR</b>	<b>A077449 007</b>	Dec 06, 2017
-----------	-------	-------------------	--------------------	--------------

**FENTANYL-75**

<b>AB</b>	3M DRUG DELIVERY	<b>75MCG/HR</b>	<b>A202097 004</b>	Nov 04, 2016
<b>AB</b>	ACTAVIS LABS UT INC	<b>75MCG/HR</b>	<b>A076709 003</b>	Aug 20, 2007
<b>AB</b>	AVEVA	<b>75MCG/HR</b>	<b>A077449 003</b>	Oct 20, 2008
<b>AB</b>	LAVIPHARM LABS	<b>75MCG/HR</b>	<b>A077051 003</b>	Aug 04, 2006
<b>AB</b>	MAYNE PHARMA	<b>75MCG/HR</b>	<b>A077062 003</b>	Aug 20, 2007
<b>AB</b>	MYLAN TECHNOLOGIES	<b>75MCG/HR</b>	<b>A076258 003</b>	Jan 28, 2005
<b>AB</b>	SPECGX LLC	<b>75MCG/HR</b>	<b>A077154 003</b>	Feb 09, 2011

**FENTANYL-87**

<b>AB</b>	AVEVA	<b>87.5MCG/HR</b>	<b>A077449 008</b>	Dec 06, 2017
	FENTANYL-37			
	MYLAN TECHNOLOGIES	37.5MCG/HR	A076258 006	Dec 29, 2014
	FENTANYL-62			
	MYLAN TECHNOLOGIES	62.5MCG/HR	A076258 007	Dec 29, 2014
	FENTANYL-87			
	MYLAN TECHNOLOGIES	87.5MCG/HR	A076258 008	Dec 29, 2014

SPRAY; SUBLINGUAL

SUBSYS

+	INSYS DEV CO INC	0.1MG	N202788 001	Jan 04, 2012
+		0.2MG	N202788 002	Jan 04, 2012
+!		0.4MG	N202788 003	Jan 04, 2012
+		0.6MG	N202788 004	Jan 04, 2012
+		0.8MG	N202788 005	Jan 04, 2012
+		1.2MG	N202788 006	Aug 30, 2012
+		1.6MG	N202788 007	Aug 30, 2012

**FENTANYL CITRATE**

INJECTABLE; INJECTION

**FENTANYL CITRATE**

<b>AP</b>	HOSPIRA	<b>EQ 0.05MG BASE/ML</b>	<b>N019115 001</b>	Jan 12, 1985
	FENTANYL CITRATE PRESERVATIVE FREE			
<b>AP</b>	HOSPIRA	<b>EQ 0.05MG BASE/ML</b>	<b>A072786 001</b>	Sep 24, 1991

<b>AP</b>	+!	WEST-WARD PHARMS INTL	<b>EQ 0.05MG BASE/ML</b>	<b>N019101 001</b>	Jul 11, 1984
-----------	----	-----------------------	--------------------------	--------------------	--------------

**SUBLIMAZE PRESERVATIVE FREE**

<b>AP</b>	+!	AKORN	<b>EQ 0.05MG BASE/ML</b>	<b>N016619 001</b>
	SPRAY, METERED;NASAL			
	LAZANDA			
+	ELEFSEE PHARMS INTL	EQ 0.1MG BASE	N022569 001	Jun 30, 2011
+		EQ 0.3MG BASE	N022569 003	Dec 21, 2015
+!		EQ 0.4MG BASE	N022569 002	Jun 30, 2011

TABLET;BUCCAL, SUBLINGUAL

FENTORA

+	CEPHALON	EQ 0.1MG BASE	N021947 001	Sep 25, 2006
+		EQ 0.2MG BASE	N021947 002	Sep 25, 2006
+!		EQ 0.4MG BASE	N021947 003	Sep 25, 2006
+		EQ 0.6MG BASE	N021947 004	Sep 25, 2006
+		EQ 0.8MG BASE	N021947 005	Sep 25, 2006

TABLET;SUBLINGUAL

**ABSTRAL**

<b>AB</b>	+	SENTYNL THERAPS INC	<b>EQ 0.1IMG BASE</b>	<b>N022510 001</b>	Jan 07, 2011
<b>AB</b>	+		<b>EQ 0.2MG BASE</b>	<b>N022510 002</b>	Jan 07, 2011
<b>AB</b>	+		<b>EQ 0.3MG BASE</b>	<b>N022510 003</b>	Jan 07, 2011
<b>AB</b>	+!		<b>EQ 0.4MG BASE</b>	<b>N022510 004</b>	Jan 07, 2011
<b>AB</b>	+		<b>EQ 0.6MG BASE</b>	<b>N022510 005</b>	Jan 07, 2011
<b>AB</b>	+		<b>EQ 0.8MG BASE</b>	<b>N022510 006</b>	Jan 07, 2011

**FENTANYL CITRATE**

<b>AB</b>	ACTAVIS LABS FL INC	<b>EQ 0.1IMG BASE</b>	<b>A207338 001</b>	Nov 17, 2017
<b>AB</b>		<b>EQ 0.2MG BASE</b>	<b>A207338 002</b>	Nov 17, 2017
<b>AB</b>		<b>EQ 0.3MG BASE</b>	<b>A207338 003</b>	Nov 17, 2017
<b>AB</b>		<b>EQ 0.4MG BASE</b>	<b>A207338 004</b>	Nov 17, 2017
<b>AB</b>		<b>EQ 0.6MG BASE</b>	<b>A207338 005</b>	Nov 17, 2017
<b>AB</b>		<b>EQ 0.8MG BASE</b>	<b>A207338 006</b>	Nov 17, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-179 (of 436)

FENTANYL CITRATE

TROCHE/LOZENGE; TRANSMUCOSAL

<u>ACTION</u>					
<u>AB</u>	+	CEPHALON	<u>EQ 0.2MG BASE</u>	<u>N020747 001</u>	Nov 04, 1998
<u>AB</u>	+!		<u>EQ 0.4MG BASE</u>	<u>N020747 002</u>	Nov 04, 1998
<u>AB</u>	+		<u>EQ 0.6MG BASE</u>	<u>N020747 003</u>	Nov 04, 1998
<u>AB</u>	+		<u>EQ 0.8MG BASE</u>	<u>N020747 004</u>	Nov 04, 1998
<u>AB</u>	+		<u>EQ 1.2MG BASE</u>	<u>N020747 005</u>	Nov 04, 1998
<u>AB</u>	+		<u>EQ 1.6MG BASE</u>	<u>N020747 006</u>	Nov 04, 1998
<u>FENTANYL CITRATE</u>					
<u>AB</u>		PAR PHARM	<u>EQ 0.2MG BASE</u>	<u>A077312 001</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 0.4MG BASE</u>	<u>A077312 002</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 0.6MG BASE</u>	<u>A077312 003</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 0.8MG BASE</u>	<u>A077312 004</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 1.2MG BASE</u>	<u>A077312 005</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 1.6MG BASE</u>	<u>A077312 006</u>	Oct 30, 2009
<u>AB</u>		SPECGX LLC	<u>EQ 0.2MG BASE</u>	<u>A078907 001</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 0.4MG BASE</u>	<u>A078907 002</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 0.6MG BASE</u>	<u>A078907 003</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 0.8MG BASE</u>	<u>A078907 004</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 1.2MG BASE</u>	<u>A078907 005</u>	Oct 30, 2009
<u>AB</u>			<u>EQ 1.6MG BASE</u>	<u>A078907 006</u>	Oct 30, 2009

FENTANYL HYDROCHLORIDE

SYSTEM; IONTOPHORESIS, TRANSDERMAL

IONSYS				
+!	THE MEDICINES CO	EQ 40MCG BASE/ACTIVATION	N021338 001	May 22, 2006

FERRIC CARBOXYMALTOSA

INJECTABLE; INTRAVENOUS

INJECTAFER				
+!	LUITPOLD PHARMS INC	750MG IRON/15ML (50MG IRON/ML)	N203565 001	Jul 25, 2013

FERRIC CITRATE

TABLET; ORAL

AURYXIA				
+!	KERYX BIOPHARMS	EQ 210MG IRON	N205874 001	Sep 05, 2014

FERRIC HEXACYANOFERRATE (II)

CAPSULE; ORAL

RADIOGARDASE (PRUSSIAN BLUE)				
+!	HEYL CHEMISCHE	500MG	N021626 001	Oct 02, 2003

FERRIC PYROPHOSPHATE CITRATE

FOR SOLUTION; INTRAVENOUS

TRIFERIC				
+!	ROCKWELL MEDICAL INC	272MG IRON/PACKET	N208551 001	Apr 25, 2016
SOLUTION; IV (INFUSION)				
TRIFERIC				
+!	ROCKWELL MEDICAL INC	27.2MG IRON/5ML (5.44MG IRON/ML)	N206317 001	Jan 23, 2015
+		272MG IRON/50ML (5.44MG IRON/ML)	N206317 002	Sep 04, 2015

FERUMOXYTOL

SOLUTION; INTRAVENOUS

FERAHEME				
+!	AMAG PHARMS INC	EQ 510MG IRON/17ML (EQ 30MG IRON/ML)	N022180 001	Jun 30, 2009

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

<u>FESOTERODINE FUMARATE</u>					
<u>AB</u>	AUROBINDO PHARMA LTD	<u>4MG</u>	<u>A205007 001</u>	Feb 17, 2017	
<u>AB</u>		<u>8MG</u>	<u>A205007 002</u>	Feb 17, 2017	
<u>AB</u>	ZYDUS PHARMS USA INC	<u>4MG</u>	<u>A204946 001</u>	Oct 03, 2017	
<u>AB</u>		<u>8MG</u>	<u>A204946 002</u>	Oct 03, 2017	
<u>TOVIAZ</u>					
<u>AB</u>	+	Pfizer	<u>4MG</u>	<u>N022030 001</u>	Oct 31, 2008
<u>AB</u>	+		<u>8MG</u>	<u>N022030 002</u>	Oct 31, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-180 (of 436)

FEXOFENADINE HYDROCHLORIDE

SUSPENSION;ORAL

**ALLEGRA**

**AB** +! SANOFI AVENTIS US **30MG/5ML** **N021963 001** Oct 16, 2006

**FEXOFENADINE HYDROCHLORIDE**

**AB** ACTAVIS MID ATLANTIC **30MG/5ML** **A201311 001** Jul 25, 2012

TABLET;ORAL

**FEXOFENADINE HYDROCHLORIDE**

<b>AB</b>	BARR	<b>30MG</b>	<b>A076191 001</b>	Aug 31, 2005
<b>AB</b>		<b>60MG</b>	<b>A076191 002</b>	Aug 31, 2005
<b>AB</b>		<b>180MG</b>	<b>A076191 003</b>	Aug 31, 2005
<b>AB</b>	DR REDDYS LABS LTD	<b>30MG</b>	<b>A076502 001</b>	Apr 11, 2006
<b>AB</b>		<b>60MG</b>	<b>A076502 002</b>	Apr 11, 2006
<b>AB</b>		<b>180MG</b>	<b>A076502 003</b>	Apr 11, 2006
<b>AB</b>	MYLAN	<b>30MG</b>	<b>A077081 002</b>	Apr 11, 2008
<b>AB</b>		<b>60MG</b>	<b>A077081 003</b>	Apr 11, 2008
<b>AB</b>		<b>180MG</b>	<b>A077081 001</b>	Apr 16, 2007
<b>AB</b>	TEVA	<b>30MG</b>	<b>A076447 001</b>	Sep 01, 2005
<b>AB</b>		<b>60MG</b>	<b>A076447 002</b>	Sep 01, 2005
<b>AB</b>		<b>180MG</b>	<b>A076447 003</b>	Sep 01, 2005

FIDAXOMICIN

TABLET;ORAL

DIFICID

+! CUBIST PHARMS LLC 200MG **N201699 001** May 27, 2011

FINAFLOXACIN

SUSPENSION/DROPS;OTIC

XTORO

+! NOVARTIS PHARMS CORP 0.3% **N206307 001** Dec 17, 2014

FINASTERIDE

TABLET;ORAL

**FINASTERIDE**

<b>AB</b>	ACCORD HLTHCARE	<b>1MG</b>	<b>A091643 001</b>	Nov 05, 2013
<b>AB</b>		<b>5MG</b>	<b>A090121 001</b>	Feb 23, 2010
<b>AB</b>	ACTAVIS TOTOWA	<b>1MG</b>	<b>A078371 001</b>	Nov 05, 2013
<b>AB</b>	ACTAVIS TOTOWA TEVA	<b>5MG</b>	<b>A077914 001</b>	Mar 28, 2007
<b>AB</b>	ALKEM LABS LTD	<b>1MG</b>	<b>A207750 001</b>	Jan 06, 2017
<b>AB</b>		<b>5MG</b>	<b>A204304 001</b>	Jan 05, 2017
<b>AB</b>	AUROBINDO PHARMA	<b>5MG</b>	<b>A078341 001</b>	Oct 30, 2007
<b>AB</b>	AUROBINDO PHARMA LTD	<b>1MG</b>	<b>A203687 001</b>	Nov 05, 2013
<b>AB</b>	CIPILA LTD	<b>1MG</b>	<b>A077335 001</b>	Nov 20, 2014
<b>AB</b>	DR REDDYS LABS INC	<b>1MG</b>	<b>A076436 001</b>	Jul 28, 2006
<b>AB</b>	DR REDDYS LABS LTD	<b>5MG</b>	<b>A076437 001</b>	Feb 28, 2007
<b>AB</b>	GEDEON RICHTER USA	<b>5MG</b>	<b>A077251 001</b>	Dec 22, 2006
<b>AB</b>	HETERO LABS LTD III	<b>1MG</b>	<b>A090060 001</b>	Jul 01, 2013
<b>AB</b>		<b>5MG</b>	<b>A090061 001</b>	Jun 07, 2010
<b>AB</b>	MYLAN	<b>5MG</b>	<b>A077578 001</b>	Dec 18, 2006
<b>AB</b>	MYLAN PHARMS INC	<b>1MG</b>	<b>A078161 001</b>	Nov 05, 2013
<b>AB</b>	SUN PHARMA GLOBAL	<b>1MG</b>	<b>A090508 001</b>	Jul 01, 2013
<b>AB</b>		<b>5MG</b>	<b>A090507 001</b>	Aug 16, 2011
<b>AB</b>	TEVA	<b>1MG</b>	<b>A076905 001</b>	Nov 05, 2013
<b>AB</b>		<b>5MG</b>	<b>A076511 001</b>	Dec 15, 2006
<b>AB</b>	ZYDUS PHARMS USA INC	<b>5MG</b>	<b>A078900 001</b>	Dec 28, 2009

**PROPECIA**

**AB** +! MERCK **1MG** **N020788 001** Dec 19, 1997

**PROSCAR**

**AB** +! MERCK **5MG** **N020180 001** Jun 19, 1992

FINGOLIMOD

CAPSULE;ORAL

GILENYA

+! NOVARTIS 0.5MG **N022527 001** Sep 21, 2010

FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL

EMULSION;INTRAVENOUS

SMOFLIPID 20%

+! FRESENIUS KABI USA 3GM/100ML;6GM/100ML;5GM/100ML;6GM/100ML (100ML) **N207648 001** Jul 13, 2016

+! 3GM/100ML;6GM/100ML;5GM/100ML;6GM/100ML (250ML) **N207648 002** Jul 13, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-181 (of 436)

FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL

EMULSION; INTRAVENOUS

SMOFLIPID 20%

+!

3GM/100ML; 6GM/100ML; 5GM/100ML; 6GM/100ML (500ML) N207648 003 Jul 13, 2016

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

**FLAVOXATE HYDROCHLORIDE**

<b>AB</b>	EPIC PHARMA	<b><u>100MG</u></b>	
<b>AB</b>	! PADDOCK LLC	<b><u>100MG</u></b>	

<b>A076835 001</b>	Nov 30, 2005
<b>A076831 001</b>	Dec 16, 2004

FLECAINIDE ACETATE

TABLET; ORAL

**FLECAINIDE ACETATE**

<b>AB</b>	AMNEAL PHARM	<b><u>50MG</u></b>	
<b>AB</b>		<b><u>100MG</u></b>	
<b>AB</b>		<b><u>150MG</u></b>	
<b>AB</b>	ANI PHARMS INC	<b><u>50MG</u></b>	
<b>AB</b>		<b><u>100MG</u></b>	
<b>AB</b>		<b><u>150MG</u></b>	
<b>AB</b>	AUROBINDO PHARMA LTD	<b><u>50MG</u></b>	
<b>AB</b>		<b><u>100MG</u></b>	
<b>AB</b>		<b><u>150MG</u></b>	
<b>AB</b>	SUN PHARM INDs LTD	<b><u>50MG</u></b>	
<b>AB</b>		<b><u>100MG</u></b>	
<b>AB</b>		<b><u>150MG</u></b>	
<b>AB</b>	WEST-WARD PHARMS INT	<b><u>50MG</u></b>	
<b>AB</b>		<b><u>100MG</u></b>	
<b>AB</b>	!	<b><u>150MG</u></b>	
<b>TAMBOCOR</b>			
<b>AB</b>	+	CNTY LINE PHARMS	<b><u>50MG</u></b>
<b>AB</b>	+		<b><u>100MG</u></b>
<b>AB</b>	+		<b><u>150MG</u></b>

<b>A075442 001</b>	Jul 31, 2001
<b>A075442 002</b>	Jul 31, 2001
<b>A075442 003</b>	Jul 31, 2001
<b>A075882 001</b>	Oct 28, 2002
<b>A075882 002</b>	Oct 28, 2002
<b>A075882 003</b>	Oct 28, 2002
<b>A202821 001</b>	Nov 03, 2017
<b>A202821 002</b>	Nov 03, 2017
<b>A202821 003</b>	Nov 03, 2017
<b>A076421 001</b>	Mar 28, 2003
<b>A076421 002</b>	Mar 28, 2003
<b>A076421 003</b>	Mar 28, 2003
<b>A076278 001</b>	Jan 14, 2003
<b>A076278 002</b>	Jan 14, 2003
<b>A076278 003</b>	Jan 14, 2003
<b>N018830 004</b>	Aug 23, 1988
<b>N018830 001</b>	Oct 31, 1985
<b>N018830 003</b>	Jun 03, 1988

FLIBANSERIN

TABLET; ORAL

ADDYI

+! SPROUT PHARMS

100MG

N022526 001 Aug 18, 2015

FLORBETABEN F-18

SOLUTION; INTRAVENOUS

NEURACEQ

+! PIRAMAL IMAGING

30ML (1.4-135mCi/ML)

N204677 001 Mar 19, 2014

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

+! AVID RADIOPHARMS INC

10-30ML (13.5-51mCi/ML)

N202008 002 Apr 06, 2012

+!

10-50ML (13.5-51mCi/ML)

N202008 003 Apr 06, 2012

FLOXURIDINE

INJECTABLE; INJECTION

**FLOXURIDINE**

<b>AP</b>	FRESENIUS KABI USA	<b><u>500MG/VIAL</u></b>	
<b>AP</b>	PHARMAFORCE	<b><u>500MG/VIAL</u></b>	
<b>AP</b>	! WEST-WARD PHARMS INT	<b><u>500MG/VIAL</u></b>	

<b>A075837 001</b>	Feb 22, 2001
<b>A203008 001</b>	Nov 22, 2017
<b>A075387 001</b>	Apr 16, 2000

FLUCICLOVINE F-18

SOLUTION; INTRAVENOUS

AXUMIN

+! BLUE EARTH

9-221mCi/ML

N208054 001 May 27, 2016

FLUCONAZOLE

FOR SUSPENSION; ORAL

**DIFLUCAN**

<b>AB</b>	+	PFIZER	<b><u>50MG/5ML</u></b>	
<b>AB</b>	+	!	<b><u>200MG/5ML</u></b>	

<b>N020090 001</b>	Dec 23, 1993
<b>N020090 002</b>	Dec 23, 1993

**FLUCONAZOLE**

<b>AB</b>	AUROBINDO PHARMA LTD	<b><u>50MG/5ML</u></b>	
<b>AB</b>		<b><u>200MG/5ML</u></b>	
<b>AB</b>	IVAX SUB TEVA PHARMS	<b><u>50MG/5ML</u></b>	

<b>A079150 001</b>	Sep 18, 2009
<b>A079150 002</b>	Sep 18, 2009
<b>A077523 001</b>	Sep 12, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-182 (of 436)

FLUCONAZOLE

FOR SUSPENSION;ORAL

FLUCONAZOLE

<u>AB</u>		<u>200MG/5ML</u>	<u>A077523 002</u>	Sep 12, 2007
<u>AB</u>	WEST-WARD PHARMS INT	<u>50MG/5ML</u>	<u>A076246 001</u>	Jul 29, 2004
<u>AB</u>		<u>200MG/5ML</u>	<u>A076246 002</u>	Jul 29, 2004

INJECTABLE;INJECTION

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML (2MG/ML)</u>	<u>A078764 001</u>	Jan 30, 2012
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A078764 002</u>	Jan 30, 2012
<u>AP</u>	HOSPIRA	<u>200MG/100ML (2MG/ML)</u>	<u>A076304 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076304 002</u>	Jul 29, 2004
<u>AP</u>	RENAISSANCE SSA LLC	<u>200MG/100ML (2MG/ML)</u>	<u>A077988 001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077988 002</u>	May 26, 2010

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

<u>AP</u>	BAXTER HLTHCARE CORP	<u>200MG/100ML (2MG/ML)</u>	<u>A077947 001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077947 002</u>	May 26, 2010
<u>AP</u>	FRESENIUS KABI USA	<u>200MG/100ML (2MG/ML)</u>	<u>A076145 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076145 002</u>	Jul 29, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML (2MG/ML)</u>	<u>A076736 001</u>	Aug 23, 2005
<u>AP</u>	WEST-WARD PHARMS INT	<u>200MG/100ML (2MG/ML)</u>	<u>A076087 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076087 003</u>	Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	! ACS DOBFAR INFO SA	<u>200MG/100ML (2MG/ML)</u>	<u>A079104 001</u>	Jul 30, 2009
<u>AP</u>	!	<u>400MG/200ML (2MG/ML)</u>	<u>A079104 002</u>	Jul 30, 2009
<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML (2MG/ML)</u>	<u>A076766 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076766 002</u>	Jul 29, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML (2MG/ML)</u>	<u>A078698 001</u>	Jan 30, 2012
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A078698 002</u>	Jan 30, 2012
<u>AP</u>	HOSPIRA	<u>200MG/100ML (2MG/ML)</u>	<u>A076303 001</u>	Jul 29, 2004
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A076303 002</u>	Jul 29, 2004
<u>AP</u>	RENAISSANCE SSA LLC	<u>200MG/100ML (2MG/ML)</u>	<u>A077909 001</u>	May 26, 2010
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A077909 002</u>	May 26, 2010
<u>AP</u>	WEST-WARD PHARMS INT	<u>200MG/100ML (2MG/ML)</u>	<u>A078107 001</u>	Jul 30, 2008
<u>AP</u>		<u>400MG/200ML (2MG/ML)</u>	<u>A078107 002</u>	Jul 30, 2008

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

WEST-WARD PHARMS  
INT

A076087 002 Sep 26, 2008

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

RENAISSANCE SSA LLC 100MG/50ML (2MG/ML)

A077909 003 Apr 20, 2015

TABLET;ORAL

DIFLUCAN

<u>AB</u>	+	PFIZER	<u>50MG</u>	<u>N019949 001</u>	Jan 29, 1990
<u>AB</u>	+		<u>100MG</u>	<u>N019949 002</u>	Jan 29, 1990
<u>AB</u>	+		<u>150MG</u>	<u>N019949 004</u>	Jun 30, 1994
<u>AB</u>	++!		<u>200MG</u>	<u>N019949 003</u>	Jan 29, 1990

FLUCONAZOLE

<u>AB</u>		APOTEX	<u>50MG</u>	<u>A076665 001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076665 002</u>	Jul 29, 2004
<u>AB</u>			<u>150MG</u>	<u>A076665 003</u>	Jul 29, 2004
<u>AB</u>			<u>200MG</u>	<u>A076665 004</u>	Jul 29, 2004
<u>AB</u>		AUROBINDO PHARMA	<u>50MG</u>	<u>A077731 001</u>	Oct 07, 2008
<u>AB</u>			<u>100MG</u>	<u>A077731 002</u>	Oct 07, 2008
<u>AB</u>			<u>150MG</u>	<u>A077731 003</u>	Oct 07, 2008
<u>AB</u>			<u>200MG</u>	<u>A077731 004</u>	Oct 07, 2008
<u>AB</u>		DR REDDYS LABS INC	<u>50MG</u>	<u>A076658 001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076658 002</u>	Jul 29, 2004
<u>AB</u>			<u>150MG</u>	<u>A076658 003</u>	Jul 29, 2004
<u>AB</u>			<u>200MG</u>	<u>A076658 004</u>	Jul 29, 2004
<u>AB</u>		GLENMARK GENERICS	<u>50MG</u>	<u>A077253 001</u>	Jan 25, 2006
<u>AB</u>			<u>100MG</u>	<u>A077253 002</u>	Jan 25, 2006
<u>AB</u>			<u>150MG</u>	<u>A077253 003</u>	Jan 25, 2006
<u>AB</u>			<u>200MG</u>	<u>A077253 004</u>	Jan 25, 2006
<u>AB</u>		HARRIS PHARM	<u>50MG</u>	<u>A078423 001</u>	Mar 07, 2011
<u>AB</u>			<u>100MG</u>	<u>A078423 002</u>	Mar 07, 2011
<u>AB</u>			<u>150MG</u>	<u>A078423 003</u>	Mar 07, 2011
<u>AB</u>			<u>200MG</u>	<u>A078423 004</u>	Mar 07, 2011
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>50MG</u>	<u>A076077 001</u>	Jul 29, 2004
<u>AB</u>			<u>100MG</u>	<u>A076077 002</u>	Jul 29, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-183 (of 436)

**FLUCONAZOLE**

TABLET;ORAL

**FLUCONAZOLE**

<b><u>AB</u></b>		<b><u>150MG</u></b>	<b><u>A076077 003</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>200MG</u></b>	<b><u>A076077 004</u></b>	Jul 29, 2004
<b><u>AB</u></b>	MYLAN	<b><u>50MG</u></b>	<b><u>A076351 001</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>100MG</u></b>	<b><u>A076351 002</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>150MG</u></b>	<b><u>A076351 003</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>200MG</u></b>	<b><u>A076351 004</u></b>	Jul 29, 2004
<b><u>AB</u></b>	TARO	<b><u>50MG</u></b>	<b><u>A076507 001</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>100MG</u></b>	<b><u>A076507 002</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>150MG</u></b>	<b><u>A076507 003</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>200MG</u></b>	<b><u>A076507 004</u></b>	Jul 29, 2004
<b><u>AB</u></b>	TEVA	<b><u>50MG</u></b>	<b><u>A074681 001</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>100MG</u></b>	<b><u>A074681 002</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>150MG</u></b>	<b><u>A074681 003</u></b>	Jul 29, 2004
<b><u>AB</u></b>		<b><u>200MG</u></b>	<b><u>A074681 004</u></b>	Jul 29, 2004
<b><u>AB</u></b>	UNIQUE PHARM LABS	<b><u>50MG</u></b>	<b><u>A076957 001</u></b>	Sep 28, 2005
<b><u>AB</u></b>		<b><u>100MG</u></b>	<b><u>A076957 002</u></b>	Sep 28, 2005
<b><u>AB</u></b>		<b><u>150MG</u></b>	<b><u>A076957 004</u></b>	Feb 27, 2017
<b><u>AB</u></b>		<b><u>200MG</u></b>	<b><u>A076957 003</u></b>	Sep 28, 2005
<b><u>AB</u></b>	ZYDUS PHARMS USA INC	<b><u>50MG</u></b>	<b><u>A208963 001</u></b>	Feb 16, 2017
<b><u>AB</u></b>		<b><u>100MG</u></b>	<b><u>A208963 002</u></b>	Feb 16, 2017
<b><u>AB</u></b>		<b><u>150MG</u></b>	<b><u>A208963 003</u></b>	Feb 16, 2017
<b><u>AB</u></b>		<b><u>200MG</u></b>	<b><u>A208963 004</u></b>	Feb 16, 2017

**FLUCYTOSINE**

CAPSULE;ORAL

**ANCOBON**

<b><u>AB</u></b>	+	VALEANT	<b><u>250MG</u></b>	<b><u>N017001 001</u></b>
<b><u>AB</u></b>	+!		<b><u>500MG</u></b>	<b><u>N017001 002</u></b>
<b><u>FLUCYTOSINE</u></b>				
<b><u>AB</u></b>		NOVEL LABS INC	<b><u>250MG</u></b>	<b><u>A204652 001</u></b>
<b><u>AB</u></b>			<b><u>500MG</u></b>	<b><u>A204652 002</u></b>
<b><u>AB</u></b>		SIGMAPHARM LABS LLC	<b><u>250MG</u></b>	<b><u>A201566 001</u></b>
<b><u>AB</u></b>			<b><u>500MG</u></b>	<b><u>A201566 002</u></b>
<b><u>AB</u></b>		WEST-WARD PHARMS INT	<b><u>250MG</u></b>	<b><u>A206550 001</u></b>
<b><u>AB</u></b>			<b><u>500MG</u></b>	<b><u>A206550 002</u></b>
<b><u>FLUDARABINE PHOSPHATE</u></b>				
INJECTABLE;INJECTION				
<b><u>FLUDARABINE PHOSPHATE</u></b>				
<b><u>AP</u></b>		ACTAVIS LLC	<b><u>50MG/2ML (25MG/ML)</u></b>	<b><u>A203738 001</u></b>
<b><u>AP</u></b>		ACTAVIS TOTOWA	<b><u>50MG/VIAL</u></b>	<b><u>A078610 001</u></b>
<b><u>AP</u></b>		CUSTOPHARM INC	<b><u>50MG/VIAL</u></b>	<b><u>A076349 001</u></b>
<b><u>AP</u></b>	!	FRESENIUS KABI USA	<b><u>50MG/2ML (25MG/ML)</u></b>	<b><u>A078393 001</u></b>
<b><u>AP</u></b>			<b><u>50MG/VIAL</u></b>	<b><u>A078544 001</u></b>
<b><u>AP</u></b>	!	HOSPIRA	<b><u>50MG/VIAL</u></b>	<b><u>A077790 001</u></b>
<b><u>AP</u></b>		MUSTAFA NEVZAT ILAC	<b><u>50MG/2ML (25MG/ML)</u></b>	<b><u>A090724 001</u></b>
<b><u>AP</u></b>		MYLAN LABS LTD	<b><u>50MG/2ML (25MG/ML)</u></b>	<b><u>A200647 001</u></b>
<b><u>AP</u></b>			<b><u>50MG/VIAL</u></b>	<b><u>A200648 001</u></b>
<b><u>AP</u></b>		SAGENT PHARMS	<b><u>50MG/2ML (25MG/ML)</u></b>	<b><u>A076661 001</u></b>
<b><u>AP</u></b>	+	SANDOZ	<b><u>50MG/2ML (25MG/ML)</u></b>	<b><u>N022137 001</u></b>
<b><u>FLUDEOXYGLUCOSE F-18</u></b>				
INJECTABLE;INTRAVENOUS				
<b><u>FLUDEOXYGLUCOSE F18</u></b>				
<b><u>AP</u></b>		3D IMAGING DRUG	<b><u>20-300mCi/ML</u></b>	<b><u>A203778 001</u></b>
<b><u>AP</u></b>		BIOMEDCL RES FDN	<b><u>20-300mCi/ML</u></b>	<b><u>A203710 001</u></b>
<b><u>AP</u></b>			<b><u>20-300mCi/ML</u></b>	<b><u>A203837 001</u></b>
<b><u>AP</u></b>		BRIGHAM WOMENS	<b><u>20-300mCi/ML</u></b>	<b><u>A203816 001</u></b>
<b><u>AP</u></b>		CARDINAL HEALTH 414	<b><u>20-300mCi/ML</u></b>	<b><u>A203603 001</u></b>
<b><u>AP</u></b>		CHILDRENS HOSP MI	<b><u>20-300mCi/ML</u></b>	<b><u>A204385 001</u></b>
<b><u>AP</u></b>		CPDC	<b><u>20-300mCi/ML</u></b>	<b><u>A204525 001</u></b>
<b><u>AP</u></b>		ESSENTIAL ISOTOPES	<b><u>20-300mCi/ML</u></b>	<b><u>A203946 001</u></b>
<b><u>AP</u></b>	+	FEINSTAIN	<b><u>20-200mCi/ML</u></b>	<b><u>N021870 001</u></b>
<b><u>AP</u></b>	+		<b><u>20-400mCi/ML</u></b>	<b><u>N021870 002</u></b>
<b><u>AP</u></b>		GLOBAL ISOTOPES LLC	<b><u>20-300mCi/ML</u></b>	<b><u>A204463 001</u></b>
<b><u>AP</u></b>	!	HOUSTON CYCLOTRON	<b><u>20-500mCi/ML</u></b>	<b><u>A203665 001</u></b>
<b><u>AP</u></b>		JUBILANT DRAXIMAGE	<b><u>20-300mCi/ML</u></b>	<b><u>A203920 001</u></b>
<b><u>AP</u></b>		KETTERING MEDCTR	<b><u>4-40mCi/ML</u></b>	<b><u>A204759 001</u></b>
<b><u>AP</u></b>		KREITCHMAN PET CTR	<b><u>10-100mCi/ML</u></b>	<b><u>A203942 001</u></b>

<b><u>A203778 001</u></b>	Oct 30, 2015
<b><u>A203710 001</u></b>	May 01, 2015
<b><u>A203837 001</u></b>	May 01, 2015
<b><u>A203816 001</u></b>	Oct 30, 2014
<b><u>A203603 001</u></b>	Nov 13, 2015
<b><u>A204385 001</u></b>	Oct 29, 2014
<b><u>A204525 001</u></b>	Oct 29, 2014
<b><u>A203946 001</u></b>	Feb 05, 2014
<b><u>N021870 001</u></b>	Aug 19, 2005
<b><u>N021870 002</u></b>	Nov 21, 2008
<b><u>A204463 001</u></b>	Oct 21, 2014
<b><u>A203665 001</u></b>	Feb 14, 2013
<b><u>A203920 001</u></b>	Jun 23, 2015
<b><u>A204759 001</u></b>	Oct 27, 2015
<b><u>A203942 001</u></b>	Apr 11, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-184 (of 436)

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F18

<u>AP</u>	LANTHEUS MEDICAL	<u>20-200mCi/ML</u>	<u>A203664</u> <u>001</u>	Feb 04, 2014
<u>AP</u>	MA GENERAL HOSP	<u>20-300mCi/ML</u>	<u>A204333</u> <u>001</u>	Sep 25, 2014
<u>AP</u>	MCPRF	<u>20-240mCi/ML</u>	<u>A203612</u> <u>001</u>	Aug 05, 2013
<u>AP</u>	MEM SLOAN-KETTERING	<u>20-300mCi/ML</u>	<u>A208679</u> <u>001</u>	Dec 08, 2016
<u>AP</u>	METHODIST HOSP RES	<u>20-300mCi/ML</u>	<u>A203904</u> <u>001</u>	Apr 23, 2015
<u>AP</u>	MIDWEST MEDCL	<u>20-200mCi/ML</u>	<u>A203736</u> <u>001</u>	Nov 19, 2015
<u>AP</u>	MIPS CRF	<u>20-300mCi/ML</u>	<u>A204472</u> <u>001</u>	Sep 11, 2015
<u>AP</u>	NCM USA BRONX LLC	<u>20-300mCi/ML</u>	<u>A204512</u> <u>001</u>	Jan 07, 2015
<u>AP</u>	PETNET	<u>20-200mCi/ML</u>	<u>A079086</u> <u>001</u>	Feb 25, 2011
<u>AP</u>	QUEEN HAMAMATSU PET	<u>10-100mCi/ML</u>	<u>A203771</u> <u>001</u>	Aug 31, 2015
<u>AP</u>	SHERTECH LABS LLC	<u>20-300mCi/ML</u>	<u>A204264</u> <u>001</u>	Dec 18, 2014
<u>AP</u>	SOFIE	<u>20-300mCi/ML</u>	<u>A203591</u> <u>001</u>	Aug 31, 2015
<u>AP</u>	TRUSTEES UNIV PA	<u>20-200mCi/ML</u>	<u>A203801</u> <u>001</u>	Oct 29, 2014
<u>AP</u>	UCLA BIOMEDICAL	<u>4-40mCi/ML</u>	<u>A203811</u> <u>001</u>	Jun 27, 2013
<u>AP</u>	UCSF RODIOPHARM	<u>20-300mCi/ML</u>	<u>A203902</u> <u>001</u>	May 09, 2014
<u>AP</u>	UIHC PET IMAGING	<u>20-300mCi/ML</u>	<u>A203990</u> <u>001</u>	Aug 06, 2014
<u>AP</u>	UNIV MICHIGAN	<u>20-300mCi/ML</u>	<u>A204531</u> <u>001</u>	Jul 17, 2015
<u>AP</u>	UNIV TX MD ANDERSON	<u>20-300mCi/ML</u>	<u>A203246</u> <u>002</u>	Jan 13, 2014
<u>AP</u>	UNIV UTAH CYCLOTRON	<u>20-300mCi/ML</u>	<u>A204498</u> <u>001</u>	Jun 23, 2015
<u>AP</u>	WI MEDCL CYCLOTRON	<u>20-500mCi/ML</u>	<u>A203709</u> <u>001</u>	Oct 23, 2013
<u>AP</u>	WUSM CYCLOTRON	<u>20-300mCi/ML</u>	<u>A203935</u> <u>001</u>	Feb 05, 2014
	HOT SHOTS NM LLC	4-500mCi/ML	A203937	001 Oct 30, 2014
	PRECISION NUCLEAR	20-500mCi/ML	A204546	001 Apr 07, 2015
	SPECTRON MRC LLC	4-500mCi/ML	A203911	001 Apr 22, 2015
	UNIV NORTH DAKOTA	4-500mCi/ML	A203994	001 Feb 04, 2015
	UNIV TX MD ANDERSON	20-150mCi/ML	A203246	001 Jan 13, 2014

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLUDROCORTISONE ACETATE

<u>AB</u>	BARR	<u>0.1MG</u>	<u>A040425</u> <u>001</u>	Jan 21, 2003
<u>AB</u>	HIKMA PHARMS	<u>0.1MG</u>	<u>A091302</u> <u>001</u>	Jul 22, 2011
<u>AB</u>	! IMPAX LABS	<u>0.1MG</u>	<u>A040431</u> <u>001</u>	Mar 18, 2002

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	FRESENIUS KABI USA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076955</u> <u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076955</u> <u>001</u>	Oct 12, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078527</u> <u>001</u>	Mar 23, 2009
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A078527</u> <u>002</u>	Mar 23, 2009
<u>AP</u>	MYLAN LABS LTD	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A078595</u> <u>001</u>	May 13, 2008
<u>AP</u>	!	<u>1MG/10ML (0.1MG/ML)</u>	<u>A078595</u> <u>002</u>	May 13, 2008
<u>AP</u>	SAGENT PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A090584</u> <u>001</u>	Aug 28, 2012
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A090584</u> <u>002</u>	Aug 28, 2012
<u>AP</u>	SANDOZ INC	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A077071</u> <u>001</u>	May 03, 2005
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A077071</u> <u>002</u>	May 03, 2005
<u>AP</u>	WEST-WARD PHARMS INT	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076256</u> <u>002</u>	Oct 12, 2004
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>A076787</u> <u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076256</u> <u>001</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>A076787</u> <u>001</u>	Oct 12, 2004

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROSOL HFA

+! MYLAN SPECIALITY LP 0.078MG/INH

N021247 001 Jan 27, 2006

SPRAY, METERED; NASAL

FLUNISOLIDE

<u>AB</u>	! BAUSCH AND LOMB	<u>0.025MG/SPRAY</u>	<u>A074805</u> <u>001</u>	Feb 20, 2002
<u>AB</u>	HI TECH PHARMA CO	<u>0.025MG/SPRAY</u>	<u>A077704</u> <u>001</u>	Aug 03, 2006
	! APOTEX INC	0.029MG/SPRAY	A077436	001 Aug 09, 2007

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA PHARMS INC	<u>0.01%</u>	<u>A088170</u> <u>001</u>	Dec 16, 1982
<u>AT</u>		<u>0.025%</u>	<u>A088169</u> <u>001</u>	Dec 16, 1982
<u>AT</u>	G AND W LABS	<u>0.01%</u>	<u>A089526</u> <u>001</u>	Jul 26, 1988
<u>AT</u>		<u>0.025%</u>	<u>A089525</u> <u>001</u>	Jul 26, 1988
<u>AT</u>	TARO	<u>0.025%</u>	<u>A087104</u> <u>001</u>	Apr 27, 1982

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-185 (of 436)

FLUOCINOLONE ACETONIDE

CREAM;TOPICAL

**SYNALAR**

<b>AT</b>	+!	MEDIMETRIKS PHARMS	<u>0.01%</u>
<b>AT</b>	+!		<u>0.025%</u>
<b>AT</b>	+!		<u>0.025%</u>

IMPLANT;INTRAVITREAL

ILUVIEN

+!	ALIMERA SCIENCES INC	0.19MG
----	----------------------	--------

<b>N012787</b>	<b>004</b>
<b>N012787</b>	<b>002</b>
<b>N012787</b>	<b>005</b>

RETISERT

+!	BAUSCH AND LOMB	0.59MG
----	-----------------	--------

N201923 001 Sep 26, 2014

OIL;TOPICAL

**DERMA-SMOOTH/FS**

<b>AT</b>	+!	HILL DERMAC	<u>0.01%</u>
<b>AT</b>	+!		<u>0.01%</u>

**N019452** **001** Feb 03, 1988  
**N019452** **002** Nov 09, 2005

**FLUOCINOLONE ACETONIDE**

<b>AT</b>	AKORN	<u>0.01%</u>
<b>AT</b>	IDENTI PHARMS INC	<u>0.01%</u>
<b>AT</b>		<u>0.01%</u>
<b>AT</b>	LYNE	<u>0.01%</u>
<b>AT</b>		<u>0.01%</u>
<b>AT</b>	PERRIGO ISRAEL	<u>0.01%</u>
<b>AT</b>		<u>0.01%</u>
<b>AT</b>	TARO	<u>0.01%</u>
<b>AT</b>		<u>0.01%</u>

**A091514** **001** Jun 25, 2015  
**A201759** **001** Oct 17, 2011  
**A201764** **001** Oct 17, 2011  
**A090982** **001** Apr 25, 2016  
**A203377** **001** Apr 25, 2016  
**A202847** **001** Aug 09, 2013  
**A202848** **001** Aug 09, 2013  
**A202368** **001** May 19, 2016  
**A209336** **001** May 19, 2016

OIL/DROPS;OTIC

**DERMOTIC**

<b>AT</b>	+!	HILL DERMAC	<u>0.01%</u>
-----------	----	-------------	--------------

**N019452** **003** Nov 09, 2005

**FLUOCINOLONE ACETONIDE**

<b>AT</b>	AKORN	<u>0.01%</u>
<b>AT</b>	IDENTI PHARMS INC	<u>0.01%</u>
<b>AT</b>	LYNE	<u>0.01%</u>
<b>AT</b>	PERRIGO ISRAEL	<u>0.01%</u>

**A202705** **001** Sep 09, 2016  
**A091306** **001** Oct 17, 2011  
**A203378** **001** Apr 25, 2016  
**A202849** **001** Jul 17, 2017

OINTMENT;TOPICAL

**FLUOCINOLONE ACETONIDE**

<b>AT</b>	FOUGERA PHARMS INC	<u>0.025%</u>
<b>AT</b>	G AND W LABS	<u>0.025%</u>
<b>AT</b>	TARO	<u>0.025%</u>

**A088168** **001** Dec 16, 1982  
**A089524** **001** Jul 26, 1988  
**A040041** **001** Sep 15, 1994

**SYNALAR**

<b>AT</b>	+!	MEDIMETRIKS PHARMS	<u>0.025%</u>
-----------	----	--------------------	---------------

**N013960** **001**

SHAMPOO;TOPICAL

CAPEX

+!	GALDERMA LABS LP	0.01%
----	------------------	-------

N020001 001 Aug 27, 1990

SOLUTION;TOPICAL

**FLUOCINOLONE ACETONIDE**

<b>AT</b>	ACTAVIS LABS UT INC	<u>0.01%</u>
<b>AT</b>	FOUGERA PHARMS INC	<u>0.01%</u>
<b>AT</b>	G AND W LABS INC	<u>0.01%</u>
<b>AT</b>	GAVIS PHARMS LLC	<u>0.01%</u>
<b>AT</b>	GLASSHOUSE PHARMS	<u>0.01%</u>
<b>AT</b>	TARO	<u>0.01%</u>

**A208386** **001** Oct 21, 2016  
**A088167** **001** Dec 16, 1982  
**A207441** **001** Sep 28, 2016  
**A206422** **001** Sep 02, 2015  
**A209596** **001** Dec 26, 2017  
**A089124** **001** Sep 11, 1985

**SYNALAR**

<b>AT</b>	+!	MEDIMETRIKS PHARMS	<u>0.01%</u>
-----------	----	--------------------	--------------

**N015296** **001**

FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM;TOPICAL

TRI-LUMA

+!	GALDERMA LABS LP	0.01%;4%;0.05%
----	------------------	----------------

N021112 001 Jan 18, 2002

FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM;TOPICAL

NEO-SYNALAR

!	MEDIMETRIKS PHARMS	0.025%;EQ 3.5MG BASE/GM
---	--------------------	-------------------------

A060700 001

FLUOCINONIDE

CREAM;TOPICAL

**FLUOCINONIDE**

<b>AB</b>	FOUGERA PHARMS INC	<u>0.1%</u>
<b>AB</b>	GLENMARK GENERICS	<u>0.1%</u>
<b>AB</b>	PERRIGO ISRAEL	<u>0.1%</u>
<b>AB</b>	TARO	<u>0.1%</u>

**A200735** **001** Jul 14, 2014  
**A091282** **001** Jul 14, 2014  
**A090256** **001** Jan 14, 2014  
**A200734** **001** Jul 14, 2014

**VANOS**

<b>AB</b>	+!	MEDICIS	<u>0.1%</u>
-----------	----	---------	-------------

**N021758** **001** Feb 11, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-186 (of 436)

FLUOCINONIDE

CREAM;TOPICAL

FLUOCINONIDE

<u>AB1</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A073030</u> <u>001</u>	Oct 17, 1994
<u>AB1</u>	G AND W LABS INC	<u>0.05%</u>	<u>A073085</u> <u>001</u>	Feb 14, 1992
<u>AB1</u>	TARO	<u>0.05%</u>	<u>A071500</u> <u>001</u>	Jun 10, 1987
<u>AB1</u> +!		<u>0.05%</u>	<u>N019117</u> <u>001</u>	Jun 26, 1984
<u>AB1</u>	TEVA	<u>0.05%</u>	<u>A072488</u> <u>001</u>	Feb 06, 1989

FLUOCINONIDE EMULSIFIED BASE

<u>AB2</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A076586</u> <u>001</u>	Jun 23, 2004
<u>AB2</u>	G AND W LABS INC	<u>0.05%</u>	<u>A074204</u> <u>001</u>	Jun 13, 1995
<u>AB2</u> !	TARO	<u>0.05%</u>	<u>A072494</u> <u>001</u>	Jan 19, 1989
<u>AB2</u>	TEVA	<u>0.05%</u>	<u>A072490</u> <u>001</u>	Feb 07, 1989

GEL;TOPICAL

FLUOCINONIDE

<u>AB</u> +	CNTY LINE PHARMS	<u>0.05%</u>	<u>N017373</u> <u>001</u>	
<u>AB</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072933</u> <u>001</u>	Dec 30, 1994
<u>AB</u>	G AND W LABS INC	<u>0.05%</u>	<u>A072537</u> <u>001</u>	Feb 07, 1989
<u>AB</u> !	TARO	<u>0.05%</u>	<u>A074935</u> <u>001</u>	Jul 29, 1997

OINTMENT;TOPICAL

FLUOCINONIDE

<u>AB</u>	FOUGERA PHARMS	<u>0.05%</u>	<u>A074905</u> <u>001</u>	Aug 26, 1997
<u>AB</u>	NOVEL LABS INC	<u>0.05%</u>	<u>A207538</u> <u>001</u>	Jul 31, 2017
<u>AB</u> !	TARO	<u>0.05%</u>	<u>A075008</u> <u>001</u>	Jun 30, 1999
<u>AB</u>	TEVA	<u>0.05%</u>	<u>A073481</u> <u>001</u>	Dec 27, 1991

LIDEX

<u>AB</u> +	CNTY LINE PHARMS	<u>0.05%</u>	<u>N016909</u> <u>002</u>	
-------------	------------------	--------------	---------------------------	--

SOLUTION;TOPICAL

FLUOCINONIDE

<u>AT</u>	FOUGERA PHARMS INC	<u>0.05%</u>	<u>A072934</u> <u>001</u>	Feb 27, 1995
<u>AT</u>	G AND W LABS INC	<u>0.05%</u>	<u>A071535</u> <u>001</u>	Dec 02, 1988
<u>AT</u>	NOVEL LABS INC	<u>0.05%</u>	<u>A206003</u> <u>001</u>	Jul 21, 2017
<u>AT</u> !	TARO	<u>0.05%</u>	<u>A074799</u> <u>001</u>	Dec 31, 1996
<u>AT</u>	TEVA	<u>0.05%</u>	<u>A072511</u> <u>001</u>	Feb 07, 1989

LIDEX

<u>AT</u> +	CNTY LINE PHARMS	<u>0.05%</u>	<u>N018849</u> <u>001</u>	Apr 06, 1984
-------------	------------------	--------------	---------------------------	--------------

FLUORESCIN SODIUM

INJECTABLE;INTRAVENOUS

AK-FLUOR 10%

<u>AP</u> +	AKORN	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N022186</u> <u>001</u>	Aug 08, 2008
-------------	-------	---	---------------------------	--------------

FLUORESCITE

<u>AP</u> +!	NOVARTIS PHARMS CORP	<u>EQ 500MG BASE/5ML (EQ 100MG BASE/ML)</u>	<u>N021980</u> <u>001</u>	Mar 28, 2006
AK-FLUOR 25%				

+! AKORN

EQ 500MG BASE/2ML (EQ 250MG BASE/ML)

N022186 002 Aug 08, 2008

FLUOROMETHOLONE

OINTMENT;OPHTHALMIC

FML

+! ALLERGAN

0.1%

N017760 001 Sep 04, 1985

SUSPENSION/DROPS;OPHTHALMIC

FML

+! ALLERGAN

0.1%

N016851 002 Jul 28, 1982

FML FORTE

ALLERGAN

0.25%

N019216 001 Apr 23, 1986

FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

FLAREX

+! NOVARTIS PHARMS

0.1%

N019079 001 Feb 11, 1986

CORP

FLUOROURACIL

CREAM;TOPICAL

CARAC

<u>AB</u> +!	VALEANT PHARMS NORTH	<u>0.5%</u>	<u>N020985</u> <u>001</u>	Oct 27, 2000
--------------	----------------------	-------------	---------------------------	--------------

EFUDEX

<u>AB</u> +!	VALEANT PHARM INTL	<u>5%</u>	<u>N016831</u> <u>003</u>	
--------------	--------------------	-----------	---------------------------	--

FLUOROURACIL

<u>AB</u>	MYLAN PHARMS INC	<u>0.5%</u>	<u>A203122</u> <u>001</u>	Apr 20, 2015
<u>AB</u>	SPEAR PHARMS	<u>5%</u>	<u>A077524</u> <u>001</u>	Apr 11, 2008
<u>AB</u>	TARO	<u>5%</u>	<u>A090368</u> <u>001</u>	Mar 05, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-187 (of 436)

**FLUOROURACIL**

CREAM;TOPICAL

FLUOROPLEX

+! AQUA PHARMS

1%

N016988 001

TOLAK

+!

HILL DERMACEUTICALS

4%

N022259 001 Sep 18, 2015

INJECTABLE;INJECTION

**FLUOROURACIL**

<b>AP</b>	!	ACCORD HLTHCARE	<u>500MG/10ML (50MG/ML)</u>	<b>A040743 002</b>	Apr 26, 2007
<b>AP</b>	!		<u>1GM/20ML (50MG/ML)</u>	<b>A040743 001</b>	Apr 26, 2007
<b>AP</b>	!		<u>2.5GM/50ML (50MG/ML)</u>	<b>A040798 002</b>	Apr 26, 2007
<b>AP</b>	!		<u>5GM/100ML (50MG/ML)</u>	<b>A040798 001</b>	Apr 26, 2007
<b>AP</b>	!	FRESENIUS KABI USA	<u>500MG/10ML (50MG/ML)</u>	<b>A040279 002</b>	Sep 30, 1998
<b>AP</b>	!		<u>1GM/20ML (50MG/ML)</u>	<b>A040279 001</b>	Sep 30, 1998
<b>AP</b>	!		<u>2.5GM/50ML (50MG/ML)</u>	<b>A040278 001</b>	Sep 30, 1998
<b>AP</b>	!		<u>5GM/100ML (50MG/ML)</u>	<b>A040278 002</b>	Sep 30, 1998
<b>AP</b>		GLAND PHARMA LTD	<u>500MG/10ML (50MG/ML)</u>	<b>A210123 001</b>	Oct 27, 2017
<b>AP</b>			<u>1GM/20ML (50MG/ML)</u>	<b>A210123 002</b>	Oct 27, 2017
<b>AP</b>			<u>2.5GM/50ML (50MG/ML)</u>	<b>A210124 001</b>	Dec 26, 2017
<b>AP</b>			<u>5GM/100ML (50MG/ML)</u>	<b>A210124 002</b>	Dec 26, 2017
<b>AP</b>		MYLAN LABS LTD	<u>500MG/10ML (50MG/ML)</u>	<b>A202668 001</b>	Jul 17, 2012
<b>AP</b>			<u>1GM/20ML (50MG/ML)</u>	<b>A202668 002</b>	Jul 17, 2012
<b>AP</b>			<u>2.5GM/50ML (50MG/ML)</u>	<b>A202669 001</b>	Jul 17, 2012
<b>AP</b>			<u>5GM/100ML (50MG/ML)</u>	<b>A202669 002</b>	Jul 17, 2012
<b>AP</b>		SAGENT PHARMS	<u>500MG/10ML (50MG/ML)</u>	<b>A203608 001</b>	May 11, 2017
<b>AP</b>			<u>1GM/20ML (50MG/ML)</u>	<b>A203608 002</b>	May 11, 2017
<b>AP</b>			<u>2.5GM/50ML (50MG/ML)</u>	<b>A203609 001</b>	Feb 17, 2016
<b>AP</b>			<u>5GM/100ML (50MG/ML)</u>	<b>A203609 002</b>	Feb 17, 2016
<b>AP</b>	!	TEVA PHARMS USA	<u>500MG/10ML (50MG/ML)</u>	<b>A040333 001</b>	Jan 27, 2000
<b>AP</b>	!		<u>2.5GM/50ML (50MG/ML)</u>	<b>A040334 001</b>	Feb 25, 2000
<b>AP</b>	!		<u>5GM/100ML (50MG/ML)</u>	<b>A040334 002</b>	Feb 25, 2000

SOLUTION;TOPICAL

**EFDUDEX**

<b>AT</b>	+	VALEANT PHARM INTL	<u>2%</u>	<b>N016831 001</b>
<b>AT</b>	+		<u>5%</u>	<b>N016831 002</b>

**FLUOROURACIL**

<b>AT</b>		TARO PHARM	<u>2%</u>	<b>A076526 001</b>	Nov 05, 2003
<b>AT</b>			<u>5%</u>	<b>A076526 002</b>	Nov 05, 2003

**FLUOXETINE HYDROCHLORIDE**

CAPSULE;ORAL

**FLUOXETINE HYDROCHLORIDE**

<b>AB</b>		ALEMBIC PHARMS LTD	<u>EQ 40MG BASE</u>	<b>A090223 003</b>	Mar 19, 2009
<b>AB</b>		AUROBINDO PHARMA	<u>EQ 40MG BASE</u>	<b>A078619 003</b>	Jan 31, 2008
<b>AB</b>		DR REDDYS LABS LTD	<u>EQ 40MG BASE</u>	<b>A075465 003</b>	Aug 02, 2001
<b>AB</b>		HERITAGE PHARMS INC	<u>EQ 40MG BASE</u>	<b>A201336 003</b>	Oct 01, 2012
<b>AB</b>		IVAX SUB TEVA PHARMS	<u>EQ 40MG BASE</u>	<b>A075245 003</b>	Sep 28, 2004
<b>AB</b>		PAR PHARM	<u>EQ 40MG BASE</u>	<b>A076922 003</b>	Dec 16, 2004
<b>AB</b>		SANDOZ	<u>EQ 40MG BASE</u>	<b>A075049 003</b>	Jan 29, 2002
<b>AB</b>		SCIEGEN PHARMS INC	<u>EQ 40MG BASE</u>	<b>A204597 003</b>	Mar 16, 2015
<b>AB</b>		SUN PHARM INDs LTD	<u>EQ 40MG BASE</u>	<b>A076990 001</b>	Dec 13, 2004
<b>AB</b>		TEVA	<u>EQ 40MG BASE</u>	<b>A075452 003</b>	Jan 29, 2002

**PROZAC**

<b>AB</b>	+	ELI LILLY AND CO	<u>EQ 40MG BASE</u>	<b>N018936 003</b>	Jun 15, 1999
-----------	---	------------------	---------------------	--------------------	--------------

**FLUOXETINE HYDROCHLORIDE**

<b>AB1</b>		ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<b>A090223 001</b>	Mar 19, 2009
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A090223 002</b>	Mar 19, 2009
<b>AB1</b>		AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<b>A078619 001</b>	Jan 31, 2008
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A078619 002</b>	Jan 31, 2008
<b>AB1</b>		BARR	<u>EQ 10MG BASE</u>	<b>A074803 002</b>	Jan 30, 2002
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A074803 001</b>	Aug 02, 2001
<b>AB1</b>		DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<b>A075465 001</b>	Jan 29, 2002
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A075465 002</b>	Jan 29, 2002
<b>AB1</b>		HERITAGE PHARMS INC	<u>EQ 10MG BASE</u>	<b>A201336 001</b>	Oct 01, 2012
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A201336 002</b>	Oct 01, 2012
<b>AB1</b>		IVAX SUB TEVA PHARMS	<u>EQ 10MG BASE</u>	<b>A075245 002</b>	Jan 31, 2002
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A075245 001</b>	Jan 31, 2002
<b>AB1</b>		LANDELA PHARM	<u>EQ 10MG BASE</u>	<b>A075464 001</b>	Jan 30, 2002
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A075464 002</b>	Jan 30, 2002
<b>AB1</b>		SANDOZ	<u>EQ 10MG BASE</u>	<b>A075049 001</b>	Aug 02, 2001
<b>AB1</b>			<u>EQ 20MG BASE</u>	<b>A075049 002</b>	Jan 29, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-188 (of 436)

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB1</u>	SCIEGEN PHARMS INC	<u>EQ 10MG BASE</u>	<u>A204597 001</u>	Mar 16, 2015
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A204597 002</u>	Mar 16, 2015
<u>AB1</u>	SPECGX LLC	<u>EQ 10MG BASE</u>	<u>A075658 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075658 002</u>	Jan 29, 2002
<u>AB1</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075452 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A075452 002</u>	Jan 29, 2002
<u>AB1</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A076001 001</u>	Jan 29, 2002
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A076001 002</u>	Jan 29, 2002

PROZAC

<u>AB1</u> +	ELI LILLY AND CO	<u>EQ 10MG BASE</u>	<u>N018936 006</u>	Dec 23, 1992
<u>AB1</u> +		<u>EQ 20MG BASE</u>	<u>N018936 001</u>	Dec 29, 1987

FLUOXETINE HYDROCHLORIDE

MYLAN	<u>EQ 10MG BASE</u>
!	<u>EQ 20MG BASE</u>

CAPSULE, DELAYED REL PELLETS; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	BARR	<u>EQ 90MG BASE</u>	<u>A076237 001</u>	Mar 24, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 90MG BASE</u>	<u>A078572 001</u>	Mar 22, 2010

PROZAC WEEKLY

<u>AB</u> +!	LILLY	<u>EQ 90MG BASE</u>	<u>N021235 001</u>	Feb 26, 2001
	SOLUTION; ORAL			

FLUOXETINE HYDROCHLORIDE

<u>AA</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE/5ML</u>	<u>A079209 001</u>	Mar 20, 2009
<u>AA</u> !	PHARM ASSOC	<u>EQ 20MG BASE/5ML</u>	<u>A076015 001</u>	Jan 30, 2002
<u>AA</u>	SILARX	<u>EQ 20MG BASE/5ML</u>	<u>A077849 001</u>	Feb 09, 2007
<u>AA</u>	SPECGX LLC	<u>EQ 20MG BASE/5ML</u>	<u>A075920 001</u>	Jan 29, 2002
<u>AA</u>	TEVA	<u>EQ 20MG BASE/5ML</u>	<u>A075506 001</u>	Aug 02, 2001
<u>AA</u>	WOCKHARDT BIO AG	<u>EQ 20MG BASE/5ML</u>	<u>A075514 001</u>	Aug 29, 2002

TABLET; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A208698 001</u>	Apr 05, 2017
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A208698 002</u>	Apr 05, 2017
<u>AB</u> +!	ALVOGEN	<u>EQ 60MG BASE</u>	<u>N202133 001</u>	Oct 06, 2011
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 10MG BASE</u>	<u>A076006 001</u>	Jan 30, 2002
<u>AB</u>	INVENTIA HLTHCARE	<u>EQ 60MG BASE</u>	<u>A209695 001</u>	Nov 20, 2017
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A075755 001</u>	Aug 02, 2001
<u>AB</u> !		<u>EQ 20MG BASE</u>	<u>A075755 002</u>	Aug 02, 2001
<u>AB</u>	PAR FORM	<u>EQ 10MG BASE</u>	<u>A203836 001</u>	Aug 19, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203836 002</u>	Aug 19, 2016
<u>AB</u>	PAR PHARM INC	<u>EQ 60MG BASE</u>	<u>A209419 001</u>	Nov 16, 2017
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A075872 001</u>	Jan 29, 2002
<u>AB1</u>	TORRENT PHARMS LTD	<u>EQ 10MG BASE</u>	<u>A206937 001</u>	Oct 21, 2016
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A206937 002</u>	Oct 21, 2016

SARAFEM

<u>AB1</u> +	APIL	<u>EQ 10MG BASE</u>	<u>N021860 001</u>	May 19, 2006
<u>AB1</u> +		<u>EQ 15MG BASE</u>	<u>N021860 002</u>	May 19, 2006
<u>AB1</u> +!		<u>EQ 20MG BASE</u>	<u>N021860 003</u>	May 19, 2006

SELFEMRA

<u>AB1</u>	TEVA PHARMS USA	<u>EQ 10MG BASE</u>	<u>A200151 001</u>	Feb 03, 2014
<u>AB1</u>		<u>EQ 15MG BASE</u>	<u>A200151 002</u>	Feb 03, 2014
<u>AB1</u>		<u>EQ 20MG BASE</u>	<u>A200151 003</u>	Feb 03, 2014

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

<u>AB</u>	PAR PHARM	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A077742 001</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077742 002</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077742 003</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077742 004</u>	Nov 02, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077742 005</u>	Nov 02, 2012
<u>AB</u>	SANDOZ	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A078901 005</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A078901 001</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A078901 003</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A078901 002</u>	Nov 16, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A078901 004</u>	Nov 16, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>A202074 001</u>	Mar 25, 2013
<u>AB</u>		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>A077528 001</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>A077528 002</u>	Jun 19, 2012
<u>AB</u>		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>A077528 003</u>	Jun 19, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-189 (of 436)

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

OLANZAPINE AND FLUOXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>A077528 004</u>	Jun 19, 2012
	<u>SYMBYAX</u>			
<u>AB</u>	+	LILLY	<u>EQ 25MG BASE;EQ 3MG BASE</u>	<u>N021520 001</u> Apr 09, 2007
<u>AB</u>	+		<u>EQ 25MG BASE;EQ 6MG BASE</u>	<u>N021520 002</u> Dec 24, 2003
<u>AB</u>	+		<u>EQ 25MG BASE;EQ 12MG BASE</u>	<u>N021520 004</u> Dec 24, 2003
<u>AB</u>	+		<u>EQ 50MG BASE;EQ 6MG BASE</u>	<u>N021520 003</u> Dec 24, 2003
<u>AB</u>	+		<u>EQ 50MG BASE;EQ 12MG BASE</u>	<u>N021520 005</u> Dec 24, 2003

FLUOXYMESTERONE

TABLET; ORAL

FLUOXYMESTERONE

! USL PHARMA

10MG

A088342 001 Oct 21, 1983

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

<u>AO</u>	AUROBINDO PHARMA LTD	<u>25MG/ML</u>	<u>A207739 001</u>	Oct 17, 2017
<u>AO</u>	! FRESENIUS KABI USA	<u>25MG/ML</u>	<u>A071413 001</u>	Jul 14, 1987
<u>AO</u>	PAR STERILE PRODUCTS	<u>25MG/ML</u>	<u>A203732 001</u>	Jul 03, 2014
<u>AO</u>	WEST-WARD PHARMS INT	<u>25MG/ML</u>	<u>A074531 001</u>	Aug 30, 1996

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

5MG/ML

A074725 001 Sep 16, 1996

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

! PHARM ASSOC

2.5MG/5ML

A040146 001 Aug 21, 1996

INJECTABLE; INJECTION

FLUPHENAZINE HYDROCHLORIDE

! FRESENIUS KABI USA

2.5MG/ML

A089556 001 Apr 16, 1987

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>	LANNETT	<u>1MG</u>	<u>A089743 002</u>	Aug 25, 1988
<u>AB</u>		<u>2.5MG</u>	<u>A089743 003</u>	Aug 25, 1988
<u>AB</u>		<u>5MG</u>	<u>A089743 004</u>	Aug 25, 1988
<u>AB</u>		<u>10MG</u>	<u>A089743 001</u>	Aug 25, 1988
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A089804 002</u>	Aug 12, 1988
<u>AB</u>		<u>2.5MG</u>	<u>A089804 003</u>	Aug 12, 1988
<u>AB</u>		<u>5MG</u>	<u>A089804 004</u>	Aug 12, 1988
<u>AB</u>	!	<u>10MG</u>	<u>A089804 001</u>	Aug 12, 1988
<u>AB</u>	SANDOZ	<u>1MG</u>	<u>A089586 002</u>	Oct 16, 1987
<u>AB</u>		<u>2.5MG</u>	<u>A089586 003</u>	Oct 16, 1987
<u>AB</u>		<u>5MG</u>	<u>A089586 004</u>	Oct 16, 1987
<u>AB</u>		<u>10MG</u>	<u>A089586 001</u>	Oct 16, 1987

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

<u>AT</u>	! AQUA PHARMS	<u>0.05%</u>	<u>N012806 002</u>	
<u>AT</u>	CINTEX SVCS	<u>0.05%</u>	<u>A205342 001</u>	Apr 13, 2016

CORDRAN SP

+! AQUA PHARMS

0.025%

N012806 003

LOTION; TOPICAL

CORDRAN

<u>AT</u>	! AQUA PHARMS	<u>0.05%</u>	<u>N013790 001</u>	
<u>AT</u>	CINTEX SVCS	<u>0.05%</u>	<u>A205343 001</u>	Dec 22, 2016

PERRIGO UK FINCO

0.05%

A207133 001 Aug 30, 2016

OINTMENT; TOPICAL

CORDRAN

<u>AT</u>	! AQUA PHARMS	<u>0.05%</u>	<u>N012806 001</u>	
<u>AT</u>	FLURANDRENOLIDE		<u>A207851 001</u>	Dec 30, 2016

TELIGENT PHARMA INC

0.05%

A207851 001 Dec 30, 2016

TAPE; TOPICAL

CORDRAN

+! ALLERGAN SALES LLC

0.004MG/SQ CM

N016455 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-190 (of 436)

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL  
 FLURAZEPAM HYDROCHLORIDE  
 MYLAN PHARMS INC  
 !  
 15MG  
 30MG

A070345 002 Nov 27, 1985  
 A070345 001 Nov 27, 1985

FLURBIPROFEN

TABLET; ORAL

**FLURBIPROFEN**

<b>AB</b>	MYLAN	<b>50MG</b>	
<b>AB</b>	!	<b>100MG</b>	
<b>AB</b>	SUN PHARM INDNS INC	<b>50MG</b>	
<b>AB</b>		<b>100MG</b>	
<b>AB</b>	TEVA	<b>100MG</b>	

<b>A074358 001</b>	Jun 20, 1994
<b>A074358 002</b>	Jun 20, 1994
<b>A075058 001</b>	Apr 27, 2001
<b>A075058 002</b>	Apr 27, 2001
<b>A074431 001</b>	May 31, 1995

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

**FLURBIPROFEN SODIUM**

<b>AT</b>	BAUSCH AND LOMB	<b>0.03%</b>	
<b>OCUFEN</b>			<b>A074447 001</b> Jan 04, 1995
<b>AT</b>	+!	ALLERGAN	<b>N019404 001</b> Dec 31, 1986

FLUTAMIDE

CAPSULE; ORAL

**FLUTAMIDE**

<b>AB</b>	ACTAVIS LABS FL INC	<b>125MG</b>	
<b>AB</b>	!	<b>125MG</b>	
<b>AB</b>	PAR PHARM	<b>125MG</b>	

<b>A075820 001</b>	Sep 18, 2001
<b>A075780 001</b>	Sep 19, 2001
<b>A075298 001</b>	Sep 18, 2001

FLUTEMETAMOL F-18

INJECTABLE; INTRAVENOUS

VIZAMYL		
+!	GE HEALTHCARE	40.5mCi/10ML (4.05mCi/ML)
+!		121.5mCi/30ML (4.05mCi/ML)

N203137 001	Oct 25, 2013
N203137 002	Oct 25, 2013

FLUTICASONE FUROATE

POWDER; INHALATION

ARNUNITY ELLIPTA		
+!	GLAXOSMITHKLINE	0.1MG/INH
+!		0.2MG/INH

N205625 001	Aug 20, 2014
N205625 002	Aug 20, 2014

FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER; INHALATION

TRELEGY ELLIPTA		
+!	GLAXOSMITHKLINE	0.1MG/INH; EQ 0.0625MG BASE/INH; EQ 0.025MG BASE/INH

N209482 001	Sep 18, 2017
-------------	--------------

FLUTICASONE FUROATE; VILANTEROL TRIFENATATE

POWDER; INHALATION

BREO ELLIPTA		
+!	GLAXO GRP LTD	0.1MG/INH; EQ 0.025MG BASE/INH
+!		0.2MG/INH; EQ 0.025MG BASE/INH

N204275 001	May 10, 2013
N204275 002	Apr 30, 2015

FLUTICASONE PROPIONATE

AEROSOL, METERED; INHALATION

FLOVENT HFA		
+!	GLAXO GRP LTD	0.044MG/INH
+!		0.11MG/INH
+!		0.22MG/INH

N021433 003	May 14, 2004
N021433 002	May 14, 2004
N021433 001	May 14, 2004

CREAM; TOPICAL

**FLUTICASONE PROPIONATE**

<b>AB</b>	FOUGERA PHARMS	<b>0.05%</b>	
<b>AB</b>	G AND W LABS	<b>0.05%</b>	
<b>AB</b>	!	<b>PERRIGO NEW YORK</b>	<b>0.05%</b>
<b>AB</b>	TOLMAR	<b>0.05%</b>	

<b>A076451 001</b>	May 14, 2004
<b>A077055 001</b>	Jun 30, 2006
<b>A076793 001</b>	May 14, 2004
<b>A076633 001</b>	May 14, 2004

LOTION; TOPICAL

**CUTIVATE**

<b>AB</b>	+!	FOUGERA PHARMS	<b>0.05%</b>	
<b>FLUTICASONE PROPIONATE</b>				

<b>N021152 001</b>	Mar 31, 2005
--------------------	--------------

<b>AB</b>	GLENMARK GENERICS	<b>0.05%</b>	
<b>AB</b>	PERRIGO ISRAEL	<b>0.05%</b>	

<b>A090759 001</b>	May 02, 2011
<b>A091553 001</b>	Jul 30, 2013

OINTMENT; TOPICAL

**CUTIVATE**

<b>AB</b>	+!	FOUGERA PHARMS	<b>0.005%</b>	
<b>FLUTICASONE PROPIONATE</b>				

<b>N019957 001</b>	Dec 14, 1990
--------------------	--------------

<b>AB</b>	FOUGERA PHARMS	<b>0.005%</b>	
<b>AB</b>	G AND W LABS	<b>0.005%</b>	

<b>A076300 001</b>	May 14, 2004
<b>A077168 001</b>	Mar 03, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-191 (of 436)

**FLUTICASONE PROPIONATE**

OINTMENT;TOPICAL

**FLUTICASONE PROPIONATE**

<b>AB</b>	PERRIGO NEW YORK	<b><u>0.005%</u></b>	<b>A076668 001</b>	May 14, 2004
POWDER;INHALATION				
ARMONAIR RESPICLICK				
+ TEVA PHARM	0.055MG/INH		N208798 001	Jan 27, 2017
+!	0.113MG/INH		N208798 002	Jan 27, 2017
+!	0.232MG/INH		N208798 003	Jan 27, 2017
FLOVENT DISKUS 100				
+! GLAXO GRP LTD	0.1MG/INH		N020833 002	Sep 29, 2000
FLOVENT DISKUS 250				
+! GLAXO GRP LTD	0.25MG/INH		N020833 003	Sep 29, 2000
FLOVENT DISKUS 50				
+! GLAXO GRP LTD	0.05MG/INH		N020833 001	Sep 29, 2000
SPRAY, METERED;NASAL				
<b><u>FLUTICASONE PROPIONATE</u></b>				
<b>AB</b>	APOTEX INC	<b><u>0.05MG/SPRAY</u></b>	<b>A077538 001</b>	Sep 12, 2007
<b>AB</b>	HI TECH PHARMA	<b><u>0.05MG/SPRAY</u></b>	<b>A077570 001</b>	Jan 16, 2008
<b>AB</b> !	WEST-WARD PHARMS INT	<b><u>0.05MG/SPRAY</u></b>	<b>A076504 001</b>	Feb 22, 2006
<b>AB</b>	WOCKHARDT BIO AG	<b><u>0.05MG/SPRAY</u></b>	<b>A078492 001</b>	Jan 09, 2012
XHANCE				
+! OPTNOSE US	0.093MG		N209022 001	Sep 18, 2017

**FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE**

AEROSOL, METERED;INHALATION

ADVAIR HFA				
+! GLAXO GRP LTD	0.045MG/INH;EQ 0.021MG BASE/INH		N021254 001	Jun 08, 2006
+!	0.115MG/INH;EQ 0.021MG BASE/INH		N021254 002	Jun 08, 2006
+!	0.23MG/INH;EQ 0.021MG BASE/INH		N021254 003	Jun 08, 2006
POWDER;INHALATION				
ADVAIR DISKUS 100/50				
+! GLAXO GRP LTD	0.1MG/INH;EQ 0.05MG BASE/INH		N021077 001	Aug 24, 2000
ADVAIR DISKUS 250/50				
+! GLAXO GRP LTD	0.25MG/INH;EQ 0.05MG BASE/INH		N021077 002	Aug 24, 2000
ADVAIR DISKUS 500/50				
+! GLAXO GRP LTD	0.5MG/INH;EQ 0.05MG BASE/INH		N021077 003	Aug 24, 2000
AIRDUO RESPICLICK				
+ TEVA PHARM	0.055MG/INH;EQ 0.014MG BASE/INH		N208799 001	Jan 27, 2017
+!	0.113MG/INH;EQ 0.014MG BASE/INH		N208799 002	Jan 27, 2017
+!	0.232MG/INH;EQ 0.014MG BASE/INH		N208799 003	Jan 27, 2017

**FLUVASTATIN SODIUM**

CAPSULE;ORAL

**FLUVASTATIN SODIUM**

<b>AB</b>	MYLAN PHARMS INC	<b><u>EQ 20MG BASE</u></b>	<b>A090595 001</b>	Apr 11, 2012
<b>AB</b> !		<b><u>EQ 40MG BASE</u></b>	<b>A090595 002</b>	Apr 11, 2012
<b>AB</b>	TEVA PHARMS	<b><u>EQ 20MG BASE</u></b>	<b>A078407 001</b>	Jun 12, 2012
<b>AB</b>		<b><u>EQ 40MG BASE</u></b>	<b>A078407 002</b>	Jun 12, 2012
TABLET, EXTENDED RELEASE;ORAL				
<b><u>FLUVASTATIN SODIUM</u></b>				
<b>AB</b>	MYLAN PHARMS INC	<b><u>EQ 80MG BASE</u></b>	<b>A202458 001</b>	Sep 11, 2015
<b>AB</b>	TEVA PHARMS USA	<b><u>EQ 80MG BASE</u></b>	<b>A079011 001</b>	Jan 27, 2016
<b><u>LESCOL XL</u></b>				
<b>AB</b> +! NOVARTIS		<b><u>EQ 80MG BASE</u></b>	<b>N021192 001</b>	Oct 06, 2000

**FLUVOXAMINE MALEATE**

CAPSULE, EXTENDED RELEASE;ORAL

**FLUVOXAMINE MALEATE**

<b>AB</b>	ACTAVIS ELIZABETH	<b><u>100MG</u></b>	<b>A091482 001</b>	Apr 23, 2013
<b>AB</b> !		<b><u>150MG</u></b>	<b>A091482 002</b>	Nov 18, 2013
<b>AB</b>	ANCHEM PHARMS	<b><u>100MG</u></b>	<b>A091476 001</b>	Mar 13, 2013
<b>AB</b>		<b><u>150MG</u></b>	<b>A091476 002</b>	Mar 13, 2013
<b>AB</b>	TORRENT PHARMS LTD	<b><u>100MG</u></b>	<b>A203240 001</b>	Oct 31, 2014
<b>AB</b>		<b><u>150MG</u></b>	<b>A203240 002</b>	Oct 31, 2014
TABLET;ORAL				
<b><u>FLUVOXAMINE MALEATE</u></b>				
<b>AB</b>	ANI PHARMS INC	<b><u>25MG</u></b>	<b>A075897 001</b>	Jan 25, 2001
<b>AB</b>		<b><u>50MG</u></b>	<b>A075897 002</b>	Jan 25, 2001
<b>AB</b>		<b><u>100MG</u></b>	<b>A075897 003</b>	Jan 25, 2001
<b>AB</b>	APOTEX	<b><u>25MG</u></b>	<b>A075902 001</b>	May 07, 2001
<b>AB</b>		<b><u>50MG</u></b>	<b>A075902 002</b>	May 07, 2001
<b>AB</b>		<b><u>100MG</u></b>	<b>A075902 003</b>	May 07, 2001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-192 (of 436)

FLUVOXAMINE MALEATE

TABLET;ORAL

FLUVOXAMINE MALEATE

<u>AB</u>	MYLAN	<u>25MG</u>	<u>A075889 001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>A075889 002</u>	Nov 29, 2000
<u>AB</u>		<u>100MG</u>	<u>A075889 003</u>	Nov 29, 2000
<u>AB</u>	TEVA	<u>25MG</u>	<u>A075893 001</u>	Sep 10, 2002
<u>AB</u>		<u>50MG</u>	<u>A075893 002</u>	Sep 10, 2002
<u>AB</u>		<u>100MG</u>	<u>A075893 003</u>	Sep 10, 2002
<u>AB</u>	UPSHER-SMITH LABS	<u>25MG</u>	<u>A075888 001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>A075888 002</u>	Nov 29, 2000
<u>AB</u>	!	<u>100MG</u>	<u>A075888 003</u>	Nov 29, 2000
<u>LUVOX</u>				
<u>AB</u>	ANI PHARMS	<u>25MG</u>	<u>N021519 001</u>	Dec 20, 2007
<u>AB</u>		<u>50MG</u>	<u>N021519 002</u>	Dec 20, 2007
<u>AB</u>		<u>100MG</u>	<u>N021519 003</u>	Dec 20, 2007

FOLIC ACID

INJECTABLE;INJECTION

FOLIC ACID

! FRESENIUS KABI USA 5MG/ML

A089202 001 Feb 18, 1986

TABLET;ORAL

FOLIC ACID

<u>AA</u>	AIPING PHARM INC	<u>1MG</u>	<u>A091145 001</u>	Jul 12, 2013
<u>AA</u>	! AMNEAL PHARM	<u>1MG</u>	<u>A040625 001</u>	Jul 21, 2005
<u>AA</u>	CADILA PHARMS LTD	<u>1MG</u>	<u>A202437 001</u>	Jan 27, 2014
<u>AA</u>	HIKMA PHARMS	<u>1MG</u>	<u>A080600 001</u>	
<u>AA</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A090035 001</u>	Jun 09, 2009
<u>AA</u>	LEADING PHARMA LLC	<u>1MG</u>	<u>A040796 001</u>	Jan 12, 2009
<u>AA</u>	NUVO PHARM INC	<u>1MG</u>	<u>A204418 001</u>	Jul 28, 2015
<u>AA</u>	VINTAGE	<u>1MG</u>	<u>A040756 001</u>	Jun 04, 2010
<u>AA</u>	! WATSON LABS	<u>1MG</u>	<u>A080680 001</u>	

FOLLITROPIN ALFA/BETA

INJECTABLE;SUBCUTANEOUS

FOLLISTIM AQ

+! ORGANON USA INC 300 IU/0.36ML  
+! 600 IU/0.72ML  
+! 900 IU/1.08ML

N021211 001 Mar 23, 2004  
N021211 002 Mar 23, 2004  
N021211 004 Feb 11, 2005

GONAL-F

+! EMD SERONO 450 IU/VIAL  
+ 1,050 IU/VIAL

N020378 005 Mar 26, 2004  
N020378 004 Feb 28, 2001

GONAL-F RFF

+! EMD SERONO 75 IU/VIAL

N021765 002 Mar 25, 2004

GONAL-F RFF REDI-JECT

+! EMD SERONO 300 IU/0.5ML  
+! 450 IU/0.75ML  
+! 900 IU/1.5ML

N021684 001 May 25, 2004  
N021684 002 May 25, 2004  
N021684 003 May 25, 2004

FOMEPIZOLE

INJECTABLE;INJECTION

ANTIZOL

<u>AP</u>	+! PAR PHARM INC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>N020696 001</u>	Dec 04, 1997
-----------	------------------	-----------------------------	--------------------	--------------

FOMEPIZOLE

<u>AP</u>	LUITPOLD	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078368 001</u>	Dec 14, 2007
<u>AP</u>	MYLAN INSTITUTIONAL	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078639 001</u>	Mar 03, 2008
<u>AP</u>	NAVINTA LLC	<u>1.5GM/1.5ML (1GM/ML)</u>	<u>A078537 001</u>	Mar 06, 2008

FONDAPARINUX SODIUM

INJECTABLE;SUBCUTANEOUS

ARIIXTRA

<u>AP</u>	+! MYLAN IRELAND LTD	<u>2.5MG/0.5ML</u>	<u>N021345 001</u>	Dec 07, 2001
<u>AP</u>	+!	<u>5MG/0.4ML</u>	<u>N021345 002</u>	May 28, 2004
<u>AP</u>	+!	<u>7.5MG/0.6ML</u>	<u>N021345 003</u>	May 28, 2004
<u>AP</u>	+!	<u>10MG/0.8ML</u>	<u>N021345 004</u>	May 28, 2004

FONDAPARINUX SODIUM

<u>AP</u>	AUROBINDO PHARMA LTD	<u>2.5MG/0.5ML</u>	<u>A206918 001</u>	Dec 26, 2017
<u>AP</u>		<u>5MG/0.4ML</u>	<u>A206918 002</u>	Dec 26, 2017
<u>AP</u>		<u>7.5MG/0.6ML</u>	<u>A206918 003</u>	Dec 26, 2017
<u>AP</u>		<u>10MG/0.8ML</u>	<u>A206918 004</u>	Dec 26, 2017
<u>AP</u>	DR REDDYS LABS LTD	<u>2.5MG/0.5ML</u>	<u>A091316 001</u>	Jul 11, 2011
<u>AP</u>		<u>5MG/0.4ML</u>	<u>A091316 002</u>	Jul 11, 2011
<u>AP</u>		<u>7.5MG/0.6ML</u>	<u>A091316 003</u>	Jul 11, 2011
<u>AP</u>		<u>10MG/0.8ML</u>	<u>A091316 004</u>	Jul 11, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-193 (of 436)

FORMOTEROL FUMARATE

SOLUTION; INHALATION  
 PERFOROMIST  
 +! MYLAN SPECLT 0.02MG/2ML N022007 001 May 11, 2007

FORMOTEROL FUMARATE; GLYCOPYRROLATE

AEROSOL, METERED; INHALATION  
 BEVESPI AEROSPHERE  
 +! ASTRazeneca PHARMS 0.0048MG/INH;0.0090MG/INH N208294 001 Apr 25, 2016

FORMOTEROL FUMARATE; MOMETASONE FUBRATE

AEROSOL, METERED; INHALATION  
 DULERA  
 +! MERCK SHARP DOHME 0.005MG/INH;0.1MG/INH N022518 001 Jun 22, 2010  
 +! 0.005MG/INH;0.2MG/INH N022518 002 Jun 22, 2010

FOSAMPRENAVIR CALCIUM

SUSPENSION; ORAL  
 LEXIVA  
 +! VIIV HLTHCARE EQ 50MG BASE/ML N022116 001 Jun 14, 2007

TABLET; ORAL  
FOSAMPRENAVIR CALCIUM  
AB MYLAN PHARMS INC EQ 700MG BASE A204060 001 Apr 15, 2016  
LEXIVA  
AB +! VIIV HLTHCARE EQ 700MG BASE N021548 001 Oct 20, 2003

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS  
EMEND  
AP +! MERCK AND CO INC EQ 150MG BASE/VIAL N022023 002 Nov 12, 2010  
FOSAPREPITANT DIMEGLUMINE  
AP FRESENIUS KABI USA EQ 150MG BASE/VIAL A206197 001 Jun 09, 2016

FOSCARNET SODIUM

INJECTABLE; INJECTION  
 FOSCAVIR  
 +! CLINIGEN HLTHCARE 2.4GM/100ML N020068 001 Sep 27, 1991

FOSFOMYCIN TROMETHAMINE

FOR SUSPENSION; ORAL  
 MONUROL  
 +! ZAMBON SPA EQ 3GM BASE/PACKET N050717 001 Dec 19, 1996

FOSINOPRIL SODIUM

TABLET; ORAL  
FOSINOPRIL SODIUM  
AB APOTEX INC 10MG A076906 001 May 17, 2005  
AB 20MG A076906 002 May 17, 2005  
AB 40MG A076906 003 May 17, 2005  
AB AUROBINDO PHARMA LTD 10MG A091163 001 Mar 30, 2011  
AB 20MG A091163 002 Mar 30, 2011  
AB 40MG A091163 003 Mar 30, 2011  
AB INVAGEN PHARMS 10MG A077222 001 Apr 20, 2005  
AB 20MG A077222 002 Apr 20, 2005  
AB 40MG A077222 003 Apr 20, 2005  
AB PRINSTON INC 10MG A205670 001 Aug 29, 2016  
AB 20MG A205670 002 Aug 29, 2016  
AB 40MG A205670 003 Aug 29, 2016  
AB TEVA 10MG A076139 001 Nov 25, 2003  
AB 20MG A076139 002 Nov 25, 2003  
AB ! 40MG A076139 003 Nov 25, 2003  
AB UPSHER-SMITH LABS 10MG A076483 001 Apr 23, 2004  
AB 20MG A076483 002 Apr 23, 2004  
AB 40MG A076483 003 Apr 23, 2004

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL  
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE  
AB AUROBINDO PHARMA 10MG;12.5MG A079245 001 Jul 09, 2009  
AB 20MG;12.5MG A079245 002 Jul 09, 2009  
AB EMCURE PHARMS INDIA 10MG;12.5MG A079025 001 Sep 17, 2010  
AB ! 20MG;12.5MG A079025 002 Sep 17, 2010  
AB INVAGEN PHARMS 10MG;12.5MG A090228 001 Jul 09, 2009  
AB 20MG;12.5MG A090228 002 Jul 09, 2009  
AB SANDOZ 10MG;12.5MG A076961 001 Sep 28, 2005  
AB 20MG;12.5MG A076961 002 Sep 28, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-194 (of 436)

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYX

<u>AP</u>	+!	PARKE DAVIS	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>N020450 001</u>	Aug 05, 1996
<u>FOSPHENYTOIN SODIUM</u>					
<u>AP</u>		AMNEAL PHARMS CO	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078476 001</u>	Mar 18, 2008
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078052 001</u>	Aug 06, 2007
<u>AP</u>		HIKMA FARMACEUTICA	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078765 001</u>	Dec 02, 2009
<u>AP</u>		LUITPOLD	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078277 001</u>	Aug 06, 2007
<u>AP</u>		MYLAN LABS LTD	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A090099 001</u>	May 13, 2010
<u>AP</u>		SUN PHARMA GLOBAL	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078736 001</u>	Jun 08, 2010
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078417 001</u>	Mar 18, 2008
<u>AP</u>		WOCKHARDT	<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077481 001</u>	Aug 06, 2007
<u>AP</u>			<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A077989 001</u>	Aug 06, 2007
<u>AP</u>			<u>EQ 50MG PHENYTOIN NA/ML</u>	<u>A078137 001</u>	Aug 06, 2007

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

<u>AB</u>	+!	ENDO PHARMS	<u>EQ 2.5MG BASE</u>	<u>N021006 001</u>	Nov 08, 2001
<u>FROVATRIPTAN SUCCINATE</u>					
<u>AB</u>		GLENMARK PHARMS LTD	<u>EQ 2.5MG BASE</u>	<u>A204730 001</u>	Mar 11, 2016
<u>AB</u>		MYLAN PHARMS INC	<u>EQ 2.5MG BASE</u>	<u>A202931 001</u>	Aug 28, 2014

FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FASLODEX

+! ASTRAZENECA

50MG/ML

N021344 001 Apr 25, 2002

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>		AMNEAL PHARMS CO	<u>10MG/ML</u>	<u>A207552 001</u>	Jul 20, 2016
<u>AP</u>		BAXTER HLTHCARE CORP	<u>10MG/ML</u>	<u>A202747 001</u>	Jan 27, 2014
<u>AP</u>		EMCURE PHARMS LTD	<u>10MG/ML</u>	<u>A203428 001</u>	Aug 26, 2014
<u>AP</u>		FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N018902 001</u>	May 22, 1984
<u>AP</u>		HOSPIRA	<u>10MG/ML</u>	<u>A075241 001</u>	May 28, 1999
<u>AP</u>			<u>10MG/ML</u>	<u>N018667 001</u>	May 28, 1982
<u>AP</u>		WOCKHARDT	<u>10MG/ML</u>	<u>A077941 001</u>	Mar 22, 2007

SOLUTION; ORAL

FUROSEMIDE

<u>AA</u>	!	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A070434 001</u>	Apr 22, 1987
<u>AA</u>		WOCKHARDT BIO AG WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A070655 001</u>	Oct 02, 1987

TABLET; ORAL

FUROSEMIDE

<u>AB</u>		IPCA LABS LTD	<u>20MG</u>	<u>A078010 001</u>	Sep 18, 2006
<u>AB</u>			<u>40MG</u>	<u>A078010 002</u>	Sep 18, 2006
<u>AB</u>			<u>80MG</u>	<u>A078010 003</u>	Sep 18, 2006
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>20MG</u>	<u>N018413 001</u>	Nov 30, 1983
<u>AB</u>			<u>40MG</u>	<u>N018413 002</u>	Nov 30, 1983
<u>AB</u>		LEADING PHARMA LLC	<u>20MG</u>	<u>A077293 001</u>	Nov 09, 2005
<u>AB</u>			<u>40MG</u>	<u>A077293 002</u>	Nov 09, 2005
<u>AB</u>			<u>80MG</u>	<u>A077293 003</u>	Nov 09, 2005
<u>AB</u>		MYLAN	<u>20MG</u>	<u>N018487 001</u>	
<u>AB</u>			<u>40MG</u>	<u>N018487 002</u>	
<u>AB</u>			<u>80MG</u>	<u>A070082 001</u>	Oct 29, 1986
<u>AB</u>		PRINSTON INC	<u>20MG</u>	<u>A076796 001</u>	Mar 26, 2004
<u>AB</u>			<u>40MG</u>	<u>A076796 002</u>	Mar 26, 2004
<u>AB</u>			<u>80MG</u>	<u>A076796 003</u>	Mar 26, 2004
<u>AB</u>		SANDOZ	<u>20MG</u>	<u>N018569 002</u>	
<u>AB</u>			<u>40MG</u>	<u>N018569 001</u>	
<u>AB</u>			<u>80MG</u>	<u>N018569 005</u>	Aug 14, 1984
<u>AB</u>		WEST-WARD PHARMS INT	<u>20MG</u>	<u>N018823 001</u>	Nov 10, 1983
<u>AB</u>			<u>40MG</u>	<u>N018823 002</u>	Nov 10, 1983
<u>AB</u>			<u>80MG</u>	<u>A070086 001</u>	Jan 24, 1986
<u>LASIX</u>					
<u>AB</u>	+	US PHARM HOLDINGS	<u>20MG</u>	<u>N016273 002</u>	
<u>AB</u>	+		<u>40MG</u>	<u>N016273 001</u>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-195 (of 436)

FUROSEMIDE

TABLET;ORAL

LASIX

AB +! 80MG

N016273 003

GABAPENTIN

CAPSULE;ORAL

GABAPENTIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A075350 001</u>	Sep 12, 2003
<u>AB</u>		<u>300MG</u>	<u>A075350 002</u>	Sep 12, 2003
<u>AB</u>		<u>400MG</u>	<u>A075350 003</u>	Sep 12, 2003
<u>AB</u>	ALKEM	<u>100MG</u>	<u>A090858 001</u>	Dec 17, 2010
<u>AB</u>		<u>300MG</u>	<u>A090858 002</u>	Dec 17, 2010
<u>AB</u>		<u>400MG</u>	<u>A090858 003</u>	Dec 17, 2010
<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>	<u>A078428 001</u>	Jul 25, 2007
<u>AB</u>		<u>300MG</u>	<u>A078428 002</u>	Jul 25, 2007
<u>AB</u>		<u>400MG</u>	<u>A078428 003</u>	Jul 25, 2007
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A075360 001</u>	Apr 06, 2005
<u>AB</u>		<u>300MG</u>	<u>A075360 002</u>	Apr 06, 2005
<u>AB</u>		<u>400MG</u>	<u>A075360 003</u>	Apr 06, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A078787 001</u>	Jan 31, 2008
<u>AB</u>		<u>300MG</u>	<u>A078787 002</u>	Jan 31, 2008
<u>AB</u>		<u>400MG</u>	<u>A078787 003</u>	Jan 31, 2008
<u>AB</u>	EPIC PHARMA LLC	<u>100MG</u>	<u>A207099 001</u>	Mar 24, 2017
<u>AB</u>		<u>300MG</u>	<u>A207099 002</u>	Mar 24, 2017
<u>AB</u>		<u>400MG</u>	<u>A207099 003</u>	Mar 24, 2017
<u>AB</u>	INVAGEN PHARMS	<u>100MG</u>	<u>A090705 001</u>	Dec 30, 2009
<u>AB</u>		<u>300MG</u>	<u>A090705 002</u>	Dec 30, 2009
<u>AB</u>		<u>400MG</u>	<u>A090705 003</u>	Dec 30, 2009
<u>AB</u>	JIANGSU HENGRI MED	<u>100MG</u>	<u>A091008 001</u>	Oct 26, 2017
<u>AB</u>		<u>300MG</u>	<u>A091008 002</u>	Oct 26, 2017
<u>AB</u>		<u>400MG</u>	<u>A091008 003</u>	Oct 26, 2017
<u>AB</u>	MARKSANS PHARMA	<u>100MG</u>	<u>A090007 001</u>	Jul 21, 2011
<u>AB</u>		<u>300MG</u>	<u>A090007 002</u>	Jul 21, 2011
<u>AB</u>		<u>400MG</u>	<u>A090007 003</u>	Jul 21, 2011
<u>AB</u>	MYLAN	<u>100MG</u>	<u>A090158 001</u>	Feb 14, 2011
<u>AB</u>		<u>300MG</u>	<u>A090158 002</u>	Feb 14, 2011
<u>AB</u>		<u>400MG</u>	<u>A090158 003</u>	Feb 14, 2011
<u>AB</u>	SCIEGEN PHARMS INC	<u>100MG</u>	<u>A204989 001</u>	Feb 18, 2016
<u>AB</u>		<u>300MG</u>	<u>A204989 002</u>	Feb 18, 2016
<u>AB</u>		<u>400MG</u>	<u>A204989 003</u>	Feb 18, 2016
<u>AB</u>	SUN PHARM INDs LTD	<u>100MG</u>	<u>A077242 001</u>	Aug 24, 2006
<u>AB</u>		<u>300MG</u>	<u>A077242 002</u>	Aug 24, 2006
<u>AB</u>		<u>400MG</u>	<u>A077242 003</u>	Aug 24, 2006
<u>AB</u>	TARO PHARM	<u>100MG</u>	<u>A077261 001</u>	Aug 02, 2013
<u>AB</u>		<u>300MG</u>	<u>A077261 002</u>	Aug 02, 2013
<u>AB</u>		<u>400MG</u>	<u>A077261 003</u>	Aug 02, 2013
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>A075435 001</u>	Oct 08, 2004
<u>AB</u>		<u>300MG</u>	<u>A075435 002</u>	Oct 08, 2004
<u>AB</u>		<u>400MG</u>	<u>A075435 003</u>	Oct 08, 2004
<u>NEURONTIN</u>				
<u>AB</u>	+ PFIZER PHARMS	<u>100MG</u>	<u>N020235 001</u>	Dec 30, 1993
<u>AB</u>	+	<u>300MG</u>	<u>N020235 002</u>	Dec 30, 1993
<u>AB</u>	+!	<u>400MG</u>	<u>N020235 003</u>	Dec 30, 1993

SOLUTION;ORAL

GABAPENTIN

<u>AA</u>	ACELLA PHARMS LLC	<u>250MG/5ML</u>	<u>A076403 001</u>	May 01, 2012
<u>AA</u>	AMNEAL PHARMS	<u>250MG/5ML</u>	<u>A202024 001</u>	Mar 23, 2012
<u>AA</u>	HI TECH PHARMA	<u>250MG/5ML</u>	<u>A078974 001</u>	Feb 18, 2011
<u>AA</u>	TARO	<u>250MG/5ML</u>	<u>A076672 001</u>	Jul 03, 2013
<u>AA</u>	TRIS PHARMA INC	<u>250MG/5ML</u>	<u>A091286 001</u>	Mar 14, 2016

NEURONTIN

<u>AA</u>	+! PARKE DAVIS	<u>250MG/5ML</u>	<u>N021129 001</u>	Mar 02, 2000
-----------	----------------	------------------	--------------------	--------------

TABLET;ORAL

GABAPENTIN

<u>AB</u>	ACI HEALTHCARE LTD	<u>600MG</u>	<u>A203244 002</u>	Jul 12, 2013
<u>AB</u>		<u>800MG</u>	<u>A203244 001</u>	Jul 12, 2013
<u>AB</u>	ACTAVIS ELIZABETH	<u>600MG</u>	<u>A075694 001</u>	Oct 21, 2004
<u>AB</u>		<u>800MG</u>	<u>A075694 002</u>	Oct 21, 2004
<u>AB</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A206402 001</u>	Dec 23, 2015
<u>AB</u>		<u>800MG</u>	<u>A206402 002</u>	Dec 23, 2015
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077894 001</u>	Oct 10, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-196 (of 436)

GABAPENTIN

TABLET;ORAL

GABAPENTIN

<u>AB</u>		<u>300MG</u>	<u>A077894</u> <u>002</u>	Oct 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077894</u> <u>003</u>	Oct 10, 2006
<u>AB</u>		<u>600MG</u>	<u>A077661</u> <u>004</u>	Sep 13, 2006
<u>AB</u>		<u>800MG</u>	<u>A077661</u> <u>005</u>	Sep 13, 2006
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A200651</u> <u>001</u>	Oct 06, 2011
<u>AB</u>		<u>800MG</u>	<u>A200651</u> <u>002</u>	Oct 06, 2011
<u>AB</u>	CSPC OUYI PHARM CO	<u>600MG</u>	<u>A207057</u> <u>001</u>	Oct 26, 2017
<u>AB</u>		<u>800MG</u>	<u>A207057</u> <u>002</u>	Oct 26, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>600MG</u>	<u>A077662</u> <u>001</u>	Aug 18, 2006
<u>AB</u>		<u>800MG</u>	<u>A077662</u> <u>002</u>	Aug 18, 2006
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A202764</u> <u>001</u>	Oct 16, 2012
<u>AB</u>		<u>800MG</u>	<u>A202764</u> <u>002</u>	Oct 16, 2012
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>100MG</u>	<u>A076017</u> <u>001</u>	Apr 28, 2004
<u>AB</u>		<u>300MG</u>	<u>A076017</u> <u>002</u>	Apr 28, 2004
<u>AB</u>		<u>400MG</u>	<u>A076017</u> <u>003</u>	Apr 28, 2004
<u>AB</u>		<u>600MG</u>	<u>A076017</u> <u>004</u>	Apr 29, 2005
<u>AB</u>		<u>800MG</u>	<u>A076017</u> <u>005</u>	Apr 29, 2005
<u>AB</u>	MYLAN PHARMS INC	<u>600MG</u>	<u>A090335</u> <u>001</u>	Jun 01, 2010
<u>AB</u>		<u>800MG</u>	<u>A090335</u> <u>002</u>	Jun 01, 2010
<u>AB</u>	SCIEGEN PHARMS INC	<u>600MG</u>	<u>A205101</u> <u>001</u>	Feb 04, 2016
<u>AB</u>		<u>800MG</u>	<u>A205101</u> <u>002</u>	Feb 04, 2016
<u>AB</u>	SUN PHARM INDs LTD	<u>600MG</u>	<u>A077525</u> <u>001</u>	Aug 24, 2006
<u>AB</u>		<u>800MG</u>	<u>A077525</u> <u>002</u>	Aug 24, 2006
<u>AB</u>	TEVA PHARMS USA	<u>600MG</u>	<u>A205807</u> <u>001</u>	Mar 10, 2017
<u>AB</u>		<u>800MG</u>	<u>A205807</u> <u>002</u>	Mar 10, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>600MG</u>	<u>A078926</u> <u>001</u>	Feb 11, 2011
<u>AB</u>		<u>800MG</u>	<u>A078926</u> <u>002</u>	Feb 11, 2011
<b>NEURONTIN</b>				
<u>AB</u>	+ PFIZER PHARMS	<u>600MG</u>	<u>N020882</u> <u>001</u>	Oct 09, 1998
<u>AB</u>	+!	<u>800MG</u>	<u>N020882</u> <u>002</u>	Oct 09, 1998
GRALISE				
BX	+! DEPOMED INC	300MG	N022544 001	Jan 28, 2011
BX	+!	600MG	N022544 002	Jan 28, 2011

GABAPENTIN ENACARBIL

TABLET, EXTENDED RELEASE;ORAL

HORIZANT

+ ARBOR PHARMS LLC	300MG	N022399 002	Dec 13, 2011
+!	600MG	N022399 001	Apr 06, 2011

GADOBENATE DIMEGLUMINE

INJECTABLE;INTRAVENOUS

MULTIHANCE

+! BRACCO	2.645GM/5ML (529MG/ML)	N021357 001	Nov 23, 2004
+!	5.29GM/10ML (529MG/ML)	N021357 002	Nov 23, 2004
+!	7.935GM/15ML (529MG/ML)	N021357 003	Nov 23, 2004
+!	10.58GM/20ML (529MG/ML)	N021357 004	Nov 23, 2004

MULTIHANCE MULTIPACK

+! BRACCO	26.45GM/50ML (529MG/ML)	N021358 001	Nov 23, 2004
+!	52.9GM/100ML (529MG/ML)	N021358 002	Nov 23, 2004

GADOBUTROL

SOLUTION;INTRAVENOUS

GADAVIST

+! BAYER HLTHCARE	1.20944GM/2ML (604.72MG/ML)	N201277 006	Dec 18, 2013
+!	4.5354GM/7.5ML (604.72MG/ML)	N201277 001	Mar 14, 2011
+!	6.0472GM/10ML (604.72MG/ML)	N201277 002	Mar 14, 2011
+!	9.0708GM/15ML (604.72MG/ML)	N201277 003	Mar 14, 2011
+!	18.1416GM/30ML (604.72MG/ML)	N201277 004	Mar 14, 2011
+!	39.3068GM/65ML (604.72MG/ML)	N201277 005	Mar 14, 2011

GADODIAMIDE

INJECTABLE;INJECTION

OMNISCAN

+! GE HEALTHCARE	287MG/ML	N020123 001	Jan 08, 1993
+!	28.7GM/100ML (287MG/ML)	N022066 002	Sep 05, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-197 (of 436)

GADOPENTETATE DIMEGLUMINE

INJECTABLE; INJECTION

MAGNEVIST

+!	BAYER HLTHCARE	469.01MG/ML
+!		469.01MG/ML

N019596	001	Jun 02, 1988
N021037	001	Mar 10, 2000

GADOTERATE MEGLUMINE

SOLUTION; INTRAVENOUS

DOTAREM

+!	GUERBET	37.69GM/100ML (376.9MG/ML)
+!		1.8845GM/5ML (376.9MG/ML)
+!		3.769GM/10ML (376.9MG/ML)
+!		5.6535GM/15ML (376.9MG/ML)
+!		7.538GM/20ML (376.9MG/ML)

N204781	001	Mar 20, 2013
N204781	005	Mar 31, 2017
N204781	002	Mar 20, 2013
N204781	003	Mar 20, 2013
N204781	004	Mar 20, 2013

GADOTERIDOL

INJECTABLE; INJECTION

PROHANCE

+!	BRACCO	279.3MG/ML
PROHANCE MULTIPACK		
+!	BRACCO	279.3MG/ML

N020131	001	Nov 16, 1992
N021489	001	Oct 09, 2003

GADOVERSETAMIDE

INJECTABLE; INJECTION

OPTIMARK

+!	LIEBEL-FLARSHEIM	1654.5MG/5ML (330.9MG/ML)
+!		3309MG/10ML (330.9MG/ML)
+!		4963.5MG/15ML (330.9MG/ML)
+!		6618MG/20ML (330.9MG/ML)
+!		16.545GM/50ML (330.9MG/ML)
OPTIMARK IN PLASTIC CONTAINER		
+!	LIEBEL-FLARSHEIM	3309MG/10ML (330.9MG/ML)
+!		4963.5MG/15ML (330.9MG/ML)
+!		6618MG/20ML (330.9MG/ML)
+!		9927MG/30ML (330.9MG/ML)

N020937	001	Dec 08, 1999
N020937	002	Dec 08, 1999
N020937	003	Dec 08, 1999
N020937	004	Dec 08, 1999
N020975	001	Dec 08, 1999
N020976	002	Dec 08, 1999
N020976	003	Dec 08, 1999
N020976	004	Dec 08, 1999
N020976	001	Dec 08, 1999

GADOXETATE DISODIUM

SOLUTION; INTRAVENOUS

EOVIST

+!	BAYER HLTHCARE	1.8143GM/10ML (181.43MG/ML)
+		2.72145GM/15ML (181.43MG/ML)

N022090	001	Jul 03, 2008
N022090	002	Feb 04, 2013

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 8MG BASE</u>	<u>A204895 001</u>	Aug 05, 2016
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A204895 002</u>	Aug 05, 2016
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A204895 003</u>	Aug 05, 2016
<u>AB</u>	BARR	<u>EQ 8MG BASE</u>	<u>A078189 001</u>	Sep 15, 2008
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A078189 002</u>	Sep 15, 2008
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A078189 003</u>	Sep 15, 2008
<u>AB</u>	MYLAN	<u>EQ 8MG BASE</u>	<u>A090900 001</u>	Jan 24, 2011
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A090900 002</u>	Jan 24, 2011
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A090900 003</u>	Jan 24, 2011
<u>AB</u>	SUN PHARMA GLOBAL	<u>EQ 8MG BASE</u>	<u>A090178 001</u>	Feb 02, 2011
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A090178 002</u>	Feb 02, 2011
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A090178 003</u>	Feb 02, 2011
<u>AB</u>	WATSON LABS	<u>EQ 8MG BASE</u>	<u>A079028 001</u>	Dec 15, 2008
<u>AB</u>		<u>EQ 16MG BASE</u>	<u>A079028 002</u>	Dec 15, 2008
<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A079028 003</u>	Dec 15, 2008

RAZADYNE ER

<u>AB</u>	+!	JANSSEN PHARMS	<u>EQ 8MG BASE</u>	<u>N021615 001</u>	Apr 01, 2005
<u>AB</u>	+		<u>EQ 16MG BASE</u>	<u>N021615 002</u>	Apr 01, 2005
<u>AB</u>	+		<u>EQ 24MG BASE</u>	<u>N021615 003</u>	Apr 01, 2005

SOLUTION; ORAL

GALANTAMINE HYDROBROMIDE

!	WEST-WARD PHARMS INT	4MG/ML
---	----------------------	--------

A078185	001	Jan 30, 2009
---------	-----	--------------

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

<u>AB</u>	APOTEK INC	<u>EQ 4MG BASE</u>	<u>A077781 001</u>	Sep 27, 2011
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>A077781 002</u>	Sep 27, 2011
<u>AB</u>		<u>EQ 12MG BASE</u>	<u>A077781 003</u>	Sep 27, 2011
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 4MG BASE</u>	<u>A090957 001</u>	Mar 29, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-198 (of 436)

GALANTAMINE HYDROBROMIDE

TABLET;ORAL

**GALANTAMINE HYDROBROMIDE**

LTD

<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A090957 002</b>	Mar 29, 2011
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A090957 003</b>	Mar 29, 2011
<b>AB</b>	BARR	<b>EQ 4MG BASE</b>	<b>A077605 001</b>	Aug 28, 2008
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077605 002</b>	Aug 28, 2008
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077605 003</b>	Aug 28, 2008
<b>AB</b>	DR REDDYS LABS LTD	<b>EQ 4MG BASE</b>	<b>A077593 001</b>	Sep 11, 2008
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077593 002</b>	Sep 11, 2008
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077593 003</b>	Sep 11, 2008
<b>AB</b>	MYLAN	<b>EQ 4MG BASE</b>	<b>A077590 001</b>	May 29, 2009
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077590 002</b>	May 29, 2009
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077590 003</b>	May 29, 2009
<b>AB</b>	SANDOZ	<b>EQ 4MG BASE</b>	<b>A077589 001</b>	Jun 22, 2009
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077589 002</b>	Jun 22, 2009
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077589 003</b>	Jun 22, 2009
<b>AB</b>	TEVA PHARMS	<b>EQ 4MG BASE</b>	<b>A077587 001</b>	Jul 09, 2009
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077587 002</b>	Jul 09, 2009
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077587 003</b>	Jul 09, 2009
<b>AB</b>	WEST-WARD PHARMS INT	<b>EQ 4MG BASE</b>	<b>A077608 001</b>	Feb 11, 2009
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077608 002</b>	Feb 11, 2009
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077608 003</b>	Feb 11, 2009
<b>AB</b>	YABAO PHARM	<b>EQ 4MG BASE</b>	<b>A077604 001</b>	Feb 06, 2009
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A077604 002</b>	Feb 06, 2009
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A077604 003</b>	Feb 06, 2009
<b>AB</b>	ZYDUS PHARMS USA INC	<b>EQ 4MG BASE</b>	<b>A078898 001</b>	Feb 17, 2011
<b>AB</b>		<b>EQ 8MG BASE</b>	<b>A078898 002</b>	Feb 17, 2011
<b>AB</b>		<b>EQ 12MG BASE</b>	<b>A078898 003</b>	Feb 17, 2011
<b>RAZADYNE</b>				
<b>AB</b>	+! JANSSEN PHARMS	<b>EQ 4MG BASE</b>	<b>N021169 001</b>	Feb 28, 2001
<b>AB</b>	+	<b>EQ 8MG BASE</b>	<b>N021169 002</b>	Feb 28, 2001
<b>AB</b>	+	<b>EQ 12MG BASE</b>	<b>N021169 003</b>	Feb 28, 2001

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

BS	LANTHEUS MEDCL	2mCi/ML
BS	MALLINKRODT NUCLEAR	2mCi/ML

N017478 001

N018058 001

GALLIUM DOTATATE GA-68

POWDER; INTRAVENOUS

NETSPOT

+! AAA USA INC

2.1-5.5mCi/ML

N208547 001 Jun 01, 2016

GANCICLOVIR

GEL; OPHTHALMIC

ZIRGAN

+! BAUSCH AND LOMB

0.15%

N022211 001 Sep 15, 2009

GANCICLOVIR

SOLUTION; IV (INFUSION)

GANCICLOVIR

+! EXELA PHARMA SCS LLC

500MG/250ML (2MG/ML)

N209347 001 Feb 17, 2017

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

**CYTOVENE**

<b>AP</b>	+! ROCHE PALO	<b>EQ 500MG BASE/VIAL</b>
<b>AP</b>		<b>EQ 500MG BASE/VIAL</b>

**N019661 001** Jun 23, 1989

**GANCICLOVIR**

<b>AP</b>	FRESENIUS KABI USA	<b>EQ 500MG BASE/VIAL</b>
<b>AP</b>	LUITPOLD PHARMS INC	<b>EQ 500MG BASE/VIAL</b>
<b>AP</b>	MYLAN LABS LTD	<b>EQ 500MG BASE/VIAL</b>
<b>AP</b>	PAR STERILE PRODUCTS	<b>EQ 500MG BASE/VIAL</b>

**A090658 001** Jun 21, 2010

**A202624 001** Sep 18, 2013

**A204560 001** Nov 17, 2017

**A204950 001** Dec 06, 2016

**GANCICLOVIR SODIUM**

<b>AP</b>	PHARMASCIENCE INC	<b>EQ 500MG BASE/VIAL</b>
-----------	-------------------	---------------------------

**A207645 001** Dec 08, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-199 (of 436)

GANIRELIX ACETATE

INJECTABLE; INJECTION

GANIRELIX ACETATE

+! ORGANON USA INC

EQ 250MCG BASE/0.5ML

N021057 001 Jul 29, 1999

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

**AT** HI-TECH PHARMA CO 0.5%

**A203189 001** Sep 03, 2014

**AT** LUPIN LTD 0.5%

**A202653 001** Aug 28, 2013

**AT** SANDOZ INC 0.5%

**A204227 001** Jul 11, 2016

ZYMAXID

**AT** +! ALLERGAN 0.5%

**N022548 001** May 18, 2010

ZYMAR

+! ALLERGAN

0.3%

N021493 001 Mar 28, 2003

GEFITINIB

TABLET; ORAL

IRESSA

+! ASTRAZENECA PHARMS 250MG

N206995 001 Jul 13, 2015

GEMCITABINE HYDROCHLORIDE

GEMCITABINE HYDROCHLORIDE

**AP** ACCORD HLTHCARE EQ 200MG BASE/VIAL

**A091594 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A091594 002** Jul 25, 2011

**AP** EQ 2GM BASE/VIAL

**A091594 003** Jul 25, 2011

**AP** ACTAVIS INC 200MG/5.26ML (38MG/ML)

**A204549 001** Apr 11, 2016

**AP** 1GM/26.3ML (38MG/ML)

**A204549 002** Apr 11, 2016

**AP** 2GM/52.6ML (38MG/ML)

**A204549 003** Apr 11, 2016

**AP** ACTAVIS TOTOWA EQ 200MG BASE/VIAL

**A079160 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A079160 002** Jul 25, 2011

**AP** EQ 2GM BASE/VIAL

**A079160 003** Jul 28, 2016

**AP** APOTEX INC 200MG/5.26ML (38MG/ML)

**A206776 001** May 23, 2017

**AP** 1GM/26.3ML (38MG/ML)

**A206776 002** May 23, 2017

**AP** 2GM/52.6ML (38MG/ML)

**A206776 003** May 23, 2017

**AP** CIPLA LTD EQ 200MG BASE/VIAL

**A078759 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A078759 002** Jul 25, 2011

**AP** DR REDDYS LABS LTD EQ 200MG BASE/VIAL

**A091365 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A091365 002** Jul 25, 2011

**AP** EQ 2GM BASE/VIAL

**A202997 001** May 07, 2013

**AP** EMCURE PHARMS LTD EQ 200MG BASE/VIAL

**A202063 001** Sep 11, 2012

**AP** EQ 1GM BASE/VIAL

**A202063 002** Sep 11, 2012

**AP** FRESENIUS KABI ONCOL EQ 200MG BASE/VIAL

**A090799 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A090799 002** Jul 25, 2011

**AP** EQ 2GM BASE/VIAL

**A090799 003** May 16, 2011

**AP** FRESENIUS KABI USA EQ 2GM BASE/VIAL

**A090242 003** May 16, 2011

**AP** GLAND PHARMA LTD EQ 200MG BASE/VIAL

**A204520 001** Jan 05, 2016

**AP** EQ 1GM BASE/VIAL

**A204520 002** Jan 05, 2016

**AP** HAMELN RDS GMBH EQ 200MG BASE/VIAL

**A090663 001** Sep 10, 2012

**AP** EQ 1GM BASE/VIAL

**A090663 002** Sep 10, 2012

**AP** HOSPIRA EQ 200MG BASE/VIAL

**A078339 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A078339 002** Jul 25, 2011

**AP** +! HOSPIRA INC 200MG/5.26ML (38MG/ML)

**N200795 001** Aug 04, 2011

**AP** +! 1GM/26.3ML (38MG/ML)

**N200795 002** Aug 04, 2011

**AP** ! EQ 2GM BASE/VIAL

**A079183 001** Nov 15, 2010

**AP** +! 2GM/52.6ML (38MG/ML)

**N200795 003** Aug 04, 2011

**AP** JIANGSU HANSOH PHARM EQ 200MG BASE/VIAL

**A202485 001** May 07, 2013

**AP** EQ 1GM BASE/VIAL

**A202485 002** May 07, 2013

**AP** LUITPOLD PHARMS INC EQ 200MG BASE/VIAL

**A202031 001** May 07, 2013

**AP** EQ 1GM BASE/VIAL

**A202031 002** May 07, 2013

**AP** MYLAN LABS LTD EQ 200MG BASE/VIAL

**A200145 001** Jul 25, 2011

**AP** 200MG/5.26ML (38MG/ML)

**A205242 001** Dec 06, 2017

**AP** EQ 1GM BASE/VIAL

**A200145 002** Jul 25, 2011

**AP** 1GM/26.3ML (38MG/ML)

**A205242 002** Dec 06, 2017

**AP** EQ 2GM BASE/VIAL

**A200145 003** Jul 25, 2011

**AP** 2GM/52.6ML (38MG/ML)

**A205242 003** Dec 06, 2017

**AP** SUN PHARMA GLOBAL EQ 200MG BASE/VIAL

**A078433 001** Jul 25, 2011

**AP** EQ 1GM BASE/VIAL

**A078433 002** Jul 25, 2011

**AP** TEVA PHARMS EQ 200MG BASE/VIAL

**A077983 002** Jan 25, 2011

**AP** EQ 1GM BASE/VIAL

**A077983 001** Jan 25, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-200 (of 436)

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMZAR

<u>AP</u>	+!	LILLY	<u>EQ 200MG BASE/VIAL</u>	<u>N020509 001</u>	May 15, 1996
<u>AP</u>	+!		<u>EQ 1GM BASE/VIAL</u>	<u>N020509 002</u>	May 15, 1996
SOLUTION;IV (INFUSION)					
GEMCITABINE HYDROCHLORIDE					
+!	ACCORD HLTHCARE		1GM/10ML (100MG/ML)	N209604 002	Aug 03, 2017
+!			1.5GM/15ML (100MG/ML)	N209604 003	Aug 03, 2017
+!			2GM/20ML (100MG/ML)	N209604 004	Aug 03, 2017
+!			200MG/2ML (100MG/ML)	N209604 001	Aug 03, 2017

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

<u>AB</u>	APOTEX	<u>600MG</u>	<u>A075034 001</u>	Jul 20, 1998
<u>AB</u>	AUROBINDO PHARMA LTD	<u>600MG</u>	<u>A202726 001</u>	Sep 16, 2015
<u>AB</u>	CADILA PHARMS LTD	<u>600MG</u>	<u>A203266 001</u>	Jun 17, 2016
<u>AB</u>	CARIBE HOLDINGS	<u>600MG</u>	<u>A078012 001</u>	Mar 26, 2007
<u>AB</u>	CHARTWELL MOLECULES	<u>600MG</u>	<u>A074270 001</u>	Sep 27, 1993
<u>AB</u>	HIKMA PHARMS	<u>600MG</u>	<u>A078599 001</u>	Aug 16, 2010
<u>AB</u>	IMPAX PHARMS	<u>600MG</u>	<u>A078207 001</u>	Jun 01, 2007
<u>AB</u>	INVAGEN PHARMS	<u>600MG</u>	<u>A077836 001</u>	Jul 27, 2006
<u>AB</u>	NORTHSTAR HLTHCARE	<u>600MG</u>	<u>A079072 001</u>	Sep 13, 2010
<u>AB</u>	SUN PHARM INDs INC	<u>600MG</u>	<u>A079239 001</u>	Dec 29, 2008
<u>AB</u>	TEVA	<u>600MG</u>	<u>A074256 001</u>	Oct 31, 1993
	<u>LOPID</u>			
<u>AB</u>	+! PFIZER PHARMS	<u>600MG</u>	<u>N018422 003</u>	Nov 20, 1986

GEMIFLOXACIN MESYLATE

TABLET; ORAL

FACTIVE

<u>AB</u>	+! LG CHEM LTD	<u>EQ 320MG BASE</u>	<u>N021158 001</u>	Apr 04, 2003
<u>AB</u>	GEMIFLOXACIN MESYLATE	<u>EQ 320MG BASE</u>	<u>A090466 001</u>	Jun 15, 2015

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>	G AND W LABS INC	<u>EQ 0.1% BASE</u>	<u>A064056 001</u>	Apr 29, 1994
<u>AT</u>	! PERRIGO NEW YORK	<u>EQ 0.1% BASE</u>	<u>A062307 001</u>	

INJECTABLE; INJECTION

GENTAMICIN SULFATE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 10MG BASE/ML</u>	<u>A062366 002</u>	Feb 06, 1986
<u>AP</u>	!	<u>EQ 40MG BASE/ML</u>	<u>A062366 001</u>	Aug 04, 1983
<u>AP</u>	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A062420 001</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>A062612 004</u>	Feb 20, 1986
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>A062420 002</u>	Aug 15, 1983

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 1.2MG BASE/ML</u>	<u>A062373 007</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>A062373 008</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>A062373 002</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>A062373 005</u>	Sep 07, 1982
<u>AP</u>	HOSPIRA	<u>EQ 1.2MG BASE/ML</u>	<u>A062414 001</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>A062414 003</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>A062414 008</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>A062414 010</u>	Aug 15, 1983
	! BAXTER HLTHCARE	EQ 2MG BASE/ML	A062373 009	Sep 07, 1982
		EQ 120MG BASE/100ML	A062373 006	Sep 07, 1982

OINTMENT; OPHTHALMIC

GENTAMICIN SULFATE

<u>AT</u>	! AKORN	<u>EQ 0.3% BASE</u>	<u>A064093 001</u>	Aug 31, 1995
<u>AT</u>	PERRIGO CO TENNESSEE	<u>EQ 0.3% BASE</u>	<u>A065024 001</u>	Jul 30, 2004

OINTMENT; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>	FOUGERA PHARMS INC	<u>EQ 0.1% BASE</u>	<u>A062533 001</u>	Oct 05, 1984
<u>AT</u>	! PERRIGO NEW YORK	<u>EQ 0.1% BASE</u>	<u>A062351 001</u>	Feb 18, 1982
<u>AT</u>	TARO	<u>EQ 0.1% BASE</u>	<u>A062477 001</u>	Dec 23, 1983

SOLUTION/DROPS; OPHTHALMIC

GENOPTIC

<u>AT</u>	! ALLERGAN	<u>EQ 0.3% BASE</u>	<u>A062452 001</u>	Oct 10, 1984
-----------	------------	---------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-201 (of 436)

GENTAMICIN SULFATE

SOLUTION/DROPS;OPHTHALMIC

GENTAK

<u>AT</u>	AKORN	<u>EQ 0.3% BASE</u>	<u>A064163 001</u>	Oct 12, 2001
<u><b>GENTAMICIN SULFATE</b></u>				
<u>AT</u>	AKORN	<u>EQ 0.3% BASE</u>	<u>A062635 001</u>	Jan 08, 1987
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.3% BASE</u>	<u>A064048 001</u>	May 11, 1994
<u>AT</u>	PERRIGO CO TENNESSEE	<u>EQ 0.3% BASE</u>	<u>A065121 001</u>	Jan 30, 2004
<u>AT</u>	SANDOZ INC	<u>EQ 0.3% BASE</u>	<u>A062196 001</u>	

GENTAMICIN SULFATE; PREDNISOLONE ACETATE

OINTMENT;OPHTHALMIC

PRED-G

+!	ALLERGAN	EQ 0.3% BASE;0.6%	N050612 001	Dec 01, 1989
SUSPENSION/DROPS;OPHTHALMIC				
PRED-G	+!	ALLERGAN	EQ 0.3% BASE;1%	N050586 001 Jun 10, 1988

GLATIRAMER ACETATE

INJECTABLE;SUBCUTANEOUS

COPAXONE

<u>AP</u>	+!	TEVA PHARMS USA	<u>20MG/ML</u>	<u>N020622 002</u>	Feb 12, 2002
<u>AP</u>	+!		<u>40MG/ML</u>	<u>N020622 003</u>	Jan 28, 2014
<u><b>GLATIRAMER ACETATE</b></u>					
<u>AP</u>		MYLAN PHARMS INC	<u>20MG/ML</u>	<u>A091646 001</u>	Oct 03, 2017
<u>AP</u>			<u>40MG/ML</u>	<u>A206936 001</u>	Oct 03, 2017

GLATOPA

<u>AP</u>	SANDOZ INC	<u>20MG/ML</u>	<u>A090218 001</u>	Apr 16, 2015
-----------	------------	----------------	--------------------	--------------

GLECAPREVIR; PIBRENTASVIR

TABLET;ORAL

MAVYRET

+!	ABBVIE INC	100MG;40MG	N209394 001	Aug 03, 2017
----	------------	------------	-------------	--------------

GLIMEPIRIDE

TABLET;ORAL

AMARYL

<u>AB</u>	+!	SANOFI AVENTIS US	<u>1MG</u>	<u>N020496 001</u>	Nov 30, 1995
<u>AB</u>	+		<u>2MG</u>	<u>N020496 002</u>	Nov 30, 1995
<u>AB</u>	+		<u>4MG</u>	<u>N020496 003</u>	Nov 30, 1995

GLIMEPIRIDE

<u>AB</u>	ACCORD HLTHCARE	<u>1MG</u>	<u>A078181 001</u>	Aug 23, 2007
<u>AB</u>		<u>2MG</u>	<u>A078181 002</u>	Aug 23, 2007
<u>AB</u>		<u>4MG</u>	<u>A078181 003</u>	Aug 23, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1MG</u>	<u>A202759 001</u>	Jun 29, 2012
<u><b>GLIMEPIRIDE</b></u>				
<u>AB</u>		<u>2MG</u>	<u>A202759 002</u>	Jun 29, 2012
<u>AB</u>		<u>4MG</u>	<u>A202759 003</u>	Jun 29, 2012
<u>AB</u>	CARLSBAD	<u>1MG</u>	<u>A077911 001</u>	Sep 22, 2009
<u>AB</u>		<u>2MG</u>	<u>A077911 002</u>	Sep 22, 2009
<u>AB</u>		<u>4MG</u>	<u>A077911 003</u>	Sep 22, 2009
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>A077091 001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A077091 002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A077091 003</u>	Oct 06, 2005
<u>AB</u>	INDOCO REMEDIES	<u>1MG</u>	<u>A202112 001</u>	Apr 17, 2013
<u>AB</u>		<u>2MG</u>	<u>A202112 002</u>	Apr 17, 2013
<u>AB</u>		<u>4MG</u>	<u>A202112 003</u>	Apr 17, 2013
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>A077295 001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A077295 002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A077295 003</u>	Oct 06, 2005
<u>AB</u>	MYLAN	<u>1MG</u>	<u>A077624 001</u>	Nov 28, 2005
<u>AB</u>		<u>2MG</u>	<u>A077624 002</u>	Nov 28, 2005
<u>AB</u>		<u>4MG</u>	<u>A077624 003</u>	Nov 28, 2005
<u>AB</u>	PRINSTON INC	<u>1MG</u>	<u>A077370 001</u>	Dec 23, 2005
<u>AB</u>		<u>2MG</u>	<u>A077370 002</u>	Dec 23, 2005
<u>AB</u>		<u>4MG</u>	<u>A077370 003</u>	Dec 23, 2005
<u>AB</u>		<u>8MG</u>	<u>A077370 004</u>	Dec 23, 2005
<u>AB</u>	TEVA	<u>1MG</u>	<u>A076802 001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>A076802 002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>A076802 003</u>	Oct 06, 2005
<u>AB</u>	VIVA HLTHCARE	<u>1MG</u>	<u>A091220 001</u>	Jun 29, 2012
<u>AB</u>		<u>2MG</u>	<u>A091220 002</u>	Jun 29, 2012
<u>AB</u>		<u>4MG</u>	<u>A091220 004</u>	Jun 29, 2012
<u>AB</u>		<u>8MG</u>	<u>A091220 006</u>	Jun 29, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-202 (of 436)

GLIMEPIRIDE

TABLET;ORAL  
 GLIMEPIRIDE

3MG	A091220 003 Jun 29, 2012
6MG	A091220 005 Jun 29, 2012

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET;ORAL  
DUETACT

<u>AB</u>	+!	TAKEDA PHARMS USA	<u>2MG;30MG</u>	<u>N021925 001</u> Jul 28, 2006
<u>AB</u>	+		<u>4MG;30MG</u>	<u>N021925 002</u> Jul 28, 2006
<u>PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE</u>				
<u>AB</u>		SANDOZ	<u>2MG;30MG</u>	<u>A201049 001</u> Jan 04, 2013
<u>AB</u>			<u>4MG;30MG</u>	<u>A201049 002</u> Jan 04, 2013

GLIMEPIRIDE; ROSIGLITAZONE MALEATE

TABLET;ORAL

ROSIGLITAZONE MALEATE AND GLIMEPIRIDE  
 ! TEVA PHARMS USA 1MG;4MG  
                           2MG;4MG  
                           2MG;8MG  
                           4MG;4MG  
                           4MG;8MG

A078709 001 Apr 01, 2016
A078709 002 Apr 01, 2016
A078709 004 Apr 01, 2016
A078709 003 Apr 01, 2016
A078709 005 Apr 01, 2016

GLIPIZIDE

TABLET;ORAL

GLIPIZIDE

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A074550 001</u> Sep 11, 1997
<u>AB</u>			<u>10MG</u>	<u>A074550 002</u> Sep 11, 1997
<u>AB</u>		ANI PHARMS INC	<u>5MG</u>	<u>A074497 001</u> Aug 31, 1995
<u>AB</u>			<u>10MG</u>	<u>A074497 002</u> Aug 31, 1995
<u>AB</u>		APOTEX	<u>5MG</u>	<u>A075795 001</u> Jun 13, 2001
<u>AB</u>			<u>10MG</u>	<u>A075795 002</u> Jun 13, 2001
<u>AB</u>		MYLAN	<u>5MG</u>	<u>A074226 001</u> May 10, 1994
<u>AB</u>			<u>10MG</u>	<u>A074226 002</u> May 10, 1994
<u>AB</u>		SANDOZ	<u>5MG</u>	<u>A074305 001</u> Apr 07, 1995
<u>AB</u>			<u>10MG</u>	<u>A074305 002</u> Apr 07, 1995
<u>AB</u>		SUN PHARM INDS INC	<u>5MG</u>	<u>A077820 001</u> Jul 11, 2006
<u>AB</u>			<u>10MG</u>	<u>A077820 002</u> Jul 11, 2006
<u>AB</u>		WATSON LABS TEVA	<u>5MG</u>	<u>A074223 001</u> Feb 27, 1995
<u>AB</u>			<u>10MG</u>	<u>A074223 002</u> Feb 27, 1995

GLUCOTROL

<u>AB</u>	+!	PFIZER	<u>5MG</u>	<u>N017783 001</u> May 08, 1984
<u>AB</u>	+		<u>10MG</u>	<u>N017783 002</u> May 08, 1984

TABLET, EXTENDED RELEASE;ORAL

GLIPIZIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>2 . 5MG</u>	<u>A206928 001</u> May 12, 2017
<u>AB</u>			<u>5MG</u>	<u>A206928 002</u> May 12, 2017
<u>AB</u>			<u>10MG</u>	<u>A206928 003</u> May 12, 2017
<u>AB</u>		MYLAN PHARMS INC	<u>2 . 5MG</u>	<u>A202298 001</u> May 19, 2015
<u>AB</u>			<u>5MG</u>	<u>A202298 002</u> May 19, 2015
<u>AB</u>			<u>10MG</u>	<u>A202298 003</u> May 19, 2015
<u>AB</u>		PAR PHARM	<u>5MG</u>	<u>A076159 002</u> Sep 20, 2013
<u>AB</u>			<u>10MG</u>	<u>A076159 001</u> Sep 20, 2013
<u>AB</u>		UNIQUE PHARM LABS	<u>2 . 5MG</u>	<u>A204720 001</u> Dec 29, 2016
<u>AB</u>			<u>5MG</u>	<u>A204720 002</u> Dec 29, 2016
<u>AB</u>			<u>10MG</u>	<u>A204720 003</u> Dec 29, 2016
<u>AB</u>		WATSON LABS	<u>2 . 5MG</u>	<u>A076467 003</u> Mar 27, 2006
<u>AB</u>			<u>5MG</u>	<u>A076467 001</u> Sep 08, 2003
<u>AB</u>			<u>10MG</u>	<u>A076467 002</u> Nov 07, 2003

GLUCOTROL XL

<u>AB</u>	+	PFIZER	<u>2 . 5MG</u>	<u>N020329 003</u> Aug 10, 1999
<u>AB</u>	+		<u>5MG</u>	<u>N020329 001</u> Apr 26, 1994
<u>AB</u>	!+		<u>10MG</u>	<u>N020329 002</u> Apr 26, 1994

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET;ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>		EPIC PHARMA LLC	<u>2 . 5MG;250MG</u>	<u>A077507 001</u> Oct 27, 2005
<u>AB</u>			<u>2 . 5MG;500MG</u>	<u>A077507 002</u> Oct 27, 2005
<u>AB</u>			<u>5MG;500MG</u>	<u>A077507 003</u> Oct 27, 2005
<u>AB</u>		HERITAGE PHARMS INC	<u>2 . 5MG;250MG</u>	<u>A078728 001</u> Jun 23, 2010
<u>AB</u>			<u>2 . 5MG;500MG</u>	<u>A078728 002</u> Jun 23, 2010
<u>AB</u>			<u>5MG;500MG</u>	<u>A078728 003</u> Jun 23, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-203 (of 436)

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>2.5MG;250MG</u>	<u>A078083 001</u>	Apr 12, 2007
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078083 002</u>	Apr 12, 2007
<u>AB</u>		<u>5MG;500MG</u>	<u>A078083 003</u>	Apr 12, 2007
<u>AB</u>	SUN PHARM INDNS INC	<u>2.5MG;250MG</u>	<u>A077620 001</u>	Jan 11, 2008
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077620 002</u>	Jan 11, 2008
<u>AB</u>		<u>5MG;500MG</u>	<u>A077620 003</u>	Jan 11, 2008
<u>AB</u>	TEVA PHARMS	<u>2.5MG;250MG</u>	<u>A077270 001</u>	Oct 28, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A077270 002</u>	Oct 28, 2005
<u>AB</u>	!	<u>5MG;500MG</u>	<u>A077270 003</u>	Oct 28, 2005
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2.5MG;250MG</u>	<u>A078905 001</u>	Jan 31, 2011
<u>AB</u>		<u>2.5MG;500MG</u>	<u>A078905 002</u>	Jan 31, 2011
<u>AB</u>		<u>5MG;500MG</u>	<u>A078905 003</u>	Jan 31, 2011

GLUCAGON HYDROCHLORIDE

POWDER; INTRAMUSCULAR, INTRAVENOUS  
 GLUCAGON

+! FRESENIUS KABI USA EQ 1MG BASE/VIAL

N201849 001 May 08, 2015

GLUCAGON HYDROCHLORIDE RECOMBINANT

INJECTABLE; INJECTION  
 GLUCAGEN

+! NOVO NORDISK EQ 1MG BASE/VIAL

N020918 001 Jun 22, 1998

GLUCAGON RECOMBINANT

INJECTABLE; INJECTION  
 GLUCAGON

+! LILLY 1MG/VIAL

N020928 001 Sep 11, 1998

GLYBURIDE

TABLET; ORAL

GLYBURIDE (MICRONIZED)

<u>AB</u>	DAVA PHARMS INC	<u>1.5MG</u>	<u>A074591 001</u>	Dec 22, 1997
<u>AB</u>		<u>3MG</u>	<u>A074591 002</u>	Dec 22, 1997
<u>AB</u>		<u>4.5MG</u>	<u>A074591 003</u>	Dec 22, 1997
<u>AB</u>		<u>6MG</u>	<u>A074591 004</u>	Dec 22, 1997
<u>AB</u>	HIKMA	<u>1.5MG</u>	<u>A075890 001</u>	Jul 31, 2003
<u>AB</u>		<u>3MG</u>	<u>A075890 002</u>	Jul 31, 2003
<u>AB</u>		<u>6MG</u>	<u>A075890 003</u>	Jul 31, 2003
<u>AB</u>	MYLAN	<u>1.5MG</u>	<u>A074792 001</u>	Jun 26, 1998
<u>AB</u>		<u>3MG</u>	<u>A074792 002</u>	Jun 26, 1998
<u>AB</u>		<u>6MG</u>	<u>A074792 003</u>	Aug 17, 1999
<u>AB</u>	TEVA	<u>1.5MG</u>	<u>A074686 001</u>	Apr 20, 1999
<u>AB</u>		<u>3MG</u>	<u>A074686 002</u>	Apr 20, 1999
<u>AB</u>		<u>4.5MG</u>	<u>A074686 003</u>	Apr 20, 1999
<u>AB</u>		<u>6MG</u>	<u>A074686 004</u>	Apr 20, 1999

GLYNASE

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>1.5MG</u>	<u>N020051 001</u>	Mar 04, 1992
<u>AB</u>	+	<u>3MG</u>	<u>N020051 002</u>	Mar 04, 1992
<u>AB</u>	+	<u>6MG</u>	<u>N020051 004</u>	Sep 24, 1993

GLYBURIDE

<u>AB1</u>	AUROBINDO PHARMA	<u>1.25MG</u>	<u>A077537 001</u>	Oct 18, 2007
<u>AB1</u>		<u>2.5MG</u>	<u>A077537 002</u>	Oct 18, 2007
<u>AB1</u>		<u>5MG</u>	<u>A077537 003</u>	Oct 18, 2007
<u>AB1</u>	EPIC PHARMA LLC	<u>1.25MG</u>	<u>A076257 001</u>	Jun 27, 2002
<u>AB1</u>		<u>2.5MG</u>	<u>A076257 002</u>	Jun 27, 2002
<u>AB1</u>		<u>5MG</u>	<u>A076257 003</u>	Jun 27, 2002
<u>AB1</u>	HERITAGE PHARMS INC	<u>1.25MG</u>	<u>A090937 001</u>	Feb 28, 2011
<u>AB1</u>		<u>2.5MG</u>	<u>A090937 002</u>	Feb 28, 2011
<u>AB1</u>		<u>5MG</u>	<u>A090937 003</u>	Feb 28, 2011
<u>AB1</u>	PHARMADAX INC	<u>1.25MG</u>	<u>A203581 001</u>	Apr 14, 2016
<u>AB1</u>		<u>2.5MG</u>	<u>A203581 002</u>	Apr 14, 2016
<u>AB1</u>		<u>5MG</u>	<u>A203581 003</u>	Apr 14, 2016
<u>AB1</u>	TEVA	<u>1.25MG</u>	<u>A074388 001</u>	Aug 29, 1995
<u>AB1</u>		<u>2.5MG</u>	<u>A074388 002</u>	Aug 29, 1995
<u>AB1</u>		<u>5MG</u>	<u>A074388 003</u>	Aug 29, 1995
<u>AB1</u>	ZYDUS PHARMS USA INC	<u>1.25mg</u>	<u>A206749 001</u>	May 10, 2016
<u>AB1</u>		<u>2.5mg</u>	<u>A206749 002</u>	May 10, 2016
<u>AB1</u>		<u>5MG</u>	<u>A206749 003</u>	May 10, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-204 (of 436)

GLYBURIDE

TABLET;ORAL

DIABETA

<u>AB2</u>	+	SANOFI AVENTIS US	<u>1.25MG</u>
<u>AB2</u>	+		<u>2.5MG</u>
<u>AB2</u>	!+		<u>5MG</u>

GLYBURIDE

<u>AB2</u>		IMPAX LABS INC	<u>1.25MG</u>
<u>AB2</u>			<u>2.5MG</u>
<u>AB2</u>			<u>5MG</u>

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET;ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>1.25MG;250MG</u>
<u>AB</u>			<u>2.5MG;500MG</u>
<u>AB</u>			<u>5MG;500MG</u>
<u>AB</u>		AUROBINDO PHARMA	<u>1.25MG;250MG</u>
<u>AB</u>	!		<u>2.5MG;500MG</u>
<u>AB</u>			<u>5MG;500MG</u>
<u>AB</u>		HERITAGE PHARMS INC	<u>1.25MG;250MG</u>
<u>AB</u>			<u>2.5MG;500MG</u>
<u>AB</u>			<u>5MG;500MG</u>
<u>AB</u>		IMPAX LABS INC	<u>1.25MG;250MG</u>
<u>AB</u>			<u>2.5MG;500MG</u>
<u>AB</u>			<u>5MG;500MG</u>
<u>AB</u>		ZYDUS PHARMS USA INC	<u>1.25MG;250MG</u>
<u>AB</u>			<u>2.5MG;500MG</u>
<u>AB</u>			<u>5MG;500MG</u>

<u>N017532</u>	<u>001</u>	May 01, 1984
<u>N017532</u>	<u>002</u>	May 01, 1984
<u>N017532</u>	<u>003</u>	May 01, 1984

GLYCEROL PHENYLBUTYRATE

LIQUID;ORAL

RAVICTI

+! HORIZON THERAPS INC 1.1GM/ML

N203284 001 Feb 01, 2013

GLYCINE

SOLUTION;IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

<u>AT</u>		BAXTER HLTHCARE	<u>1.5GM/100ML</u>
<u>AT</u>		GLYCINE 1.5% IN PLASTIC CONTAINER	

N017865 001

GLCOPYRROLATE

INJECTABLE;INJECTION

GLCOPYRROLATE

<u>AP</u>		AMNEAL PHARMS CO	<u>0.2MG/ML</u>
<u>AP</u>		FRESENIUS KABI USA	<u>0.2MG/ML</u>
<u>AP</u>	!	HIKMA FARMACEUTICA	<u>0.2MG/ML</u>
<u>AP</u>		LUITPOLD	<u>0.2MG/ML</u>
<u>AP</u>		SOMERSET THERAPS LLC	<u>0.2MG/ML</u>

N016784 001

N018315 001

POWDER;INHALATION

SEEBRI

+! SUNOVION PHARMS INC 15.6MCG/INH

N207923 001 Oct 29, 2015

SOLUTION;INHALATION

LONHALA MAGNAIR KIT

+! SUNOVION RESP 25MCG/ML

N208437 001 Dec 05, 2017

SOLUTION;ORAL

CUVPOSA

+! MERZ PHARMS 1MG/5ML

N022571 001 Jul 28, 2010

TABLET;ORAL

GLCOPYRROLATE

<u>AA</u>		APPCO PHARMA LLC	<u>1MG</u>
<u>AA</u>			<u>2MG</u>
<u>AA</u>		AUROLIFE PHARMA LLC	<u>1MG</u>
<u>AA</u>		DR REDDYS LABS LTD	<u>1MG</u>
<u>AA</u>			<u>2MG</u>
<u>AA</u>		LEADING PHARMA LLC	<u>1MG</u>
<u>AA</u>			<u>2MG</u>
<u>AA</u>		NATCO PHARMA LTD	<u>1MG</u>
<u>AA</u>			<u>2MG</u>
<u>AA</u>		PAR PHARM	<u>1MG</u>
<u>AA</u>			<u>2MG</u>

A207201 001 Jan 03, 2017

A207201 002 Jan 03, 2017

A202675 001 Apr 15, 2013

A040847 001 Mar 21, 2008

A040847 002 Mar 21, 2008

A090195 001 Sep 21, 2012

A090195 002 Sep 21, 2012

A091413 001 Jun 20, 2016

A091413 002 Jun 20, 2016

A040653 001 Aug 31, 2006

A040653 002 Aug 31, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-205 (of 436)

GLYCOPYRROLATE

TABLET;ORAL

GLYCOPYRROLATE

<u>AA</u>	RISING PHARMS INC	<u>1MG</u>	<u>A040821 001</u>	Dec 29, 2008
<u>AA</u>		<u>2MG</u>	<u>A040821 002</u>	Dec 29, 2008
<u>AA</u>	SANTOS BIOTECH	<u>1MG</u>	<u>A091182 001</u>	Feb 03, 2014
<u>AA</u>		<u>2MG</u>	<u>A091182 002</u>	Feb 03, 2014
<u>AA</u>	SUN PHARM INDS LTD	<u>1MG</u>	<u>A040844 001</u>	Aug 18, 2009
<u>AA</u>		<u>2MG</u>	<u>A040844 002</u>	Aug 18, 2009
<u>AA</u>	VINTAGE PHARMS	<u>1MG</u>	<u>A090020 001</u>	Oct 19, 2011
<u>AA</u>		<u>2MG</u>	<u>A090020 002</u>	Oct 19, 2011
		<u><b>ROBINUL</b></u>		
<u>AA</u>	+! CASPER PHARMA LLC	<u>1MG</u>	<u>N012827 001</u>	
		<u><b>ROBINUL FORTE</b></u>		
<u>AA</u>	+! CASPER PHARMA LLC	<u>2MG</u>	<u>N012827 002</u>	
		<u>GLYCOPYRROLATE</u>		
	NEXGEN PHARMA	1.5MG		A091522 001 Mar 12, 2012

GLYCOPYRROLATE ; INDACATEROL MALEATE

POWDER;INHALATION  
 UTIBRON

+! SUNOVION PHARMS INC 15.6MCG/INH;27.5MCG/INH

N207930 001 Oct 29, 2015

GONADOTROPIN, CHORIONIC

INJECTABLE;INJECTION

CHORIONIC GONADOTROPIN

<u>AP</u>	+! FERRING	<u>10,000 UNITS/VIAL</u>	<u>N017016 007</u>
<u>AP</u>	+! FRESENIUS KABI USA	<u>10,000 UNITS/VIAL</u>	<u>N017067 002</u>
	<u>PREGNYL</u>		
<u>AP</u>	+! ORGANON USA INC	<u>10,000 UNITS/VIAL</u>	<u>N017692 001</u>
	CHORIONIC GONADOTROPIN		
	+! FERRING	5,000 UNITS/VIAL	N017016 006

GOSERELIN ACETATE

IMPLANT;IMPLANTATION  
 ZOLADEX

+! TERSERA THERAPS LLC EQ 3.6MG BASE  
 +! EQ 10.8MG BASE

N019726 001 Dec 29, 1989  
 N020578 001 Jan 11, 1996

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

<u>AT</u>	AMRING PHARMS	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>A065187 001</u>	Oct 28, 2005
<u>AT</u>	! BAUSCH AND LOMB	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>A064047 001</u>	Jan 31, 1996
	<u>NEOSPORIN</u>			
<u>AT</u>	! MONARCH PHARMS	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>A060582 001</u>	

GRANISETRON

FILM, EXTENDED RELEASE;TRANSDERMAL

SANCUSO

+! KYOWA KIRIN 3.1MG/24HR

N022198 001 Sep 12, 2008

INJECTION, EXTENDED RELEASE;SUBCUTANEOUS

SUSTOL

+! HERON THERAPS INC 10MG/0.4ML (10MG/0.4ML)

N022445 001 Aug 09, 2016

GRANISETRON HYDROCHLORIDE

INJECTABLE;INJECTION

GRANISETRON HYDROCHLORIDE

<u>AP</u>	AKORN INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A079119 001</u>	Sep 10, 2009
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A079078 001</u>	Sep 14, 2009
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A079078 002</u>	Sep 14, 2009
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A204238 001</u>	Jul 06, 2016
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A204238 002</u>	Jul 06, 2016
<u>AP</u>	BIONPHARMA INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078863 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078880 001</u>	Jun 30, 2008
<u>AP</u>	CIPLA LTD	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078262 001</u>	Dec 31, 2007
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078258 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078258 002</u>	Jun 30, 2008
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078522 001</u>	Dec 31, 2007
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078090 001</u>	Jun 30, 2008
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078629 001</u>	Dec 23, 2009
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078629 002</u>	Dec 23, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-206 (of 436)

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

GRANISETRON HYDROCHLORIDE

<u>AP</u>	LUITPOLD	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A091274 001</u>	Sep 22, 2010
<u>AP</u>	MYLAN ASI	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A091137 002</u>	Apr 09, 2010
<u>AP</u>	MYLAN LABS LTD	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A203454 001</u>	Apr 04, 2017
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A203454 002</u>	Apr 04, 2017
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A203453 001</u>	Jan 31, 2017
<u>AP</u>	SAGENT STRIDES	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A091136 001</u>	Apr 09, 2010
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A091136 002</u>	Apr 09, 2010
<u>AP</u>	SANDOZ INC	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078534 001</u>	Apr 30, 2009
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078531 001</u>	Apr 30, 2009
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078835 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078531 002</u>	Apr 30, 2009
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078835 002</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078392 001</u>	Dec 31, 2007
<u>AP</u>	TEVA PHARMS USA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077963 001</u>	Jan 03, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077297 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A077913 001</u>	Jun 26, 2008
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077186 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A077187 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A077177 001</u>	Dec 31, 2007
<u>AP</u>	WOCKHARDT USA	<u>EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)</u>	<u>A078566 001</u>	Feb 29, 2008
<u>AP</u>		<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078564 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A078565 001</u>	Jun 30, 2008
<u>GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE</u>				
<u>AP</u>	BIONPHARMA INC	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078863 002</u>	Jun 30, 2008
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML (EQ 1MG BASE/ML)</u>	<u>A078096 001</u>	Jun 30, 2008

TABLET; ORAL

GRANISETRON HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>EQ 1MG BASE</u>	<u>A078843 001</u>	Feb 27, 2008
<u>AB</u>	CIPLA LTD	<u>EQ 1MG BASE</u>	<u>A078037 001</u>	Feb 27, 2008
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 1MG BASE</u>	<u>A078846 001</u>	Feb 27, 2009
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>A078725 001</u>	Jan 30, 2008
<u>AB</u>	NATCO PHARMA	<u>EQ 1MG BASE</u>	<u>A078969 001</u>	Jun 22, 2009
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 1MG BASE</u>	<u>A078678 001</u>	Feb 13, 2008
<u>AB</u>	TARO PHARM	<u>EQ 1MG BASE</u>	<u>A090817 001</u>	May 28, 2010
<u>AB</u>	TEVA PHARMS	<u>EQ 1MG BASE</u>	<u>A078080 001</u>	Dec 31, 2007
<u>AB</u>	WEST-WARD PHARMS INT	<u>EQ 1MG BASE</u>	<u>A077842 001</u>	Dec 31, 2007

GRISEOFULVIN, MICROSIZE

SUSPENSION; ORAL

GRISEOFULVIN

<u>AB</u>	ACTAVIS MID ATLANTIC	<u>125MG/5ML</u>	<u>A065394 001</u>	Jul 06, 2007
<u>AB</u>	CHARTWELL RX	<u>125MG/5ML</u>	<u>A065200 001</u>	Mar 02, 2005
<u>AB</u>	CIPLA LTD	<u>125MG/5ML</u>	<u>A065354 001</u>	Sep 10, 2007
<u>AB</u>	VINTAGE PHARMS	<u>125MG/5ML</u>	<u>A065438 001</u>	Oct 08, 2010

TABLET; ORAL

GRISEOFULVIN

<u>AB</u>	SANDOZ INC	<u>500MG</u>	<u>A091592 002</u>	Aug 07, 2013
<u>AB</u>	SIGMAPHARM LABS LLC	<u>500MG</u>	<u>A202482 001</u>	Oct 22, 2012
	SANDOZ INC	<u>250MG</u>	A091592 001	Aug 07, 2013

GRISEOFULVIN, ULTRAMICROSIZE

TABLET; ORAL

GRIS-PEG

<u>AB</u>	VALEANT PHARMS INC	<u>125MG</u>	<u>N050475 001</u>	
<u>AB</u>	+!	<u>250MG</u>	<u>N050475 002</u>	

GRISEOFULVIN, ULTRAMICROSIZE

<u>AB</u>	COREPHARMA	<u>125MG</u>	<u>A204371 001</u>	Jan 09, 2014
<u>AB</u>		<u>250MG</u>	<u>A204371 002</u>	Jan 09, 2014

GRISEOFULVIN, ULTRAMICROSIZE

<u>AB</u>	SIGMAPHARM LABS LLC	<u>125MG</u>	<u>A202545 001</u>	Oct 22, 2012
<u>AB</u>		<u>250MG</u>	<u>A202545 002</u>	Oct 22, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-207 (of 436)

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION;ORAL

FLOWTUSS

MISSION PHARMACAL      200MG/5ML; 2.5MG/5ML  
 CO

N022424 001 May 14, 2015

GUAIFENESIN; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYCOFENIX

+! MISSION PHARMACAL      200MG/5ML; 2.5MG/5ML; 30MG/5ML  
 CO

N022279 001 May 14, 2015

GUANABENZ ACETATE

TABLET;ORAL

GUANABENZ ACETATE

ANI PHARMS INC      EQ 4MG BASE  
 !                        EQ 8MG BASE

A074149 001 Apr 07, 1995

A074149 002 Apr 07, 1995

GUANFACINE HYDROCHLORIDE

TABLET;ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARM	<u>EQ 1MG BASE</u>	<u>A075109 001</u>	Nov 25, 1998
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075109 002</u>	Nov 25, 1998
<u>AB</u>	EPIC PHARMA LLC	<u>EQ 1MG BASE</u>	<u>A074673 001</u>	Feb 28, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074673 002</u>	Feb 28, 1997
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>A074796 001</u>	Jan 27, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074796 002</u>	Jan 27, 1997
<u>AB</u>	WATSON LABS	<u>EQ 1MG BASE</u>	<u>A074145 001</u>	Oct 17, 1995
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A074145 002</u>	Oct 17, 1995

TENEX

<u>AB</u> +	PROMIUS PHARMA	<u>EQ 1MG BASE</u>	<u>N019032 001</u>	Oct 27, 1986
<u>AB</u> +!		<u>EQ 2MG BASE</u>	<u>N019032 002</u>	Nov 07, 1988

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 1MG BASE</u>	<u>A200881 001</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A200881 002</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A200881 003</u>	Oct 05, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A200881 004</u>	Oct 05, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 1MG BASE</u>	<u>A202578 001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A202578 002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A202578 003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202578 004</u>	Jun 02, 2015
<u>AB</u>	SANDOZ INC	<u>EQ 1MG BASE</u>	<u>A202568 001</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A202568 002</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A202568 003</u>	Jun 03, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A202568 004</u>	Jun 03, 2015
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 1MG BASE</u>	<u>A205689 001</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A205689 002</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A205689 003</u>	Nov 16, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A205689 004</u>	Nov 16, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 1MG BASE</u>	<u>A201382 001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A201382 002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A201382 003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201382 004</u>	Jun 02, 2015
<u>AB</u>	TWI PHARMS INC	<u>EQ 1MG BASE</u>	<u>A201408 001</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A201408 002</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A201408 003</u>	Jun 02, 2015
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A201408 004</u>	Jun 02, 2015

INTUNIV

<u>AB</u> +	SHIRE	<u>EQ 1MG BASE</u>	<u>N022037 001</u>	Sep 02, 2009
<u>AB</u> +		<u>EQ 2MG BASE</u>	<u>N022037 002</u>	Sep 02, 2009
<u>AB</u> +		<u>EQ 3MG BASE</u>	<u>N022037 003</u>	Sep 02, 2009
<u>AB</u> +!		<u>EQ 4MG BASE</u>	<u>N022037 004</u>	Sep 02, 2009

GUANIDINE HYDROCHLORIDE

TABLET;ORAL

GUANIDINE HYDROCHLORIDE

MERCK SHARP DOHME      125MG

N001546 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-208 (of 436)

HALCINONIDE

CREAM;TOPICAL  
 HALOG  
 +! RANBAXY 0.1% N017556 001  
 OINTMENT;TOPICAL  
 HALOG  
 +! SUN PHARM INDNS INC 0.1% N017824 001

HALOBETASOL PROPIONATE

CREAM;TOPICAL  
HALOBETASOL PROPIONATE  
AB FOUGERA PHARMS 0.05% A077001 001 Dec 16, 2004  
AB ! G AND W LABS 0.05% A078162 001 Apr 24, 2007  
AB PERRIGO ISRAEL 0.05% A077123 001 Dec 16, 2004  
AB TARO 0.05% A077227 001 Aug 04, 2005  
ULTRAVATE  
AB + SUN PHARM INDNS LTD 0.05% N019967 001 Dec 27, 1990  
 LOTION;TOPICAL  
 ULTRAVATE  
 +! SUN PHARM INDUSTRIES 0.05% N208183 001 Nov 06, 2015  
 OINTMENT;TOPICAL  
HALOBETASOL PROPIONATE  
AB ! G AND W LABS 0.05% A077721 001 Sep 07, 2006  
AB G AND W LABS INC 0.05% A077109 001 Jun 14, 2005  
AB PERRIGO ISRAEL 0.05% A076872 001 Dec 16, 2004  
AB TARO 0.05% A076994 001 Dec 16, 2004  
ULTRAVATE  
AB + RANBAXY 0.05% N019968 001 Dec 17, 1990

HALOPERIDOL

TABLET;ORAL  
HALOPERIDOL  
AB MYLAN 0.5MG A070278 006 Jun 10, 1986  
AB 1MG A070278 004 Jun 10, 1986  
AB 2MG A070278 001 Jun 10, 1986  
AB 5MG A070278 005 Jun 10, 1986  
AB 10MG A070278 002 Jul 16, 2009  
AB 20MG A070278 003 Jul 16, 2009  
AB SANDOZ 0.5MG A071206 001 Nov 17, 1986  
AB 1MG A071207 001 Nov 17, 1986  
AB ! 2MG A071208 001 Nov 17, 1986  
AB 5MG A071209 001 Nov 17, 1986  
AB 10MG A071210 001 Mar 11, 1988  
AB 20MG A071211 001 Mar 11, 1988  
AB ZYDUS PHARMS USA 5MG A077580 003 Nov 29, 2007  
AB 10MG A077580 004 Nov 29, 2007  
AB 20MG A077580 005 Nov 29, 2007

HALOPERIDOL DECANOATE

INJECTABLE;INJECTION  
HALDOL  
AO +! JANSSEN PHARMS EQ 50MG BASE/ML N018701 001 Jan 14, 1986  
AO +! EQ 100MG BASE/ML N018701 002 Jan 31, 1997  
HALOPERIDOL DECANOATE  
AO FRESENIUS KABI USA EQ 50MG BASE/ML A074893 001 Dec 19, 1997  
AO EQ 100MG BASE/ML A074893 002 Dec 19, 1997  
AO GLAND PHARMA LTD EQ 50MG BASE/ML A205241 001 May 12, 2017  
AO EQ 100MG BASE/ML A205241 002 May 12, 2017  
AO MYLAN LABS LTD EQ 50MG BASE/ML A075440 001 Feb 28, 2000  
AO EQ 100MG BASE/ML A075440 002 Feb 28, 2000  
AO TEVA PHARMS USA EQ 50MG BASE/ML A075393 001 May 11, 1999  
AO EQ 100MG BASE/ML A075393 002 May 11, 1999  
AO WEST-WARD PHARMS INT EQ 50MG BASE/ML A074811 001 Jan 30, 1998  
AO EQ 100MG BASE/ML A075305 001 Sep 28, 1998

HALOPERIDOL LACTATE

CONCENTRATE;ORAL  
HALOPERIDOL  
AA PHARM ASSOC EQ 2MG BASE/ML A073037 001 Feb 26, 1993  
AA SILARX EQ 2MG BASE/ML A073364 001 Sep 28, 1993  
AA ! TEVA PHARMS EQ 2MG BASE/ML A071617 001 Dec 01, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-209 (of 436)

HALOPERIDOL LACTATE

INJECTABLE; INJECTION

HALDOL

<u>AP</u>	+!	JANSSEN PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N015923 001</u>
<u>AP</u>			<u>HALOPERIDOL</u>	
<u>AP</u>		AKORN	<u>EQ 5MG BASE/ML</u>	<u>A204849 001</u> Sep 06, 2017
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A075689 001</u> Mar 09, 2001
<u>AP</u>		GLAND PHARMA LTD	<u>EQ 5MG BASE/ML</u>	<u>A076774 001</u> Aug 25, 2004
<u>AP</u>		MYLAN LABS LTD	<u>EQ 5MG BASE/ML</u>	<u>A078347 001</u> Sep 14, 2009
<u>AP</u>		SAGENT PHARMS	<u>EQ 5MG BASE/ML</u>	<u>A091637 001</u> Sep 02, 2011
<u>AP</u>			<u>EQ 5MG BASE/ML</u>	<u>A200742 001</u> Sep 02, 2011
<u>AP</u>		TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A076035 001</u> Aug 29, 2001
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 5MG BASE/ML</u>	<u>A075858 001</u> Jun 18, 2001

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

<u>AP</u>	+!	FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 001</u>
			<u>5,000 UNITS/ML</u>	<u>A206552 001</u> Jun 10, 2016
<u>AP</u>	+!		<u>5,000 UNITS/ML</u>	<u>N017651 006</u>
<u>AP</u>	+!		<u>10,000 UNITS/ML</u>	<u>N017029 003</u>
<u>AP</u>	+!		<u>20,000 UNITS/ML</u>	<u>N017029 004</u>
<u>AP</u>		GLAND PHARMA LTD	<u>5,000 UNITS/ML</u>	<u>A205323 001</u> Feb 06, 2017
<u>AP</u>		HOSPIRA INC	<u>1,000 UNITS/ML</u>	<u>A090571 001</u> Aug 31, 2009
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A090571 002</u> Aug 31, 2009
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A090571 003</u> Aug 31, 2009
<u>AP</u>		MYLAN LABS LTD	<u>1,000 UNITS/ML</u>	<u>A203851 001</u> Nov 30, 2017
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A203851 002</u> Nov 30, 2017
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A203851 003</u> Nov 30, 2017
<u>AP</u>			<u>20,000 UNITS/ML</u>	<u>A203852 001</u> Nov 30, 2017
<u>AP</u>		SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<u>A090808 001</u> Jun 30, 2010
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A090808 002</u> Jun 30, 2010
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A090808 003</u> Jun 30, 2010
<u>AP</u>			<u>20,000 UNITS/ML</u>	<u>A090809 001</u> Jun 30, 2010
<u>AP</u>		SANDOZ	<u>1,000 UNITS/ML</u>	<u>A091682 001</u> Jun 08, 2011
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A091659 001</u> Jun 08, 2011
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A091682 002</u> Jun 08, 2011
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A201002 001</u> Jun 08, 2011
<u>AP</u>		SHENZHEN TECHDOW	<u>1,000 UNITS/ML</u>	<u>A202957 001</u> Jun 12, 2014
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A202733 001</u> Jun 12, 2014
<u>AP</u>			<u>5,000 UNITS/ML</u>	<u>A202957 002</u> Jun 12, 2014
<u>AP</u>			<u>10,000 UNITS/ML</u>	<u>A203198 001</u> Jun 12, 2014
<u>AP</u>			<u>20,000 UNITS/ML</u>	<u>A203198 002</u> Jun 12, 2014
<u>AP</u>	+!	WEST-WARD PHARMS INT	<u>1,000 UNITS/ML</u>	<u>N017037 001</u>
<u>AP</u>	+!		<u>5,000 UNITS/ML</u>	<u>N017037 002</u>
<u>AP</u>	+!		<u>10,000 UNITS/ML</u>	<u>N017037 003</u>
			<u>HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>	
<u>AP</u>		BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 001</u> Apr 28, 1982
			<u>HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>	
<u>AP</u>	+!	B BRAUN	<u>200 UNITS/100ML</u>	<u>N019953 001</u> Jul 20, 1992
<u>AP</u>		HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916 010</u> Jun 23, 1989
			<u>HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	
<u>AP</u>		HOSPIRA	<u>10,000 UNITS/100ML</u>	<u>N019339 003</u> Mar 27, 1985
			<u>HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>	
<u>AP</u>		BAXTER HLTHCARE	<u>200 UNITS/100ML</u>	<u>N018609 002</u> Apr 28, 1982
			<u>HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>	
<u>AP</u>		HOSPIRA	<u>200 UNITS/100ML</u>	<u>N018916 011</u> Jun 23, 1989
			<u>HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	
<u>AP</u>	+!	B BRAUN	<u>4,000 UNITS/100ML</u>	<u>N019952 001</u> Jul 20, 1992
<u>AP</u>		HOSPIRA	<u>4,000 UNITS/100ML</u>	<u>N019805 001</u> Jan 25, 1989
			<u>HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER</u>	
<u>AP</u>	+!	B BRAUN	<u>5,000 UNITS/100ML</u>	<u>N019952 004</u> Jul 20, 1992
<u>AP</u>	+!		<u>10,000 UNITS/100ML</u>	<u>N019952 005</u> Jul 20, 1992
<u>AP</u>		HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N019339 004</u> Mar 27, 1985
<u>AP</u>			<u>5,000 UNITS/100ML</u>	<u>N019805 002</u> Jan 25, 1989
<u>AP</u>			<u>10,000 UNITS/100ML</u>	<u>N019339 002</u> Mar 27, 1985
			<u>HEPARIN SODIUM IN PLASTIC CONTAINER</u>	
<u>AP</u>	+!	FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<u>N017029 013</u> Dec 05, 1985
<u>AP</u>	+!		<u>5,000 UNITS/ML</u>	<u>N017029 014</u> Dec 05, 1985
<u>AP</u>	+!		<u>10,000 UNITS/ML</u>	<u>N017029 015</u> Dec 05, 1985
<u>AP</u>	+!		<u>20,000 UNITS/ML</u>	<u>N017029 016</u> Dec 05, 1985

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-210 (of 436)

HEPARIN SODIUM

INJECTABLE; INJECTION

**HEPARIN SODIUM PRESERVATIVE FREE**

<b>AP</b>	+!	FRESENIUS KABI USA	<u>1,000 UNITS/ML</u>	<b>N017029 010</b>	Apr 28, 1986
<b>AP</b>		SAGENT PHARMS	<u>1,000 UNITS/ML</u>	<b>A090810 001</b>	Jun 30, 2010
<b>AP</b>		SHENZHEN TECHDOW	<u>1,000 UNITS/ML</u>	<b>A202732 001</b>	Jun 12, 2014
HEPARIN SODIUM					
	+!	FRESENIUS KABI USA	10,000 UNITS/ML	N017029 020	Mar 31, 2011
	!	HOSPIRA	5,000 UNITS/ML	A088100 001	Apr 28, 1983
	+!	PFIZER	1,000 UNITS/ML	N201370 001	Jul 21, 2011
	+!		5,000 UNITS/ML	N201370 002	Jul 21, 2011
	+!		10,000 UNITS/ML	N201370 003	Jul 21, 2011
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER					
		HOSPIRA	5,000 UNITS/100ML	N019339 001	Mar 27, 1985
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER					
		HOSPIRA	5,000 UNITS/100ML	N018916 006	Jan 31, 1984
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER					
		HOSPIRA	5,000 UNITS/100ML	N018916 007	Jan 31, 1984
			10,000 UNITS/100ML	N018916 008	Jan 31, 1984
HEPARIN SODIUM PRESERVATIVE FREE					
	+!	FRESENIUS KABI USA	10,000 UNITS/ML	N017029 019	Nov 22, 2010
	!	HOSPIRA	10,000 UNITS/ML	A089522 001	May 04, 1987
	+!	PFIZER	1,000 UNITS/ML	N201370 004	Jul 21, 2011

HEXACHLOROPHENE

SPONGE; TOPICAL

**PRE-OP**

<b>AT</b>	+!	DAVIS AND GECK	<u>480MG</u>	<b>N017433 001</b>
<b>AT</b>	+	DAVIS AND GECK	<u>480MG</u>	<b>N017433 002</b>

HEXAMINOLEVULINATE HYDROCHLORIDE

FOR SOLUTION; INTRAVESICAL

CYSVIEW KIT

+!	PHOTOCURE ASA	100MG/VIAL
----	---------------	------------

N022555 001 May 28, 2010

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

SUPPRELIN LA

+!	ENDO PHARM	50MG	N022058 001	May 03, 2007
+!	ENDO PHARM	50MG	N021732 001	Oct 12, 2004

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

**HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE**

<b>AA</b>		ABHAI LLC	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A207487 001</b>	Feb 21, 2017
<b>AA</b>		ACTAVIS MID ATLANTIC	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A088017 001</b>	Jul 05, 1983
<b>AA</b>		BIO-PHARM INC	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A204765 001</b>	Mar 06, 2017
<b>AA</b>	!	HI TECH PHARMA	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A040613 001</b>	Feb 08, 2008
<b>AA</b>		NOVEL LABS INC	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A203535 001</b>	Feb 13, 2017
<b>AA</b>		PADDOCK LLC	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A205731 001</b>	Feb 15, 2017
<b>AA</b>		WOCKHARDT BIO AG	<u>1.5MG/5ML ; 5MG/5ML</u>	<b>A088008 001</b>	Mar 03, 1983

TABLET; ORAL

**HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE**

<b>AA</b>		AVANTHI INC	<u>1.5MG ; 5MG</u>	<b>A207176 001</b>	Aug 07, 2017
<b>AA</b>		NOVEL LABS INC	<u>1.5MG ; 5MG</u>	<b>A091528 001</b>	Apr 20, 2011

**TUSSIGON**

<b>AA</b>	!	KING PHARMS	<u>1.5MG ; 5MG</u>	<b>A088508 001</b>	Jul 30, 1985
-----------	---	-------------	--------------------	--------------------	--------------

HYALURONIDASE

INJECTABLE; INJECTION

AMPHADASE

+!	AMPHASTAR PHARM	150 UNITS/ML	N021665 001	Oct 26, 2004
+!	BAUSCH AND LOMB	200 UNITS/VIAL	N021640 002	Dec 02, 2004

HYALURONIDASE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HYLENEX RECOMBINANT

+!	HALOZYME THERAP	150 UNITS/ML	N021859 001	Dec 02, 2005
----	-----------------	--------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-211 (of 436)

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

<u>AP</u>	!	AKORN	<u>20MG/ML</u>	<u>A040730</u> <u>001</u>	Apr 21, 2009
<u>AP</u>		FRESENIUS KABI USA	<u>20MG/ML</u>	<u>A040388</u> <u>001</u>	Mar 13, 2001
<u>AP</u>		LUITPOLD	<u>20MG/ML</u>	<u>A040136</u> <u>001</u>	Jun 30, 1997
<u>AP</u>		MYLAN INSTITUTIONAL	<u>20MG/ML</u>	<u>A204680</u> <u>001</u>	Apr 28, 2016
<u>AP</u>		NAVINTA LLC	<u>20MG/ML</u>	<u>A202938</u> <u>001</u>	Mar 28, 2013
<u>AP</u>		X-GEN PHARMS INC	<u>20MG/ML</u>	<u>A203110</u> <u>001</u>	Jun 29, 2015

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

<u>AA</u>		ALKEM LABS LTD	<u>10MG</u>	<u>A200737</u> <u>001</u>	Dec 07, 2012
<u>AA</u>			<u>25MG</u>	<u>A200737</u> <u>002</u>	Dec 07, 2012
<u>AA</u>			<u>50MG</u>	<u>A200737</u> <u>003</u>	Dec 07, 2012
<u>AA</u>			<u>100MG</u>	<u>A200737</u> <u>004</u>	Dec 07, 2012
<u>AA</u>		CADILA PHARMS LTD	<u>25MG</u>	<u>A203845</u> <u>001</u>	Sep 18, 2014
<u>AA</u>			<u>50MG</u>	<u>A203845</u> <u>002</u>	Sep 18, 2014
<u>AA</u>			<u>100MG</u>	<u>A203845</u> <u>003</u>	Sep 18, 2014
<u>AA</u>		GLENMARK PHARMS LTD	<u>10MG</u>	<u>A090527</u> <u>001</u>	May 27, 2009
<u>AA</u>			<u>25MG</u>	<u>A090527</u> <u>002</u>	May 27, 2009
<u>AA</u>			<u>50MG</u>	<u>A090527</u> <u>003</u>	May 27, 2009
<u>AA</u>			<u>100MG</u>	<u>A090527</u> <u>004</u>	May 27, 2009
<u>AA</u>		HERITAGE PHARMS INC	<u>10MG</u>	<u>A086242</u> <u>001</u>	Feb 04, 2010
<u>AA</u>			<u>25MG</u>	<u>A086242</u> <u>003</u>	
<u>AA</u>			<u>50MG</u>	<u>A086242</u> <u>002</u>	
<u>AA</u>			<u>100MG</u>	<u>A086242</u> <u>004</u>	Feb 04, 2010
<u>AA</u>		HETERO LABS LTD III	<u>10MG</u>	<u>A040901</u> <u>001</u>	Sep 12, 2008
<u>AA</u>			<u>25MG</u>	<u>A040901</u> <u>002</u>	Sep 12, 2008
<u>AA</u>			<u>50MG</u>	<u>A040901</u> <u>003</u>	Sep 12, 2008
<u>AA</u>			<u>100MG</u>	<u>A040901</u> <u>004</u>	Sep 12, 2008
<u>AA</u>		INVAGEN PHARMS	<u>10MG</u>	<u>A090255</u> <u>001</u>	Dec 15, 2008
<u>AA</u>			<u>25MG</u>	<u>A090255</u> <u>002</u>	Dec 15, 2008
<u>AA</u>			<u>50MG</u>	<u>A090255</u> <u>003</u>	Dec 15, 2008
<u>AA</u>			<u>100MG</u>	<u>A090255</u> <u>004</u>	Dec 15, 2008
<u>AA</u>		PAR PHARM	<u>10MG</u>	<u>A087836</u> <u>001</u>	Oct 05, 1982
<u>AA</u>			<u>25MG</u>	<u>A086961</u> <u>002</u>	
<u>AA</u>			<u>50MG</u>	<u>A086962</u> <u>001</u>	
<u>AA</u>			<u>100MG</u>	<u>A088391</u> <u>001</u>	Sep 27, 1983
<u>AA</u>	!	PLIVA	<u>10MG</u>	<u>A089097</u> <u>001</u>	Dec 18, 1985
<u>AA</u>	!		<u>25MG</u>	<u>A088467</u> <u>001</u>	May 01, 1984
<u>AA</u>	!		<u>50MG</u>	<u>A088468</u> <u>001</u>	May 01, 1984
<u>AA</u>	!		<u>100MG</u>	<u>A089098</u> <u>001</u>	Dec 18, 1985
<u>AA</u>		STRIDES PHARMA	<u>25MG</u>	<u>A200770</u> <u>001</u>	May 03, 2013
<u>AA</u>			<u>50MG</u>	<u>A200770</u> <u>002</u>	May 03, 2013
<u>AA</u>			<u>100MG</u>	<u>A200770</u> <u>003</u>	May 03, 2013
<u>AA</u>		TECH ORGANIZED	<u>10MG</u>	<u>A205236</u> <u>001</u>	May 26, 2017
<u>AA</u>			<u>25MG</u>	<u>A205236</u> <u>002</u>	May 26, 2017
<u>AA</u>			<u>50MG</u>	<u>A205236</u> <u>003</u>	May 26, 2017
<u>AA</u>			<u>100MG</u>	<u>A205236</u> <u>004</u>	May 26, 2017

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRA-ZIDE

PAR PHARM	<u>25MG; 25MG</u>	<u>A088957</u> <u>001</u>	Oct 21, 1985
!	<u>50MG; 50MG</u>	<u>A088946</u> <u>001</u>	Oct 21, 1985

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

+!	ARBOR PHARMS LLC	<u>37.5MG; 20MG</u>	<u>N020727</u> <u>001</u>	Jun 23, 2005
----	------------------	---------------------	---------------------------	--------------

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>		ALEMBIC PHARMS LTD	<u>12.5MG</u>	<u>A200645</u> <u>001</u>	Nov 30, 2010
<u>AB</u>		APOTEX	<u>12.5MG</u>	<u>A078389</u> <u>001</u>	May 16, 2008
<u>AB</u>		AUROBINDO PHARMA	<u>12.5MG</u>	<u>A078164</u> <u>001</u>	Sep 18, 2007
<u>AB</u>		IPCA LABS LTD	<u>12.5MG</u>	<u>A079237</u> <u>001</u>	Apr 02, 2009
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>12.5MG</u>	<u>A077005</u> <u>001</u>	Jul 13, 2005
<u>AB</u>		JUBILANT CADISTA	<u>12.5MG</u>	<u>A078391</u> <u>001</u>	Feb 11, 2008
<u>AB</u>		MYLAN	<u>12.5MG</u>	<u>A075640</u> <u>001</u>	Jan 28, 2000
<u>AB</u>		PRINSTON INC	<u>12.5MG</u>	<u>A075907</u> <u>001</u>	Sep 17, 2002
<u>AB</u>		SUN PHARM INDNS INC	<u>12.5MG</u>	<u>A090651</u> <u>001</u>	Apr 07, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-212 (of 436)

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>	UNICHEM	<u>12.5MG</u>	<u>A090510 001</u>	Jan 19, 2010
		<u>MICROZIDE</u>		
<u>AB</u>	+! ALLERGAN SALES LLC	<u>12.5MG</u>	<u>N020504 001</u>	Dec 27, 1996
	TABLET; ORAL			
		<u>HYDROCHLOROTHIAZIDE</u>		
<u>AB</u>	ACCORD HLTHCARE	<u>12.5MG</u>	<u>A202556 001</u>	Sep 24, 2012
<u>AB</u>		<u>25MG</u>	<u>A202556 002</u>	Sep 24, 2012
<u>AB</u>		<u>50MG</u>	<u>A202556 003</u>	Sep 24, 2012
<u>AB</u>	ACTAVIS ELIZABETH	<u>12.5MG</u>	<u>A040707 001</u>	Feb 27, 2007
<u>AB</u>	APOTEX	<u>25MG</u>	<u>A040774 001</u>	Oct 03, 2007
<u>AB</u>		<u>50MG</u>	<u>A040774 002</u>	Oct 03, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A040780 001</u>	Jul 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040780 002</u>	Jul 20, 2007
<u>AB</u>	DAVA PHARMS INC	<u>25MG</u>	<u>A087059 001</u>	
<u>AB</u>		<u>50MG</u>	<u>A087068 001</u>	
<u>AB</u>	HERITAGE PHARMS INC	<u>25MG</u>	<u>A085182 002</u>	
<u>AB</u>		<u>50MG</u>	<u>A085182 001</u>	
<u>AB</u>	HIKMA INTL PHARMS	<u>50MG</u>	<u>A084878 001</u>	
<u>AB</u>	IPCA LABS LTD	<u>12.5MG</u>	<u>A040807 001</u>	Jul 20, 2007
<u>AB</u>		<u>25MG</u>	<u>A040807 002</u>	Jul 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040807 003</u>	Jul 20, 2007
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A083177 001</u>	
<u>AB</u>		<u>50MG</u>	<u>A083177 002</u>	
<u>AB</u>	LEADING PHARMA LLC	<u>12.5MG</u>	<u>A040702 003</u>	May 10, 2017
<u>AB</u>		<u>25MG</u>	<u>A040702 001</u>	Mar 16, 2007
<u>AB</u>		<u>50MG</u>	<u>A040702 002</u>	Mar 16, 2007
<u>AB</u>	MYLAN PHARMS INC	<u>25MG</u>	<u>A040735 002</u>	Jan 23, 2007
<u>AB</u>		<u>50MG</u>	<u>A040735 003</u>	Jan 23, 2007
<u>AB</u>	PRINSTON INC	<u>25MG</u>	<u>A040412 001</u>	Mar 29, 2002
<u>AB</u>		<u>50MG</u>	<u>A040412 002</u>	Mar 29, 2002
<u>AB</u>	SUN PHARM INDs INC	<u>12.5MG</u>	<u>A040857 001</u>	May 30, 2008
<u>AB</u>		<u>25MG</u>	<u>A040810 001</u>	Mar 27, 2007
<u>AB</u>		<u>50MG</u>	<u>A040810 002</u>	Mar 27, 2007
<u>AB</u>	TECH ORGANIZED	<u>25MG</u>	<u>A203018 001</u>	Jul 23, 2014
<u>AB</u>		<u>50MG</u>	<u>A203018 002</u>	Jul 23, 2014
<u>AB</u>	UNICHEM	<u>25MG</u>	<u>A040907 001</u>	Aug 15, 2008
<u>AB</u>		<u>50MG</u>	<u>A040907 002</u>	Aug 15, 2008

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

<u>AB</u>	+! SANOFI AVENTIS US	<u>12.5MG;150MG</u>	<u>N020758 002</u>	Sep 30, 1997
<u>AB</u>	+!	<u>12.5MG;300MG</u>	<u>N020758 003</u>	Aug 31, 1998
		<u>IRBESARTAN AND HYDROCHLOROTHIAZIDE</u>		
<u>AB</u>	ALEMBIC PHARMS LTD	<u>12.5MG;150MG</u>	<u>A091370 001</u>	Oct 15, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A091370 002</u>	Oct 15, 2012
<u>AB</u>		<u>25MG;300MG</u>	<u>A091370 003</u>	Oct 12, 2016
<u>AB</u>	APOTEX INC	<u>12.5MG;150MG</u>	<u>A201505 001</u>	Oct 15, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A201505 002</u>	Oct 15, 2012
<u>AB</u>	ATLAS PHARMS LLC	<u>12.5MG;150MG</u>	<u>A203036 001</u>	Jan 15, 2016
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A203036 002</u>	Jan 15, 2016
<u>AB</u>		<u>25MG;300MG</u>	<u>A203036 003</u>	Jan 15, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>12.5MG;150MG</u>	<u>A203630 001</u>	Feb 22, 2013
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A203630 002</u>	Feb 22, 2013
<u>AB</u>		<u>25MG;300MG</u>	<u>A203630 003</u>	Mar 31, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>12.5MG;150MG</u>	<u>A203500 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A203500 002</u>	Sep 27, 2012
<u>AB</u>	HISUN PHARM HANGZHOU	<u>12.5MG;150MG</u>	<u>A207896 001</u>	Oct 14, 2016
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A207896 002</u>	Oct 14, 2016
<u>AB</u>	LUPIN LTD	<u>12.5MG;150MG</u>	<u>A201524 001</u>	Feb 27, 2013
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A201524 002</u>	Feb 27, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG;150MG</u>	<u>A202414 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A202414 002</u>	Sep 27, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>12.5MG;150MG</u>	<u>A077969 001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077969 002</u>	Sep 27, 2012
<u>AB</u>		<u>25MG;300MG</u>	<u>A077969 003</u>	Jul 20, 2016
<u>AB</u>	PRINSTON INC	<u>12.5MG;150MG</u>	<u>A203072 001</u>	May 09, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-213 (of 436)

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

IRBESARTAN AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>12.5MG;300MG</u>	<u>A203072</u> <u>002</u>	May 09, 2014
<u>AB</u>	SANDOZ	<u>12.5MG;150MG</u>	<u>A077446</u> <u>001</u>	Sep 27, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077446</u> <u>002</u>	Sep 27, 2012
<u>AB</u>	TEVA	<u>12.5MG;150MG</u>	<u>A077369</u> <u>001</u>	Mar 30, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A077369</u> <u>002</u>	Mar 30, 2012
<u>AB</u>	UNICHEM LABS LTD	<u>12.5MG;150MG</u>	<u>A207018</u> <u>001</u>	Sep 19, 2017
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A207018</u> <u>002</u>	Sep 19, 2017
<u>AB</u>	WEST-WARD PHARMS INT	<u>12.5MG;150MG</u>	<u>A090351</u> <u>001</u>	Oct 15, 2012
<u>AB</u>		<u>12.5MG;300MG</u>	<u>A090351</u> <u>002</u>	Oct 15, 2012
<u>AB</u>		<u>25MG;300MG</u>	<u>A090351</u> <u>003</u>	Jun 08, 2017

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET; ORAL

LISINOPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX INC	<u>12.5MG;10MG</u>	<u>A076674</u> <u>001</u>	Oct 05, 2004
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076674</u> <u>002</u>	Oct 05, 2004
<u>AB</u>		<u>25MG;20MG</u>	<u>A076674</u> <u>003</u>	Oct 05, 2004
<u>AB</u>	AUROBINDO	<u>12.5MG;10MG</u>	<u>A077606</u> <u>001</u>	Mar 14, 2006
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A077606</u> <u>002</u>	Mar 14, 2006
<u>AB</u>		<u>25MG;20MG</u>	<u>A077606</u> <u>003</u>	Mar 14, 2006
<u>AB</u>	HIKMA INTL PHARMS	<u>12.5MG;10MG</u>	<u>A076265</u> <u>001</u>	Jul 08, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076265</u> <u>002</u>	Jul 08, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076265</u> <u>003</u>	Jul 08, 2002
<u>AB</u>	INVAGEN PHARMS	<u>12.5MG;10MG</u>	<u>A204058</u> <u>001</u>	May 23, 2017
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A204058</u> <u>002</u>	May 23, 2017
<u>AB</u>		<u>25MG;20MG</u>	<u>A204058</u> <u>003</u>	May 23, 2017
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>12.5MG;10MG</u>	<u>A075776</u> <u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A075776</u> <u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A075776</u> <u>003</u>	Jul 01, 2002
<u>AB</u>	LUPIN	<u>12.5MG;10MG</u>	<u>A077912</u> <u>001</u>	Sep 27, 2006
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A077912</u> <u>002</u>	Sep 27, 2006
<u>AB</u>		<u>25MG;20MG</u>	<u>A077912</u> <u>003</u>	Sep 27, 2006
<u>AB</u>	MYLAN	<u>12.5MG;10MG</u>	<u>A076113</u> <u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076113</u> <u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076113</u> <u>003</u>	Jul 01, 2002
<u>AB</u>	PRINSTON INC	<u>12.5MG;10MG</u>	<u>A076230</u> <u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076230</u> <u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076230</u> <u>003</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>12.5MG;10MG</u>	<u>A076262</u> <u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076262</u> <u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076262</u> <u>003</u>	Jul 01, 2002
<u>AB</u>	SUN PHARM INDs LTD	<u>12.5MG;10MG</u>	<u>A076007</u> <u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076007</u> <u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076007</u> <u>003</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>12.5MG;10MG</u>	<u>A076194</u> <u>003</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>A076194</u> <u>001</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>A076194</u> <u>002</u>	Jul 01, 2002
<b>ZESTORETIC</b>				
<u>AB</u>	+ ALVOGEN MALTA	<u>12.5MG;10MG</u>	<u>N019888</u> <u>003</u>	Nov 18, 1993
<u>AB</u>	+!	<u>12.5MG;20MG</u>	<u>N019888</u> <u>001</u>	Sep 20, 1990
<u>AB</u>	+!	<u>25MG;20MG</u>	<u>N019888</u> <u>002</u>	Jul 20, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

HYZAAR

<u>AB</u>	+	MERCK SHARP DOHME	<u>12.5MG;50MG</u>	<u>N020387</u> <u>001</u>	Apr 28, 1995
<u>AB</u>	+		<u>12.5MG;100MG</u>	<u>N020387</u> <u>003</u>	Oct 20, 2005
<u>AB</u>	+!		<u>25MG;100MG</u>	<u>N020387</u> <u>002</u>	Nov 10, 1998

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>12.5MG;50MG</u>	<u>A091617</u> <u>001</u>	Feb 17, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091617</u> <u>002</u>	Feb 17, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A091617</u> <u>003</u>	Feb 17, 2012
<u>AB</u>	APOTEX	<u>12.5MG;50MG</u>	<u>A090150</u> <u>001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A090150</u> <u>002</u>	Aug 11, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A090150</u> <u>003</u>	Oct 06, 2010
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG;50MG</u>	<u>A091629</u> <u>001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091629</u> <u>002</u>	Oct 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A091629</u> <u>003</u>	Jan 06, 2010
<u>AB</u>	CADISTA PHARMS	<u>12.5MG;50MG</u>	<u>A201845</u> <u>001</u>	Sep 18, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-214 (of 436)

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>		<u>12.5MG;100MG</u>	<u>A201845 002</u>	Sep 18, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A201845 003</u>	Sep 18, 2012
<u>AB</u>	IPCA LABS LTD	<u>12.5MG;50MG</u>	<u>A201682 001</u>	Mar 01, 2013
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A201682 002</u>	Mar 01, 2013
<u>AB</u>		<u>25MG;100MG</u>	<u>A201682 003</u>	Mar 01, 2013
<u>AB</u>	LUPIN LTD	<u>12.5MG;50MG</u>	<u>A078245 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A078245 002</u>	May 21, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078245 003</u>	Oct 06, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>12.5MG;50MG</u>	<u>A202289 001</u>	Aug 09, 2012
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A202289 002</u>	Aug 09, 2012
<u>AB</u>		<u>25MG;100MG</u>	<u>A202289 003</u>	Aug 09, 2012
<u>AB</u>	MYLAN	<u>12.5MG;50MG</u>	<u>A091652 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A091652 002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A091652 003</u>	Oct 06, 2010
<u>AB</u>	PRINSTON INC	<u>12.5MG;50MG</u>	<u>A204901 001</u>	Nov 06, 2017
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A204901 002</u>	Nov 06, 2017
<u>AB</u>		<u>25MG;100MG</u>	<u>A204901 003</u>	Nov 06, 2017
<u>AB</u>	SANDOZ	<u>12.5MG;50MG</u>	<u>A077948 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077948 003</u>	Aug 19, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077948 002</u>	Oct 06, 2010
<u>AB</u>	TEVA PHARMS	<u>12.5MG;50MG</u>	<u>A077157 001</u>	Apr 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077157 002</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077157 003</u>	Apr 06, 2010
<u>AB</u>	TORRENT PHARMS	<u>12.5MG;50MG</u>	<u>A090528 001</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A090528 003</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A090528 002</u>	Oct 06, 2010
<u>AB</u>	UNICHEM LABS LTD	<u>12.5MG;50MG</u>	<u>A204832 001</u>	Jul 21, 2017
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A204832 002</u>	Jul 21, 2017
<u>AB</u>		<u>25MG;100MG</u>	<u>A204832 003</u>	Jul 21, 2017
<u>AB</u>	WEST-WARD PHARMS INT	<u>12.5MG;50MG</u>	<u>A077732 002</u>	Oct 06, 2010
<u>AB</u>		<u>12.5MG;100MG</u>	<u>A077732 001</u>	Apr 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A077732 003</u>	Oct 06, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>12.5MG;50MG</u>	<u>A078385 001</u>	Oct 06, 2010
<u>AB</u>		<u>25MG;100MG</u>	<u>A078385 002</u>	Oct 06, 2010

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

MYLAN	15MG;250MG	A070265 002	Jan 23, 1986
!	25MG;250MG	A070265 001	Jan 23, 1986

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

DUTOPROL

+ CONCORDIA PHARMS INC	12.5MG;EQ 25MG TARTRATE	N021956 001	Aug 28, 2006
+	12.5MG;EQ 50MG TARTRATE	N021956 002	Aug 28, 2006
+	12.5MG;EQ 100MG TARTRATE	N021956 003	Aug 28, 2006

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

<u>AB</u> + US PHARMS HOLDINGS	<u>25MG;50MG</u>	<u>N018303 001</u>	Dec 31, 1984
I			
<u>AB</u> +!	<u>25MG;100MG</u>	<u>N018303 002</u>	Dec 31, 1984
<u>METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE</u>			
<u>AB</u> ALEMBIC PHARMS LTD	<u>25MG;50MG</u>	<u>A202870 001</u>	Nov 06, 2013
	<u>25MG;100MG</u>	<u>A202870 002</u>	Nov 06, 2013
<u>AB</u>	<u>50MG;100MG</u>	<u>A202870 003</u>	Nov 06, 2013
<u>AB</u> MYLAN	<u>25MG;50MG</u>	<u>A076792 001</u>	Aug 20, 2004
	<u>25MG;100MG</u>	<u>A076792 002</u>	Aug 20, 2004
<u>AB</u>	<u>50MG;100MG</u>	<u>A076792 003</u>	Aug 20, 2004
<u>AB</u> SUN PHARM INDs	<u>25MG;50MG</u>	<u>A090654 001</u>	Jan 19, 2012
	<u>25MG;100MG</u>	<u>A090654 002</u>	Jan 19, 2012
<u>AB</u>	<u>50MG;100MG</u>	<u>A090654 003</u>	Jan 19, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-215 (of 436)

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	GLENMARK PHARMS	<u>12.5MG; 7.5MG</u>	<u>A090718 001</u>	Mar 17, 2010
<u>AB</u>		<u>12.5MG; 15MG</u>	<u>A090718 002</u>	Mar 17, 2010
<u>AB</u>		<u>25MG; 15MG</u>	<u>A090718 003</u>	Mar 17, 2010
<u>AB</u>	HERITAGE PHARMS INC	<u>12.5MG; 7.5MG</u>	<u>A202150 001</u>	Mar 07, 2014
<u>AB</u>		<u>12.5MG; 15MG</u>	<u>A202150 002</u>	Mar 07, 2014
<u>AB</u>		<u>25MG; 15MG</u>	<u>A202150 003</u>	Mar 07, 2014
<u>AB</u>	TEVA	<u>12.5MG; 7.5MG</u>	<u>A076980 001</u>	Mar 07, 2007
<u>AB</u>		<u>12.5MG; 15MG</u>	<u>A076980 003</u>	Mar 07, 2007
<u>AB</u>	!	<u>25MG; 15MG</u>	<u>A076980 002</u>	Mar 07, 2007

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

<u>AB</u>	+	DAIICHI SANKYO	<u>12.5MG; 20MG</u>	<u>N021532 002</u>	Jun 05, 2003
<u>AB</u>	+		<u>12.5MG; 40MG</u>	<u>N021532 003</u>	Jun 05, 2003
<u>AB</u>	!+		<u>25MG; 40MG</u>	<u>N021532 005</u>	Jun 05, 2003

OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>12.5MG; 20MG</u>	<u>A204233 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A204233 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A204233 003</u>	Apr 24, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>12.5MG; 20MG</u>	<u>A205391 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A205391 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A205391 003</u>	Apr 24, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>12.5MG; 20MG</u>	<u>A078827 001</u>	Oct 26, 2016
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A078827 002</u>	Oct 26, 2016
<u>AB</u>		<u>25MG; 40MG</u>	<u>A078827 003</u>	Oct 26, 2016
<u>AB</u>	PRINSTON INC	<u>12.5MG; 20MG</u>	<u>A207804 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A207804 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A207804 003</u>	Apr 24, 2017
<u>AB</u>	TEVA PHARMS USA	<u>12.5MG; 20MG</u>	<u>A200532 001</u>	Apr 24, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A200532 002</u>	Apr 24, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A200532 003</u>	Apr 24, 2017
<u>AB</u>	TORRENT PHARMS LTD	<u>12.5MG; 20MG</u>	<u>A206515 001</u>	May 03, 2017
<u>AB</u>		<u>12.5MG; 40MG</u>	<u>A206515 002</u>	May 03, 2017
<u>AB</u>		<u>25MG; 40MG</u>	<u>A206515 003</u>	May 03, 2017

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ANI PHARMS INC	<u>25MG; 40MG</u>	<u>A072042 001</u>	Mar 14, 1988
<u>AB</u>		<u>25MG; 80MG</u>	<u>A072043 001</u>	Mar 14, 1988
<u>AB</u>	! MYLAN	<u>25MG; 40MG</u>	<u>A070947 002</u>	Mar 04, 1987
<u>AB</u>	!	<u>25MG; 80MG</u>	<u>A070947 001</u>	Apr 01, 1987

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCURETIC

<u>AB</u>	+	PFIZER PHARMS	<u>12.5MG; EQ 10MG BASE</u>	<u>N020125 001</u>	Dec 28, 1999
<u>AB</u>	+		<u>12.5MG; EQ 20MG BASE</u>	<u>N020125 002</u>	Dec 28, 1999
<u>AB</u>	!+		<u>25MG; EQ 20MG BASE</u>	<u>N020125 003</u>	Dec 28, 1999

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX CORP	<u>12.5MG; EQ 10MG BASE</u>	<u>A091524 001</u>	Mar 12, 2013
<u>AB</u>		<u>12.5MG; EQ 20MG BASE</u>	<u>A091524 002</u>	Mar 12, 2013
<u>AB</u>		<u>25MG; EQ 20MG BASE</u>	<u>A091524 003</u>	Mar 12, 2013
<u>AB</u>	AUROBINDO PHARMA	<u>12.5MG; EQ 10MG BASE</u>	<u>A078450 001</u>	Aug 24, 2007
<u>AB</u>		<u>12.5MG; EQ 20MG BASE</u>	<u>A078450 002</u>	Aug 24, 2007
<u>AB</u>		<u>25MG; EQ 20MG BASE</u>	<u>A078450 003</u>	Aug 24, 2007
<u>AB</u>	INVAGEN PHARMS	<u>12.5MG; EQ 10MG BASE</u>	<u>A201356 001</u>	Apr 20, 2011
<u>AB</u>		<u>12.5MG; EQ 20MG BASE</u>	<u>A201356 002</u>	Apr 20, 2011
<u>AB</u>		<u>25MG; EQ 20MG BASE</u>	<u>A201356 003</u>	Apr 20, 2011
<u>AB</u>	MYLAN	<u>12.5MG; EQ 10MG BASE</u>	<u>A077093 001</u>	Mar 28, 2005
<u>AB</u>		<u>12.5MG; EQ 20MG BASE</u>	<u>A077093 002</u>	Mar 28, 2005
<u>AB</u>		<u>25MG; EQ 20MG BASE</u>	<u>A077093 003</u>	Mar 28, 2005

QUINARETIC

<u>AB</u>	GAVIS PHARMS	<u>12.5MG; EQ 10MG BASE</u>	<u>A076374 001</u>	Mar 31, 2004
<u>AB</u>		<u>12.5MG; EQ 20MG BASE</u>	<u>A076374 002</u>	Mar 31, 2004
<u>AB</u>		<u>25MG; EQ 20MG BASE</u>	<u>A076374 003</u>	Mar 31, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-216 (of 436)

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

**ALDACTAZIDE**

<b>AB</b>	+	GD SEARLE LLC	<u>25MG;25MG</u>	<b>N012616 004</b>	Dec 30, 1982
<b>SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE</b>					
<b>AB</b>		MYLAN	<u>25MG;25MG</u>	<b>A086513 001</b>	
<b>AB</b>		SUN PHARM INDUSTRIES	<u>25MG;25MG</u>	<b>A089534 001</b>	Jul 02, 1987
		ALDACTAZIDE			
+!		GD SEARLE LLC	50MG;50MG	N012616 005	Dec 30, 1982

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL

**MICARDIS HCT**

<b>AB</b>	+	BOEHRINGER INGELHEIM	<u>12 .5MG;40MG</u>	<b>N021162 001</b>	Nov 17, 2000
<b>AB</b>	+		<u>12 .5MG;80MG</u>	<b>N021162 002</b>	Nov 17, 2000
<b>AB</b>	+!		<u>25MG;80MG</u>	<b>N021162 003</b>	Apr 19, 2004
<b>TELMISARTAN AND HYDROCHLOROTHIAZIDE</b>					
<b>AB</b>		ALEMBIC PHARMS LTD	<u>12 .5MG;40MG</u>	<b>A203010 001</b>	Feb 25, 2014
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A203010 002</b>	Feb 25, 2014
<b>AB</b>			<u>25MG;80MG</u>	<b>A203010 003</b>	Feb 25, 2014
<b>AB</b>		AUROBINDO PHARMA LTD	<u>12 .5MG;40MG</u>	<b>A208727 001</b>	Dec 15, 2016
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A208727 002</b>	Dec 15, 2016
<b>AB</b>			<u>25MG;80MG</u>	<b>A208727 003</b>	Dec 15, 2016
<b>AB</b>		LUPIN LTD	<u>12 .5MG;40MG</u>	<b>A091351 001</b>	Aug 07, 2014
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A091351 002</b>	Aug 07, 2014
<b>AB</b>			<u>25MG;80MG</u>	<b>A091351 003</b>	Aug 07, 2014
<b>AB</b>		MACLEODS PHARMS LTD	<u>12 .5MG;40MG</u>	<b>A204169 001</b>	Nov 02, 2015
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A204169 002</b>	Nov 02, 2015
<b>AB</b>			<u>25MG;80MG</u>	<b>A204169 003</b>	Nov 02, 2015
<b>AB</b>		MYLAN PHARMS INC	<u>12 .5MG;40MG</u>	<b>A091648 001</b>	Feb 25, 2014
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A091648 002</b>	Feb 25, 2014
<b>AB</b>			<u>25MG;80MG</u>	<b>A091648 003</b>	Feb 25, 2014
<b>AB</b>		PRINSTON INC	<u>12 .5MG;40MG</u>	<b>A209028 001</b>	Nov 06, 2017
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A209028 002</b>	Nov 06, 2017
<b>AB</b>			<u>25MG;80MG</u>	<b>A209028 003</b>	Nov 06, 2017
<b>AB</b>		TORRENT PHARMS LTD	<u>12 .5MG;40MG</u>	<b>A201192 001</b>	Feb 25, 2014
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A201192 002</b>	Feb 25, 2014
<b>AB</b>			<u>25MG;80MG</u>	<b>A201192 003</b>	Feb 25, 2014
<b>AB</b>		ZYDUS PHARMS USA INC	<u>12 .5MG;40MG</u>	<b>A204221 001</b>	Aug 15, 2017
<b>AB</b>			<u>12 .5MG;80MG</u>	<b>A204221 002</b>	Aug 15, 2017
<b>AB</b>			<u>25MG;80MG</u>	<b>A204221 003</b>	Aug 15, 2017

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

**DYAZIDE**

<b>AB</b>	+!	GLAXOSMITHKLINE LLC	<u>25MG;37 .5MG</u>	<b>N016042 003</b>	Mar 03, 1994
<b>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</b>					
<b>AB</b>		DURAMED PHARMS BARR	<u>25MG;37 .5MG</u>	<b>A075052 001</b>	Jun 18, 1999
<b>AB</b>		IVAX SUB TEVA PHARMS	<u>25MG;50MG</u>	<b>A074259 001</b>	Mar 30, 1995
<b>AB</b>		LANNETT HOLDINGS INC	<u>25MG;37 .5MG</u>	<b>A201407 001</b>	Dec 09, 2011
<b>AB</b>		MYLAN	<u>25MG;37 .5MG</u>	<b>A074701 001</b>	Jun 07, 1996
<b>AB</b>		SANDOZ	<u>25MG;37 .5MG</u>	<b>A074821 001</b>	Jun 05, 1997
<b>AB</b>	!		<u>25MG;50MG</u>	<b>A073191 001</b>	Jul 31, 1991

TABLET; ORAL

**MAXZIDE**

<b>AB</b>	+!	MYLAN PHARMS INC	<u>50MG;75MG</u>	<b>N019129 001</b>	Oct 22, 1984
<b>MAXZIDE-25</b>					
<b>AB</b>	+	MYLAN PHARMS INC	<u>25MG;37 .5MG</u>	<b>N019129 003</b>	May 13, 1988
<b>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</b>					
<b>AB</b>		ANI PHARMS INC	<u>50MG;75MG</u>	<b>A073467 001</b>	Jan 31, 1996
<b>AB</b>		APOTEX INC	<u>25MG;37 .5MG</u>	<b>A071251 002</b>	May 05, 1998
<b>AB</b>			<u>50MG;75MG</u>	<b>A071251 001</b>	Apr 17, 1988
<b>AB</b>		PLIVA	<u>25MG;37 .5MG</u>	<b>A074026 001</b>	Apr 26, 1996
<b>AB</b>		SANDOZ	<u>25MG;37 .5MG</u>	<b>A073281 001</b>	Apr 30, 1992
<b>AB</b>			<u>50MG;75MG</u>	<b>A072011 001</b>	Jun 17, 1988
<b>AB</b>		WATSON LABS	<u>25MG;37 .5MG</u>	<b>A073449 001</b>	Sep 23, 1993
<b>AB</b>			<u>50MG;75MG</u>	<b>A071851 001</b>	Nov 30, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-217 (of 436)

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

<u>AB</u>	+	NOVARTIS	<u>12.5MG;80MG</u>	<u>N020818 001</u>	Mar 06, 1998
<u>AB</u>	+		<u>12.5MG;160MG</u>	<u>N020818 002</u>	Mar 06, 1998
<u>AB</u>	+		<u>12.5MG;320MG</u>	<u>N020818 004</u>	Apr 28, 2006
<u>AB</u>	+		<u>25MG;160MG</u>	<u>N020818 003</u>	Jan 17, 2002
<u>AB</u>	+!		<u>25MG;320MG</u>	<u>N020818 005</u>	Apr 28, 2006
<b>VALSARTAN AND HYDROCHLOROTHIAZIDE</b>					
<u>AB</u>		ALEMBIC PHARMS LTD	<u>12.5MG;80MG</u>	<u>A201662 001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A201662 002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A201662 003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A201662 004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A201662 005</u>	Mar 21, 2013
<u>AB</u>		APOTEX INC	<u>12.5MG;80MG</u>	<u>A203026 001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A203026 002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A203026 003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A203026 004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A203026 005</u>	Mar 21, 2013
<u>AB</u>		AUROBINDO PHARMA LTD	<u>12.5MG;80MG</u>	<u>A202519 001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A202519 002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A202519 003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A202519 004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A202519 005</u>	Mar 21, 2013
<u>AB</u>		LUPIN LTD	<u>12.5MG;80MG</u>	<u>A078946 003</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A078946 004</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A078946 001</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A078946 005</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A078946 002</u>	Mar 21, 2013
<u>AB</u>		MACLEODS PHARMS LTD	<u>12.5MG;80MG</u>	<u>A203145 001</u>	Apr 19, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A203145 002</u>	Apr 19, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A203145 003</u>	Apr 19, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A203145 004</u>	Apr 19, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A203145 005</u>	Apr 19, 2013
<u>AB</u>		MYLAN PHARMS INC	<u>12.5MG;80MG</u>	<u>A078020 001</u>	Sep 21, 2012
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A078020 002</u>	Sep 21, 2012
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A078020 004</u>	Sep 21, 2012
<u>AB</u>			<u>25MG;160MG</u>	<u>A078020 003</u>	Sep 21, 2012
<u>AB</u>			<u>25MG;320MG</u>	<u>A078020 005</u>	Sep 21, 2012
<u>AB</u>		PRINSTON INC	<u>12.5MG;80MG</u>	<u>A206083 001</u>	Feb 08, 2016
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A206083 002</u>	Feb 08, 2016
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A206083 003</u>	Feb 08, 2016
<u>AB</u>			<u>25MG;160MG</u>	<u>A206083 004</u>	Feb 08, 2016
<u>AB</u>			<u>25MG;320MG</u>	<u>A206083 005</u>	Feb 08, 2016
<u>AB</u>		WATSON LABS TEVA	<u>12.5MG;80MG</u>	<u>A091519 001</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;160MG</u>	<u>A091519 002</u>	Mar 21, 2013
<u>AB</u>			<u>12.5MG;320MG</u>	<u>A091519 003</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;160MG</u>	<u>A091519 004</u>	Mar 21, 2013
<u>AB</u>			<u>25MG;320MG</u>	<u>A091519 005</u>	Mar 21, 2013

HYDROCODONE BITARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

ZOHYDRO ER

+!	PERNIX IRELAND PAIN	10MG	<u>N202880 001</u>	Oct 25, 2013
+		15MG	<u>N202880 002</u>	Oct 25, 2013
+		20MG	<u>N202880 003</u>	Oct 25, 2013
+		30MG	<u>N202880 004</u>	Oct 25, 2013
+		40MG	<u>N202880 005</u>	Oct 25, 2013
+		50MG	<u>N202880 006</u>	Oct 25, 2013

TABLET, EXTENDED RELEASE; ORAL

HYSINGLA

+!	PURDUE PHARMA LP	20MG	<u>N206627 001</u>	Nov 20, 2014
+		30MG	<u>N206627 002</u>	Nov 20, 2014
+		40MG	<u>N206627 003</u>	Nov 20, 2014
+		60MG	<u>N206627 004</u>	Nov 20, 2014
+		80MG	<u>N206627 005</u>	Nov 20, 2014
+		100MG	<u>N206627 006</u>	Nov 20, 2014
+		120MG	<u>N206627 007</u>	Nov 20, 2014

VANTRELA ER

+	TEVA BRANDED PHARM	15MG	<u>N207975 001</u>	Jan 17, 2017
+		30MG	<u>N207975 002</u>	Jan 17, 2017
+		45MG	<u>N207975 003</u>	Jan 17, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-218 (of 436)

HYDROCODONE BITARTRATE

TABLET, EXTENDED RELEASE; ORAL  
 VANTRELA ER

+ 60MG  
 + 90MG

N207975 004 Jan 17, 2017  
 N207975 005 Jan 17, 2017

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

<b>AB</b>	ACTAVIS LABS FL INC	<u>7.5MG;200MG</u>
<b>AB</b>	AMNEAL PHARMS NY	<u>5MG;200MG</u>
<b>AB</b>	!	<u>7.5MG;200MG</u>
<b>AB</b>	AUROLIFE PHARMA LLC	<u>7.5MG;200MG</u>
<b>AB</b>	SUN PHARM INDs INC	<u>2.5MG;200MG</u>
<b>AB</b>		<u>5MG;200MG</u>
<b>AB</b>		<u>7.5MG;200MG</u>
<b>AB</b>		<u>10MG;200MG</u>
<b>AB</b>	TEVA	<u>7.5MG;200MG</u>
<b>AB</b>	VINTAGE PHARMS	<u>5MG;200MG</u>
<b>AB</b>		<u>7.5MG;200MG</u>
<b>AB</b>		<u>10MG;200MG</u>

<b>A076604 001</b>	Dec 31, 2003
<b>A076642 002</b>	Mar 18, 2004
<b>A076642 001</b>	Oct 12, 2004
<b>A204575 001</b>	Jun 02, 2016
<b>A091633 001</b>	May 28, 2013
<b>A091633 002</b>	May 28, 2013
<b>A091633 003</b>	May 28, 2013
<b>A091633 004</b>	May 28, 2013
<b>A076023 001</b>	Apr 11, 2003
<b>A077727 001</b>	Nov 06, 2006
<b>A077723 001</b>	Nov 06, 2006
<b>A077723 002</b>	Nov 06, 2006

REPREXAIN

<b>AB</b>	AMNEAL PHARMS NY	<u>2.5MG;200MG</u>
<b>AB</b>		<u>10MG;200MG</u>

<b>A076642 003</b>	Oct 19, 2007
<b>A076642 004</b>	Oct 19, 2007

HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

<b>AA</b>	MAYNE PHARMA INC	<u>5MG/5ML;60MG/5ML</u>
<b>AA</b>	PADDOCK LLC	<u>5MG/5ML;60MG/5ML</u>

<b>A205658 001</b>	Nov 17, 2015
<b>A204658 001</b>	Apr 29, 2014

REZIRA

<b>AA</b>	+!	CYPRESS PHARM	<u>5MG/5ML;60MG/5ML</u>
-----------	----	---------------	-------------------------

<b>N022442 001</b>	Jun 08, 2011
--------------------	--------------

HYDROCORTISONE

CREAM; TOPICAL

ALA-CORT

<b>AT</b>	CROWN LABS	<u>2.5%</u>
<b>AT</b>		<u>1%</u>

<b>A080706 007</b>	Jan 05, 2016
<b>A080706 006</b>	

ANUSOL HC

<b>AT</b>	SALIX PHARMS	<u>2.5%</u>
-----------	--------------	-------------

<b>A088250 001</b>	Jun 06, 1984
--------------------	--------------

HYDROCORTISONE

<b>AT</b>	ACTAVIS MID ATLANTIC	<u>1%</u>
<b>AT</b>		<u>2.5%</u>
<b>AT</b>	FOUGERA PHARMS INC	<u>1%</u>
<b>AT</b>	!	<u>2.5%</u>
<b>AT</b>	PERRIGO NEW YORK	<u>2.5%</u>
<b>AT</b>	RISING PHARMS INC	<u>2.5%</u>
<b>AT</b>	TARO	<u>2.5%</u>
<b>AT</b>	VINTAGE PHARMS	<u>2.5%</u>

<b>A087795 001</b>	May 03, 1983
<b>A089682 001</b>	Mar 10, 1988
<b>A080693 003</b>	
<b>A089414 001</b>	Dec 16, 1986
<b>A085025 001</b>	
<b>A040879 001</b>	Aug 20, 2010
<b>A088799 001</b>	Nov 09, 1984
<b>A040503 001</b>	Mar 12, 2004

ENEMA; RECTAL

COLOCORT

<b>AB</b>	PADDOCK LLC	<u>100MG/60ML</u>
-----------	-------------	-------------------

<b>A075172 001</b>	Dec 03, 1999
--------------------	--------------

CORTENEMA

<b>AB</b>	+!	ANI PHARMS	<u>100MG/60ML</u>
-----------	----	------------	-------------------

<b>N016199 001</b>	
--------------------	--

HYDROCORTISONE

<b>AB</b>	TEVA PHARMS	<u>100MG/60ML</u>
-----------	-------------	-------------------

<b>A074171 001</b>	May 27, 1994
--------------------	--------------

LOTION; TOPICAL

HYDROCORTISONE

<b>AT</b>	FOUGERA PHARMS	<u>2.5%</u>
<b>AT</b>	TARO	<u>2.5%</u>
<b>AT</b>	VINTAGE PHARMS	<u>2.5%</u>

<b>A040351 001</b>	Jul 25, 2000
<b>A040247 001</b>	Jul 23, 1999
<b>A040417 001</b>	Jul 30, 2003

STIE-CORT

<b>AT</b>	PERRIGO CO	<u>2.5%</u>
-----------	------------	-------------

<b>A089074 001</b>	Nov 26, 1985
--------------------	--------------

ALA-SCALP

	CROWN LABS	<u>2%</u>
--	------------	-----------

<b>A083231 001</b>	
--------------------	--

OINTMENT; TOPICAL

HYDROCORTISONE

<b>AT</b>	ACTAVIS MID ATLANTIC	<u>1%</u>
<b>AT</b>	FOUGERA PHARMS	<u>1%</u>
<b>AT</b>	!	<u>2.5%</u>
<b>AT</b>	PERRIGO NEW YORK	<u>2.5%</u>
<b>AT</b>	TARO	<u>1%</u>

<b>A087796 001</b>	Oct 13, 1982
<b>A080692 001</b>	
<b>A081203 001</b>	May 28, 1993
<b>A085027 001</b>	
<b>A086257 001</b>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-219 (of 436)

HYDROCORTISONE

OINTMENT;TOPICAL

HYDROCORTISONE IN ABSORBASE

AT CMP PHARMA INC 1% A088138 001 Sep 06, 1985

SOLUTION;TOPICAL

TEXACORT

! MISSION PHARMA 2.5%

A081271 001 Apr 17, 1992

TABLET;ORAL

CORTEF

AB + PHARMACIA AND 5MG N008697 003

UPJOHN

AB + 10MG N008697 001

AB +! 20MG N008697 002

HYDROCORTISONE

AB HIKMA INTL PHARMS 5MG A083365 002 Feb 23, 2015

AB 10MG A083365 003 Feb 23, 2015

AB 20MG A083365 001

AB IMPAX LABS INC 5MG A040646 001 Mar 30, 2007

AB 10MG A040646 002 Mar 30, 2007

AB 20MG A040646 003 Mar 30, 2007

AB PII 5MG A207029 001 Apr 27, 2017

AB 10MG A207029 002 Apr 27, 2017

AB 20MG A207029 003 Apr 27, 2017

AB VINTAGE 5MG A040761 001 Jul 16, 2007

AB 10MG A040761 002 Jul 16, 2007

AB 20MG A040761 003 Jul 16, 2007

HYDROCORTISONE ACETATE

AEROSOL, METERED;RECTAL

CORTIFOAM

+! MYLAN SPECIALITY LP 10%

N017351 001 Feb 10, 1982

CREAM;TOPICAL

MICORT-HC

SEBELA IRELAND LTD 2.5%

A040396 001 Feb 27, 2001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM;TOPICAL

CORTISPORIN

+! MONARCH PHARMS 0.5%;EQ 3.5MG BASE/GM;10,000 UNITS/GM

N050218 001 Aug 09, 1985

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED;TOPICAL

EPIFOAM

BX MYLAN SPECIALITY LP 1%;1%

A086457 001

PROCTOFOAM HC

BX MYLAN SPECIALITY LP 1%;1%

A086195 001

CREAM;TOPICAL

PRAMOSONE

SEBELA IRELAND LTD 0.5%;1%

A083778 001

1%;1%

A085368 001

LOTION;TOPICAL

PRAMOSONE

SEBELA IRELAND LTD 1%;1%

A085980 001

2.5%;1%

A085979 001

HYDROCORTISONE ACETATE; UREA

CREAM;TOPICAL

U-CORT

TARO

1%;10%

A089472 001 Jun 13, 1988

HYDROCORTISONE BUTYRATE

CREAM;TOPICAL

HYDROCORTISONE BUTYRATE

AB1 TARO PHARM INDS 0.1%

A076654 001 Aug 03, 2005

LOCOID

AB1 +! PRECISION DERMAT 0.1%

N018514 001 Mar 31, 1982

HYDROCORTISONE BUTYRATE

AB2 ACTAVIS MID 0.1%

A205134 001 Dec 08, 2017

ATLANTIC

AB2 GLENMARK GENERICS 0.1%

A202145 001 Sep 27, 2013

LOCOID LIPOCREAM

AB2 +! PRECISION DERMAT 0.1%

N020769 001 Sep 08, 1997

LOTION;TOPICAL

HYDROCORTISONE BUTYRATE

AB TELIGENT PHARMA INC 0.1%

A209556 001 Nov 21, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-220 (of 436)

HYDROCORTISONE BUTYRATE

LOTION;TOPICAL

LOCOID

<u>AB</u>	+!	PRECISION DERMAT	<u>0.1%</u>	<u>N022076 001</u> May 18, 2007
OINTMENT;TOPICAL				

HYDROCORTISONE BUTYRATE

<u>AB</u>		TARO	<u>0.1%</u>	<u>A076842 001</u> Dec 27, 2004
-----------	--	------	-------------	---------------------------------

LOCOID

<u>AB</u>	+!	PRECISION DERMAT	<u>0.1%</u>	<u>N018652 001</u> Oct 29, 1982
SOLUTION;TOPICAL				

HYDROCORTISONE BUTYRATE

<u>AT</u>		TARO PHARM INDS	<u>0.1%</u>	<u>A076364 001</u> Jan 14, 2004
-----------	--	-----------------	-------------	---------------------------------

LOCOID

<u>AT</u>	+!	PRECISION DERMAT	<u>0.1%</u>	<u>N019116 001</u> Feb 25, 1987
-----------	----	------------------	-------------	---------------------------------

HYDROCORTISONE PROBUTATE

CREAM;TOPICAL

PANDEL

+!	FUGERA PHARMS	0.1%	N020453 001 Feb 28, 1997
----	---------------	------	--------------------------

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE;INJECTION

SOLU-CORTEF

+!	PHARMACIA AND UPJOHN	EQ 100MG BASE/VIAL	N009866 001
+!		EQ 250MG BASE/VIAL	N009866 002
+!		EQ 500MG BASE/VIAL	N009866 003
+!		EQ 1GM BASE/VIAL	N009866 004

HYDROCORTISONE VALERATE

CREAM;TOPICAL

HYDROCORTISONE VALERATE

<u>AB</u>		PERRIGO NEW YORK	<u>0.2%</u>	<u>A075666 001</u> May 24, 2000
-----------	--	------------------	-------------	---------------------------------

<u>AB</u>	!	TARO	<u>0.2%</u>	<u>A075042 001</u> Aug 25, 1998
-----------	---	------	-------------	---------------------------------

OINTMENT;TOPICAL

HYDROCORTISONE VALERATE

!	TARO	0.2%	A075043 001 Aug 25, 1998
---	------	------	--------------------------

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

CORTISPORIN

<u>AT</u>	+!	MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N050479 001</u>
-----------	----	----------------	--	--------------------

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

<u>AT</u>		AMRING PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A065216 001</u> Oct 31, 2005
-----------	--	---------------	--	---------------------------------

<u>AT</u>		BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064053 001</u> Dec 29, 1995
-----------	--	-----------------	--	---------------------------------

<u>AT</u>		SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062423 001</u> Aug 25, 1983
-----------	--	------------	--	---------------------------------

SUSPENSION/DROPS;OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

!	SANDOZ INC	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062874 001 May 11, 1988
---	------------	-------------------------------------	--------------------------

SUSPENSION/DROPS;OTIC

CASPORYN HC

<u>AT</u>	+!	CASPER PHARMA LLC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N060613 001</u>
-----------	----	-------------------	--	--------------------

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE

<u>AT</u>		AMRING PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A065219 001</u> May 01, 2006
-----------	--	---------------	--	---------------------------------

<u>AT</u>		SANDOZ INC	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A062488 001</u> Nov 06, 1985
-----------	--	------------	--	---------------------------------

OTICAIR

<u>AT</u>		BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>A064065 001</u> Aug 28, 1996
-----------	--	-----------------	--	---------------------------------

HYDROGEN PEROXIDE

SOLUTION;TOPICAL

ESKATA

+	ACLARIS THERAPS INC	40%	N209305 001 Dec 14, 2017
---	---------------------	-----	--------------------------

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE;INJECTION

DILAUDID

<u>AP</u>	+!	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>N019034 003</u> Apr 30, 2009
-----------	----	--------------------	---------------	---------------------------------

<u>AP</u>	+		<u>2MG/ML</u>	<u>N019034 004</u> Apr 30, 2009
-----------	---	--	---------------	---------------------------------

<u>AP</u>	+		<u>4MG/ML</u>	<u>N019034 005</u> Apr 30, 2009
-----------	---	--	---------------	---------------------------------

DILAUDID-HP

<u>AP</u>	+!	FRESENIUS KABI USA	<u>10MG/ML</u>	<u>N019034 001</u> Jan 11, 1984
-----------	----	--------------------	----------------	---------------------------------

HYDROMORPHONE HYDROCHLORIDE

<u>AP</u>		AKORN	<u>10MG/ML</u>	<u>A078228 001</u> Apr 14, 2010
-----------	--	-------	----------------	---------------------------------

<u>AP</u>			<u>10MG/ML</u>	<u>A078261 001</u> Apr 14, 2010
-----------	--	--	----------------	---------------------------------

<u>AP</u>		BARR	<u>10MG/ML</u>	<u>A076444 001</u> Apr 25, 2003
-----------	--	------	----------------	---------------------------------

<u>AP</u>		HOSPIRA INC	<u>1MG/ML</u>	<u>N200403 001</u> Dec 01, 2011
-----------	--	-------------	---------------	---------------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-221 (of 436)

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROMORPHONE HYDROCHLORIDE

<u>AP</u>		<u>2MG/ML</u>	<u>N200403 002</u>	Dec 01, 2011
<u>AP</u>		<u>4MG/ML</u>	<u>N200403 003</u>	Dec 01, 2011
<u>AP</u>		<u>10MG/ML</u>	<u>A078591 001</u>	Jun 17, 2008

SOLUTION; ORAL

DILAUDID

<u>AA</u>	+!	RHODES PHARMS	<u>5MG/5ML</u>	<u>N019891 001</u>	Dec 07, 1992
-----------	----	---------------	----------------	--------------------	--------------

HYDROMORPHONE HYDROCHLORIDE

<u>AA</u>		ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A210176 001</u>	Oct 27, 2017
<u>AA</u>		WEST-WARD PHARMS INT	<u>5MG/5ML</u>	<u>A074653 001</u>	Jul 29, 1998

TABLET; ORAL

DILAUDID

<u>AB</u>	+	RHODES PHARMS	<u>2MG</u>	<u>N019892 003</u>	Nov 09, 2007
<u>AB</u>	+		<u>4MG</u>	<u>N019892 002</u>	Nov 09, 2007
<u>AB</u>	+!		<u>8MG</u>	<u>N019892 001</u>	Dec 07, 1992

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>		AUROLIFE PHARMA LLC	<u>2MG</u>	<u>A205814 001</u>	May 13, 2016
<u>AB</u>			<u>4MG</u>	<u>A205814 002</u>	May 13, 2016
<u>AB</u>			<u>8MG</u>	<u>A205814 003</u>	May 13, 2016
<u>AB</u>		ELITE LABS	<u>8MG</u>	<u>A076723 001</u>	Oct 18, 2005
<u>AB</u>		LANNETT	<u>2MG</u>	<u>A077471 002</u>	Dec 09, 2009
<u>AB</u>			<u>2MG</u>	<u>A078439 001</u>	Dec 09, 2009
<u>AB</u>			<u>4MG</u>	<u>A077471 003</u>	Dec 09, 2009
<u>AB</u>			<u>4MG</u>	<u>A078439 002</u>	Dec 09, 2009
<u>AB</u>			<u>8MG</u>	<u>A077471 001</u>	Dec 09, 2009
<u>AB</u>		SPECGX LLC	<u>2MG</u>	<u>A076855 002</u>	Sep 19, 2007
<u>AB</u>			<u>4MG</u>	<u>A076855 003</u>	Sep 19, 2007
<u>AB</u>			<u>8MG</u>	<u>A076855 001</u>	Dec 23, 2004
<u>AB</u>		WEST-WARD PHARMS INT	<u>4MG</u>	<u>A074597 003</u>	May 29, 2009
<u>AB</u>			<u>8MG</u>	<u>A074597 001</u>	Jul 29, 1998

TABLET, EXTENDED RELEASE; ORAL

EXALGO

<u>AB</u>	+	SPECGX LLC	<u>8MG</u>	<u>N021217 001</u>	Mar 01, 2010
<u>AB</u>	+		<u>12MG</u>	<u>N021217 002</u>	Mar 01, 2010
<u>AB</u>	+		<u>16MG</u>	<u>N021217 003</u>	Mar 01, 2010
<u>AB</u>	+!		<u>32MG</u>	<u>N021217 004</u>	Aug 24, 2012

HYDROMORPHONE HYDROCHLORIDE

<u>AB</u>		ACTAVIS LABS FL INC	<u>8MG</u>	<u>A202144 001</u>	May 12, 2014
<u>AB</u>			<u>12MG</u>	<u>A202144 002</u>	May 12, 2014
<u>AB</u>			<u>16MG</u>	<u>A202144 003</u>	May 12, 2014
<u>AB</u>			<u>32MG</u>	<u>A202144 004</u>	Jun 30, 2016
<u>AB</u>		OSMOTICA	<u>8MG</u>	<u>A205629 001</u>	Jul 07, 2016
<u>AB</u>			<u>12MG</u>	<u>A205629 002</u>	Jul 07, 2016
<u>AB</u>			<u>16MG</u>	<u>A205629 003</u>	Jul 07, 2016
<u>AB</u>			<u>32MG</u>	<u>A205629 004</u>	Jul 07, 2016
<u>AB</u>		PADDOCK LLC	<u>8MG</u>	<u>A204278 001</u>	Apr 06, 2015
<u>AB</u>			<u>12MG</u>	<u>A204278 002</u>	Apr 06, 2015
<u>AB</u>			<u>16MG</u>	<u>A204278 003</u>	Apr 06, 2015
<u>AB</u>			<u>32MG</u>	<u>A204278 004</u>	Sep 20, 2017

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

CYANOKIT

+!	SERB SA	5GM/VIAL (5GM/KIT)		N022041 001	Apr 08, 2011
!	ACTAVIS LLC	1MG/ML		A085998 001	

HYDROXYAMPHENATE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREMYD

+!	AKORN	1%;0.25%		N019261 001	Jan 30, 1992
----	-------	----------	--	-------------	--------------

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>	HIKMA PHARMS	<u>200MG</u>	<u>A040760 001</u>	Aug 15, 2007
<u>AB</u>	IPCA LABS LTD	<u>200MG</u>	<u>A040766 001</u>	Jun 14, 2007
<u>AB</u>	MYLAN	<u>200MG</u>	<u>A040274 001</u>	May 29, 1998
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A040104 001</u>	Nov 30, 1995
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>A040081 001</u>	Sep 30, 1994

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-222 (of 436)

HYDROXYCHLOROQUINE SULFATE

TABLET;ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>A040133 001</u>	Nov 30, 1995
<u>AB</u>	ZYDUS PHARMS USA INC	<u>200MG</u>	<u>A040657 001</u>	Sep 21, 2007

PLAQUENIL

<u>AB</u>	+!	CONCORDIA PHARMS INC	<u>200MG</u>	<u>N009768 001</u>
-----------	----	----------------------	--------------	--------------------

HYDROXYPROGESTERONE CAPROATE

INJECTABLE;INJECTION

HYDROXYPROGESTERONE CAPROATE

!	ASPEN GLOBAL INC	250MG/ML	A200271 001	Aug 24, 2015
---	------------------	----------	-------------	--------------

SOLUTION;INTRAMUSCULAR

MAKENA

+	AMAG PHARMA USA	1250MG/5ML (250MG/ML)	N021945 001	Feb 03, 2011
---	-----------------	-----------------------	-------------	--------------

MAKENA PRESERVATIVE FREE

+	AMAG PHARMA USA	250MG/ML (250MG/ML)	N021945 002	Feb 19, 2016
---	-----------------	---------------------	-------------	--------------

HYDROXYPROPYL CELLULOSE

INSERT;OPHTHALMIC

LACRISERT

+	ATON	5MG	N018771 001
---	------	-----	-------------

HYDROXYUREA

CAPSULE;ORAL

HYDREA

<u>AB</u>	+!	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N016295 001</u>
-----------	----	----------------------	--------------	--------------------

HYDROXYUREA

<u>AB</u>	BARR	<u>500MG</u>	<u>A075143 001</u>	Oct 16, 1998
-----------	------	--------------	--------------------	--------------

<u>AB</u>	PAR PHARM	<u>500MG</u>	<u>A075340 001</u>	Feb 24, 1999
-----------	-----------	--------------	--------------------	--------------

DROXIA

+	BRISTOL MYERS SQUIBB	200MG	N016295 002	Feb 25, 1998
---	----------------------	-------	-------------	--------------

+		300MG	N016295 003	Feb 25, 1998
---	--	-------	-------------	--------------

+		400MG	N016295 004	Feb 25, 1998
---	--	-------	-------------	--------------

TABLET;ORAL

SIKLOS

+	ADDMEDICA SAS	100MG	N208843 001	Dec 21, 2017
---	---------------	-------	-------------	--------------

+		1GM	N208843 002	Dec 21, 2017
---	--	-----	-------------	--------------

HYDROXYZINE HYDROCHLORIDE

INJECTABLE;INJECTION

HYDROXYZINE HYDROCHLORIDE

<u>AP</u>	!	FRESENIUS KABI USA	<u>25MG/ML</u>	<u>A087329 001</u>
-----------	---	--------------------	----------------	--------------------

<u>AP</u>	!		<u>50MG/ML</u>	<u>A087329 002</u>
-----------	---	--	----------------	--------------------

<u>AP</u>	LUITPOLD		<u>25MG/ML</u>	<u>A087408 001</u>
-----------	----------	--	----------------	--------------------

<u>AP</u>			<u>50MG/ML</u>	<u>A087408 002</u>
-----------	--	--	----------------	--------------------

SYRUP;ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AA</u>	!	HI TECH PHARMA	<u>10MG/5ML</u>	<u>A040010 001</u>	Oct 28, 1994
-----------	---	----------------	-----------------	--------------------	--------------

<u>AA</u>		SILARK PHARMS INC	<u>10MG/5ML</u>	<u>A201674 001</u>	Aug 21, 2013
-----------	--	-------------------	-----------------	--------------------	--------------

<u>AA</u>	!	VINTAGE PHARMS	<u>10MG/5ML</u>	<u>A040391 001</u>	Apr 10, 2002
-----------	---	----------------	-----------------	--------------------	--------------

<u>AA</u>	!	WOCKHARDT BIO AG	<u>10MG/5ML</u>	<u>A087294 001</u>	Apr 12, 1982
-----------	---	------------------	-----------------	--------------------	--------------

TABLET;ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARM	<u>10MG</u>	<u>A040808 001</u>	Sep 24, 2008
-----------	--------------	-------------	--------------------	--------------

<u>AB</u>		<u>25MG</u>	<u>A040808 002</u>	Sep 24, 2008
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>50MG</u>	<u>A040808 003</u>	Sep 24, 2008
-----------	--	-------------	--------------------	--------------

<u>AB</u>	ECI PHARMS LLC	<u>10MG</u>	<u>A040804 001</u>	Jun 30, 2008
-----------	----------------	-------------	--------------------	--------------

<u>AB</u>		<u>25MG</u>	<u>A040804 002</u>	Jun 30, 2008
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>50MG</u>	<u>A040804 003</u>	Jun 30, 2008
-----------	--	-------------	--------------------	--------------

<u>AB</u>	ELITE LABS INC	<u>10MG</u>	<u>A040604 002</u>	Dec 28, 2004
-----------	----------------	-------------	--------------------	--------------

<u>AB</u>		<u>25MG</u>	<u>A040604 003</u>	Dec 28, 2004
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>50MG</u>	<u>A040604 001</u>	Dec 28, 2004
-----------	--	-------------	--------------------	--------------

<u>AB</u>	HERITAGE PHARMA	<u>10MG</u>	<u>A204279 001</u>	Aug 20, 2014
-----------	-----------------	-------------	--------------------	--------------

<u>AB</u>		<u>25MG</u>	<u>A204279 002</u>	Aug 20, 2014
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>50MG</u>	<u>A204279 003</u>	Aug 20, 2014
-----------	--	-------------	--------------------	--------------

<u>AB</u>	HETERO LABS LTD III	<u>10MG</u>	<u>A040805 001</u>	May 29, 2008
-----------	---------------------	-------------	--------------------	--------------

<u>AB</u>		<u>25MG</u>	<u>A040805 002</u>	May 29, 2008
-----------	--	-------------	--------------------	--------------

<u>AB</u>		<u>50MG</u>	<u>A040805 003</u>	May 29, 2008
-----------	--	-------------	--------------------	--------------

<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>A040812 001</u>	Mar 12, 2008
-----------	----------------	-------------	--------------------	--------------

<u>AB</u>		<u>25MG</u>	<u>A040812 002</u>	Mar 12, 2008
-----------	--	-------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-223 (of 436)

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>		<u>50MG</u>	<u>A040812 003</u>	Mar 12, 2008
<u>AB</u>	KVK TECH	<u>10MG</u>	<u>A040786 001</u>	Mar 20, 2007
<u>AB</u>		<u>25MG</u>	<u>A040787 001</u>	Mar 20, 2007
<u>AB</u>		<u>50MG</u>	<u>A040788 001</u>	Mar 20, 2007
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A091176 001</u>	Jun 07, 2010
<u>AB</u>		<u>25MG</u>	<u>A091176 002</u>	Jun 07, 2010
<u>AB</u>		<u>50MG</u>	<u>A091176 003</u>	Jun 07, 2010
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A040840 002</u>	Mar 31, 2008
<u>AB</u>		<u>25MG</u>	<u>A040840 003</u>	Mar 31, 2008
<u>AB</u>		<u>50MG</u>	<u>A040840 001</u>	Mar 31, 2008
<u>AB</u>	NUVO PHARM	<u>10MG</u>	<u>A207120 001</u>	Mar 29, 2017
<u>AB</u>		<u>50MG</u>	<u>A207122 001</u>	Mar 29, 2017
<u>AB</u>	NUVO PHARM INC	<u>25MG</u>	<u>A207121 001</u>	Mar 29, 2017
<u>AB</u>	PLIVA	<u>10MG</u>	<u>A088617 001</u>	Jan 10, 1986
<u>AB</u>		<u>25MG</u>	<u>A088618 001</u>	Jan 10, 1986
<u>AB</u>		<u>50MG</u>	<u>A088619 001</u>	Jan 10, 1986
<u>AB</u>	PRINSTON INC	<u>10MG</u>	<u>A040579 001</u>	May 27, 2005
<u>AB</u>		<u>25MG</u>	<u>A040574 001</u>	May 27, 2005
<u>AB</u>		<u>50MG</u>	<u>A040580 001</u>	May 27, 2005

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u>	BARR	<u>EQ 25MG HCL</u>	<u>A088496 001</u>	Jun 15, 1984
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>A088487 001</u>	Jun 15, 1984
<u>AB</u>	HERITAGE PHARMA	<u>EQ 25MG HYDROCHLORIDE</u>	<u>A201507 001</u>	Jun 03, 2013
<u>AB</u>		<u>EQ 50MG HYDROCHLORIDE</u>	<u>A201507 002</u>	Jun 03, 2013
<u>AB</u>	IMPAX LABS INC	<u>EQ 25MG HCL</u>	<u>A040156 001</u>	Jul 15, 1996
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>A040156 002</u>	Jul 15, 1996
<u>AB</u>	SANDOZ	<u>EQ 25MG HCL</u>	<u>A087479 001</u>	
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>A086183 001</u>	
	<u>VISTARIL</u>			
<u>AB</u>	+ PFIZER	<u>EQ 25MG HCL</u>	<u>N011459 002</u>	
<u>AB</u>	+!	<u>EQ 50MG HCL</u>	<u>N011459 004</u>	
	HYDROXYZINE PAMOATE			
	BARR	<u>EQ 100MG HCL</u>		
			<u>A088488 001</u>	Jun 15, 1984

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

<u>AP</u>	+! ROCHE	<u>EQ 3MG BASE/3ML</u>	<u>N021858 001</u>	Jan 06, 2006
	<u>IBANDRONATE SODIUM</u>			
<u>AP</u>	ACCORD HLTHCARE	<u>EQ 3MG BASE/3ML</u>	<u>A206058 001</u>	Feb 05, 2016
<u>AP</u>	APOTEX INC	<u>EQ 3MG BASE/3ML</u>	<u>A204222 001</u>	Oct 16, 2015
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 3MG BASE/3ML</u>	<u>A205332 001</u>	Aug 19, 2015
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 3MG BASE/3ML</u>	<u>A203987 001</u>	Sep 02, 2014
<u>AP</u>	MYLAN LABS LTD	<u>EQ 3MG BASE/3ML</u>	<u>A202671 001</u>	Sep 02, 2014
<u>AP</u>	SAGENT PHARMS	<u>EQ 3MG BASE/3ML</u>	<u>A202235 001</u>	Sep 02, 2014
<u>AP</u>	SUN PHARM INDs LTD	<u>EQ 3MG BASE/3ML</u>	<u>A090853 001</u>	Feb 14, 2014

TABLET; ORAL

BONIVA

<u>AB</u>	+! HOFFMANN LA ROCHE	<u>EQ 150MG BASE</u>	<u>N021455 002</u>	Mar 24, 2005
	<u>IBANDRONATE SODIUM</u>			
<u>AB</u>	APOTEX INC	<u>EQ 150MG BASE</u>	<u>A078948 001</u>	Mar 19, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 150MG BASE</u>	<u>A204502 001</u>	Mar 11, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A078997 001</u>	Apr 30, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A206887 001</u>	Oct 31, 2017
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 150MG BASE</u>	<u>A078998 001</u>	Mar 19, 2012
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 150MG BASE</u>	<u>A078996 001</u>	Aug 15, 2012
<u>AB</u>	WATSON LABS TEVA	<u>EQ 150MG BASE</u>	<u>A079003 001</u>	Mar 20, 2012

IBRUTINIB

CAPSULE; ORAL

IMBRUVICA

+	PHARMACYCLICS INC	70MG	N205552 002	Dec 20, 2017
+		140MG	N205552 001	Nov 13, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-224 (of 436)

IBUPROFEN

SOLUTION; INTRAVENOUS

CALDOLOR

+! CUMBERLAND PHARMS 800MG/8ML (100MG/ML)

N022348 002 Jun 11, 2009

SUSPENSION; ORAL

IBUPROFEN

<u>AB</u>	!	ACTAVIS MID ATLANTIC	<u>100MG/5ML</u>
<u>AB</u>		HI-TECH PHARMACAL	<u>100MG/5ML</u>
<u>AB</u>		PERRIGO R AND D	<u>100MG/5ML</u>
<u>AB</u>		TARO	<u>100MG/5ML</u>

TABLET; ORAL

IBU-TAB

<u>AB</u>		ALRA	<u>400MG</u>
<u>AB</u>			<u>600MG</u>

IBUPROFEN

<u>AB</u>		AIPING PHARM INC	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		AMNEAL PHARMS NY	<u>400MG</u>
<u>AB</u>			<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		CONTRACT PHARMACAL	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		DR REDDYS LA	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>	!		<u>800MG</u>
<u>AB</u>		DR REDDYS LABS INC	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		GRANULES INDIA LTD	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		MARKSANS PHARMA	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		PERRIGO R AND D	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		STRIDES PHARMA	<u>400MG</u>
<u>AB</u>			<u>600MG</u>
<u>AB</u>			<u>800MG</u>
<u>AB</u>		VINTAGE PHARMS	<u>400MG</u>

IBUPROFEN LYSINE

INJECTABLE; INTRAVENOUS

IBUPROFEN LYSINE

<u>AP</u>		EXELA PHARMA SCIENCE	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>
-----------	--	----------------------	---

A202402 001 Mar 30, 2016

NEOPROFEN

<u>AP</u>	+	RECORDATI RARE	<u>EQ 20MG BASE/2ML (EQ 10MG BASE/ML)</u>
-----------	---	----------------	---

N021903 001 Apr 13, 2006

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

<u>AB</u>		ACTAVIS ELIZABETH	<u>400MG;5MG</u>
<u>AB</u>	!	BARR LABS INC	<u>400MG;5MG</u>

A078769 001 Jan 04, 2008

A078316 001 Nov 29, 2007

IBUTILIDE FUMARATE

INJECTABLE; INJECTION

COVERT

<u>AP</u>	+	PHARMACIA AND UPJOHN	<u>0 . 1MG/ML</u>
-----------	---	----------------------	-------------------

N020491 001 Dec 28, 1995

IBUTILIDE FUMARATE

<u>AP</u>		LUITPOLD	<u>0 . 1MG/ML</u>
<u>AP</u>		MYLAN INSTITUTIONAL	<u>0 . 1MG/ML</u>
<u>AP</u>			<u>0 . 1MG/ML</u>

A090240 001 Jan 11, 2010

A090643 001 Jan 11, 2010

A090924 001 Jan 11, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-225 (of 436)

ICATIBANT ACETATE

INJECTABLE; SUBCUTANEOUS

FIRAZYR

+! SHIRE ORPHAN THERAP EQ 30MG BASE/3ML (EQ 10MG BASE/ML)

N022150 001 Aug 25, 2011

ICODEXTRIN

SOLUTION; INTRAPERITONEAL

EXTRANEAL

+! BAXTER HLTHCARE 7.5GM/100ML

N021321 001 Dec 20, 2002

ICOSAPENT ETHYL

CAPSULE; ORAL

VASCPEA

+ AMARIN PHARMS 500MG

1GM

N202057 002 Feb 16, 2017

N202057 001 Jul 26, 2012

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFS

AP +! PHARMACIA AND UPJOHN 1MG/ML

N050734 001 Feb 17, 1997

IDARUBICIN HYDROCHLORIDE

AP FRESENIUS KABI USA 1MG/ML

A065440 001 Aug 04, 2009

AP MYLAN LABS LTD 1MG/ML

A200144 001 Oct 11, 2012

AP WEST-WARD PHARMS INT 1MG/ML

A065275 001 Dec 14, 2006

AP 1MG/ML

A065288 001 May 15, 2007

IDARUBICIN HYDROCHLORIDE PFS

AP TEVA PHARMS USA 1MG/ML

A065036 001 May 01, 2002

IDEALALISIB

TABLET; ORAL

ZYDELIG

+ GILEAD SCIENCES INC 100MG

N205858 001 Jul 23, 2014

+! 150MG

N205858 002 Jul 23, 2014

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

AP +! BAXTER HLTHCARE 1GM/VIAL

N019763 001 Dec 30, 1988

AP + 3GM/VIAL

N019763 002 Dec 30, 1988

IFOSFAMIDE

AP ! FRESENIUS KABI USA 1GM/VIAL

A076078 001 May 28, 2002

AP 1GM/20ML (50MG/ML)

A090181 001 Sep 22, 2009

AP ! 3GM/VIAL

A076078 002 May 28, 2002

AP 3GM/60ML (50MG/ML)

A090181 002 Sep 22, 2009

AP MYLAN LABS LTD 1GM/20ML (50MG/ML)

A201689 001 Nov 26, 2012

AP 3GM/60ML (50MG/ML)

A201689 002 Nov 26, 2012

AP ! TEVA PHARMS USA 1GM/20ML (50MG/ML)

A076657 001 Apr 04, 2007

AP ! 3GM/60ML (50MG/ML)

A076657 002 Apr 04, 2007

AP WEST-WARD PHARMS INT 1GM/20ML (50MG/ML)

A076619 001 Jun 29, 2011

AP 3GM/60ML (50MG/ML)

A076619 002 Jun 29, 2011

ILOPERIDONE

TABLET; ORAL

FANAPT

AB +! VANDA PHARMS INC 1MG

N022192 001 May 06, 2009

AB + 2MG

N022192 002 May 06, 2009

AB + 4MG

N022192 003 May 06, 2009

AB + 6MG

N022192 004 May 06, 2009

AB + 8MG

N022192 005 May 06, 2009

AB + 10MG

N022192 006 May 06, 2009

AB + 12MG

N022192 007 May 06, 2009

ILOPERIDONE

AB INVENTIA HLTHCARE 1MG

A207231 001 Nov 28, 2016

AB 2MG

A207231 002 Nov 28, 2016

AB 4MG

A207231 003 Nov 28, 2016

AB 6MG

A207231 004 Nov 28, 2016

AB 8MG

A207231 005 Nov 28, 2016

AB 10MG

A207231 006 Nov 28, 2016

AB 12MG

A207231 007 Nov 28, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-226 (of 436)

ILOPROST

SOLUTION; INHALATION

VENTAVIS

+!	ACTELION PHARMS LTD	10MCG/ML (10MCG/ML)
+!		20MCG/ML (20MCG/ML)

N021779 002 Dec 08, 2005  
N021779 003 Aug 07, 2009

IMATINIB MESYLATE

TABLET; ORAL

**GLEEVEC**

**AB** + NOVARTIS

**EQ 100MG BASE**

**N021588 001** Apr 18, 2003

**AB** +!

**EQ 400MG BASE**

**N021588 002** Apr 18, 2003

**IMATINIB MESYLATE**

**AB** APOTEX INC

**EQ 100MG BASE**

**A079179 001** Aug 05, 2016

**AB**

**EQ 400MG BASE**

**A079179 002** Aug 05, 2016

**AB** MYLAN PHARMS INC

**EQ 100MG BASE**

**A204644 001** Jun 21, 2017

**AB**

**EQ 400MG BASE**

**A204644 002** Jun 21, 2017

**AB** SUN PHARMA GLOBAL

**EQ 100MG BASE**

**A078340 001** Dec 03, 2015

**AB**

**EQ 400MG BASE**

**A078340 002** Dec 03, 2015

**AB** TEVA PHARMS USA

**EQ 100MG BASE**

**A204285 001** Aug 04, 2016

**AB**

**EQ 400MG BASE**

**A204285 002** Aug 04, 2016

IMIGLUCERASE

INJECTABLE; INJECTION

CEREZYME

+ GENZYME

200 UNITS/VIAL

N020367 001 May 23, 1994

+!

400 UNITS/VIAL

N020367 002 Sep 22, 1999

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

**IMIPRAMINE HYDROCHLORIDE**

**AB** LEADING PHARMA LLC

**10MG**

**A040903 001** Oct 24, 2012

**AB**

**25MG**

**A040903 002** Oct 24, 2012

**AB**

**50MG**

**A040903 003** Oct 24, 2012

**AB** LUPIN LTD

**10MG**

**A090441 002** Mar 11, 2010

**AB**

**25MG**

**A090441 003** Mar 11, 2010

**AB**

**50MG**

**A090441 001** Mar 11, 2010

**AB** PAR PHARM

**10MG**

**A088292 001** Oct 21, 1983

**AB**

**25MG**

**A088262 001** Oct 21, 1983

**AB**

**50MG**

**A088276 001** Oct 21, 1983

**AB** SANDOZ

**10MG**

**A084936 002**

**AB**

**25MG**

**A083745 001**

**AB**

**50MG**

**A084937 001**

**AB** SPECGX LLC

**10MG**

**A087846 002** May 22, 1984

**AB**

**25MG**

**A087846 003** May 22, 1984

**AB** SUN PHARM INDUSTRIES

**10MG**

**A081048 001** Jun 05, 1990

**AB**

**25MG**

**A081049 001** Jun 05, 1990

**AB**

**50MG**

**A081050 001** Jun 05, 1990

**TOFRANIL**

**AB** ! SPECGX LLC

**50MG**

**A087846 001** May 22, 1984

IMIPRAMINE HYDROCHLORIDE

OXFORD PHARMS

10MG

A040751 003 Feb 28, 2008

25MG

A040751 002 Feb 28, 2008

IMIPRAMINE PAMOATE

CAPSULE; ORAL

**IMIPRAMINE PAMOATE**

**AB** LUPIN LTD

**EQ 75MG HCL**

**A090444 001** Apr 16, 2010

**AB**

**EQ 100MG HCL**

**A090444 002** Apr 16, 2010

**AB**

**EQ 125MG HCL**

**A090444 003** Apr 16, 2010

**AB**

**EQ 150MG HCL**

**A090444 004** Apr 16, 2010

**AB** ! WEST-WARD PHARMS INT

**EQ 75MG HCL**

**A091099 001** Apr 16, 2010

**AB**

**EQ 100MG HCL**

**A091099 002** Apr 16, 2010

**AB**

**EQ 125MG HCL**

**A091099 003** Apr 16, 2010

**AB**

**EQ 150MG HCL**

**A091099 004** Apr 16, 2010

IMIQUIMOD

CREAM; TOPICAL

**ALDARA**

**AB** +! MEDICIS

**5%**

**N020723 001** Feb 27, 1997

**IMIQUIMOD**

**AB** APOTEX INC

**5%**

**A091308 001** Apr 06, 2012

**AB** FOUGERA PHARMS

**5%**

**A078548 001** Feb 25, 2010

**AB** G AND W LABS INC

**5%**

**A200481 001** Apr 18, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-227 (of 436)

IMIQUIMOD

CREAM;TOPICAL

IMIQUIMOD

<u>AB</u>	GLENMARK GENERICS	<u>5%</u>	<u>A201994 001</u>	Mar 06, 2012
<u>AB</u>	PERRIGO ISRAEL	<u>5%</u>	<u>A078837 001</u>	Sep 07, 2010
<u>AB</u>	STRIDES PHARMA	<u>5%</u>	<u>A202002 001</u>	Jun 24, 2014
<u>AB</u>	TARO	<u>5%</u>	<u>A200173 001</u>	Apr 15, 2011
<u>AB</u>	TOLMAR	<u>5%</u>	<u>A091044 001</u>	Feb 28, 2011
ZYCLARA				
+!	MEDICIS	2.5%	N022483 002	Jul 15, 2011
+!		3.75%	N022483 001	Mar 25, 2010

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONE LACTATE

!	WEST-WARD PHARMS INT	EQ 5MG BASE/ML
---	----------------------	----------------

A075513 001 May 09, 2000

INDACATEROL MALEATE

POWDER; INHALATION

ARCAPTA NEOHALER

+!	SUNOVION PHARMS INC	EQ 75MCG BASE
----	---------------------	---------------

N022383 001 Jul 01, 2011

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>1.25MG</u>	<u>A074722 001</u>	Jun 17, 1996
<u>AB</u>		<u>2.5MG</u>	<u>A074722 002</u>	Jun 17, 1996
<u>AB</u>	AMERIGEN PHARMS LTD	<u>1.25MG</u>	<u>A075201 001</u>	Dec 04, 1998
<u>AB</u>		<u>2.5MG</u>	<u>A075201 002</u>	Dec 04, 1998
<u>AB</u>	ANI PHARMS INC	<u>1.25MG</u>	<u>A074299 002</u>	Apr 29, 1996
<u>AB</u>		<u>2.5MG</u>	<u>A074299 001</u>	Jul 27, 1995
<u>AB</u>	MYLAN	<u>1.25MG</u>	<u>A074461 002</u>	Mar 26, 1997
<u>AB</u> !		<u>2.5MG</u>	<u>A074461 001</u>	Mar 27, 1996

INDINAVIR SULFATE

CAPSULE; ORAL

CRIVIXAN

+	MERCK SHARP DOHME	EQ 200MG BASE
+		EQ 400MG BASE

N020685 003 Mar 13, 1996

N020685 001 Mar 13, 1996

INDIUM IN-111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+!	GE HEALTHCARE	2mCi/0.2ML
INDIUM IN 111 CHLORIDE		
+!	MALLINKRODT NUCLEAR	5mCi/0.5ML

N019862 001 Dec 29, 1992

N019841 001 Sep 27, 1994

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN 111 OXYQUINOLINE

+!	GE HEALTHCARE	1mCi/ML
----	---------------	---------

N019044 001 Dec 24, 1985

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

+!	GE HEALTHCARE	1mCi/ML
----	---------------	---------

N017707 001 Feb 18, 1982

INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

OCTREOSCAN

+!	MALLINKRODT NUCLEAR	3mCi/ML
----	---------------------	---------

N020314 001 Jun 02, 1994

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

<u>AP</u>	+!	AKORN	<u>25MG/VIAL</u>	<u>N011525 001</u>
-----------	----	-------	------------------	--------------------

INDOCYANINE GREEN

<u>AP</u>		DIAGNOSTIC GREEN	<u>25MG/VIAL</u>	<u>A040811 001</u>
-----------	--	------------------	------------------	--------------------

Nov 21, 2007

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

<u>AB</u>	GLENMARK GENERICS	<u>25MG</u>	<u>A091276 001</u>	Dec 22, 2010
<u>AB</u>		<u>50MG</u>	<u>A091276 002</u>	Dec 22, 2010
<u>AB</u>	HERITAGE PHARMS INC	<u>25MG</u>	<u>N018851 001</u>	May 18, 1984
<u>AB</u>		<u>50MG</u>	<u>N018851 002</u>	May 18, 1984
<u>AB</u>	HETERO LABS LTD III	<u>25MG</u>	<u>A091240 001</u>	Apr 12, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-228 (of 436)

INDOMETHACIN

CAPSULE;ORAL

INDOMETHACIN

<u>AB</u>		<u>50MG</u>	<u>A091240</u> <u>002</u>	Apr 12, 2011
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>25MG</u>	<u>A070719</u> <u>001</u>	Feb 12, 1986
<u>AB</u>		<u>50MG</u>	<u>A070756</u> <u>001</u>	Feb 12, 1986
<u>AB</u>	JUBILANT GENERICS	<u>25MG</u>	<u>A205215</u> <u>001</u>	Aug 25, 2017
<u>AB</u>		<u>50MG</u>	<u>A205215</u> <u>002</u>	Aug 25, 2017
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N018858</u> <u>001</u>	Apr 20, 1984
<u>AB</u>		<u>50MG</u>	<u>A070624</u> <u>001</u>	Sep 04, 1985
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A070673</u> <u>001</u>	Apr 29, 1987
<u>AB</u>		<u>50MG</u>	<u>A070674</u> <u>001</u>	Apr 29, 1987
<u>AB</u>	SUN PHARM INDs INC	<u>25MG</u>	<u>A091401</u> <u>001</u>	Mar 28, 2013
<u>AB</u>		<u>50MG</u>	<u>A091401</u> <u>002</u>	Mar 28, 2013
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A090403</u> <u>001</u>	Nov 15, 2010
<u>AB</u>		<u>50MG</u>	<u>A090403</u> <u>002</u>	Nov 15, 2010
TIVORBEX				
+ IROKO PHARMS LLC	20MG		N204768 001	Feb 24, 2014
+! !	40MG		N204768 002	Feb 24, 2014

CAPSULE, EXTENDED RELEASE;ORAL

INDOMETHACIN

<u>AB</u>	AMNEAL PHARMS	<u>75MG</u>	<u>A091549</u> <u>001</u>	Dec 01, 2010
<u>AB</u>	AUROBINDO PHARMA LTD	<u>75MG</u>	<u>A204243</u> <u>001</u>	Dec 27, 2016
<u>AB</u>	AVANTHI INC	<u>75MG</u>	<u>A079175</u> <u>001</u>	Mar 06, 2009
<u>AB</u>	CHARTWELL RX	<u>75MG</u>	<u>A200529</u> <u>001</u>	Nov 30, 2010
<u>AB</u>	GLENMARK PHARMS LTD	<u>75MG</u>	<u>A203501</u> <u>001</u>	Jun 22, 2017
<u>AB</u>	HETERO LABS LTD III	<u>75MG</u>	<u>A201807</u> <u>001</u>	Sep 28, 2012
<u>AB</u>	JUBILANT GENERICS	<u>75MG</u>	<u>A202706</u> <u>001</u>	Oct 05, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>75MG</u>	<u>A202139</u> <u>001</u>	Mar 20, 2014
<u>AB</u>	NOVAST LABS LTD	<u>75MG</u>	<u>A204853</u> <u>001</u>	May 08, 2017
<u>AB</u>	! SANDOZ	<u>75MG</u>	<u>A074464</u> <u>001</u>	May 28, 1998
<u>AB</u>	ZYDUS PHARMS USA INC	<u>75MG</u>	<u>A202711</u> <u>001</u>	Sep 25, 2017

INJECTABLE;INJECTION

INDOMETHACIN				
+! FRESENIUS KABI USA	EQ 1MG BASE/VIAL		N022536 001	Mar 17, 2010

SUPPOSITORY;RECTAL

INDOMETHACIN				
! G AND W LABS	50MG		A073314 001	Aug 31, 1992

SUSPENSION;ORAL

INDOCIN				
+! IROKO PHARMS	25MG/5ML		N018332 001	Oct 10, 1985

INDOMETHACIN SODIUM

INJECTABLE;INJECTION

<u>INDOCIN</u>				
<u>AP</u> +! RECORDATI RARE	<u>EQ 1MG BASE/VIAL</u>		<u>N018878</u> <u>001</u>	Jan 30, 1985

<u>INDOMETHACIN SODIUM</u>				
<u>AP</u>	HOSPIRA INC	<u>EQ 1MG BASE/VIAL</u>	<u>A204118</u> <u>001</u>	Apr 19, 2016
<u>AP</u>	NAVINTA LLC	<u>EQ 1MG BASE/VIAL</u>	<u>A206561</u> <u>001</u>	Jul 19, 2017
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 1MG BASE/VIAL</u>	<u>A078713</u> <u>001</u>	Jul 16, 2008

INGENOL MEQUITATE

GEL;TOPICAL

PICATO				
+! LEO LABS	0.015%		N202833 001	Jan 23, 2012
+!	0.05%		N202833 002	Jan 23, 2012

INSULIN ASPART

SOLUTION;IV (INFUSION), SUBCUTANEOUS  
 FIASP

+! NOVO NORDISK INC	1000 UNITS/10ML (100 UNITS/ML)		N208751 001	Sep 29, 2017
SOLUTION;SUBCUTANEOUS				

FIASP FLEXTOUCH				
+! NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)		N208751 002	Sep 29, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-229 (of 436)

INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE;SUBCUTANEOUS			
NOVOLOG MIX 70/30			
+! NOVO NORDISK INC	700 UNITS/10ML; 300 UNITS/10ML (70 UNITS/ML; 30 UNITS/ML)		N021172 001 Nov 01, 2001
NOVOLOG MIX 70/30 FLEXPEN			
+! NOVO NORDISK INC	210 UNITS/3ML; 90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML)		N021172 004 May 03, 2002

INSULIN ASPART RECOMBINANT

INJECTABLE;SUBCUTANEOUS			
NOVOLOG			
+! NOVO NORDISK INC	1000 UNITS/10ML (100 UNITS/ML)		N020986 001 Jun 07, 2000
NOVOLOG FLEXPEN			
+! NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)		N020986 003 Jan 19, 2001
NOVOLOG FLEXTOUCH			
+! NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)		N020986 005 Oct 31, 2013
NOVOLOG PENFILL			
+! NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)		N020986 002 Jun 07, 2000

INSULIN ASPART; INSULIN DEGLUDEC

SOLUTION;SUBCUTANEOUS			
RYZODEG 70/30			
+! NOVO NORDISK INC	90 UNITS/3ML; 210 UNITS/3ML (30 UNITS/ML; 70 UNITS/ML)		N203313 001 Sep 25, 2015

INSULIN DEGLUDEC

SOLUTION;SUBCUTANEOUS			
TRESIBA			
+! NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)		N203314 001 Sep 25, 2015
+!	600 UNITS/3ML (200 UNITS/ML)		N203314 002 Sep 25, 2015

INSULIN DEGLUDEC; LIRAGLUTIDE

SOLUTION;SUBCUTANEOUS			
XULTOPHY 100/3.6			
+! NOVO NORDISK INC	300 UNITS/3ML; 10.8MG/3ML (100 UNITS/ML; 3.6MG/ML)		N208583 001 Nov 21, 2016

INSULIN DETEMIR RECOMBINANT

INJECTABLE;SUBCUTANEOUS			
LEVEMIR			
+! NOVO NORDISK INC	1000 UNITS/10ML (100 UNITS/ML)		N021536 001 Jun 16, 2005
LEVEMIR FLEXTOUCH			
+! NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)		N021536 005 Oct 31, 2013

INSULIN GLARGINE

SOLUTION;SUBCUTANEOUS			
BASAGLAR			
ELI LILLY AND CO	300 UNITS/3ML (100 UNITS/ML)		N205692 001 Dec 16, 2015

INSULIN GLARGINE RECOMBINANT

INJECTABLE;INJECTION			
LANTUS			
+! SANOFI AVENTIS US	100 UNITS/ML		N021081 001 Apr 20, 2000
LANTUS SOLOSTAR			
+! SANOFI AVENTIS US	300 UNITS/3ML (100 UNITS/ML)		N021081 002 Apr 27, 2007
SOLUTION;SUBCUTANEOUS			
TOUJEO SOLOSTAR			
+! SANOFI US SERVICES	300 UNITS/ML (300 UNITS/ML)		N206538 001 Feb 25, 2015

INSULIN GLARGINE; LIXISENATIDE

SOLUTION;SUBCUTANEOUS			
SOLIQUA 100/33			
+! SANOFI-AVENTIS US	300 UNITS/3ML; 99MCG/3ML (100 UNITS/ML; 33MCG/ML)		N208673 001 Nov 21, 2016

INSULIN GLULISINE RECOMBINANT

INJECTABLE;IV (INFUSION), SUBCUTANEOUS			
APIDRA			
+! SANOFI AVENTIS US	1000 UNITS/10ML (100 UNITS/ML)		N021629 001 Apr 16, 2004
+!	300 UNITS/3ML (100 UNITS/ML)		N021629 002 Dec 20, 2005

INJECTABLE;SUBCUTANEOUS			
APIDRA SOLOSTAR			
+ SANOFI AVENTIS US	300 UNITS/3ML		N021629 003 Feb 24, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-230 (of 436)

INSULIN HUMAN

SOLUTION;SUBCUTANEOUS

HUMULIN R			
+! LILLY	10000 UNITS/20ML (500 UNITS/ML)	N018780 004	Mar 31, 1994
HUMULIN R KWIKPEN			
+! LILLY	1500 UNITS/3ML (500 UNITS/ML)	N018780 002	Dec 29, 2015

INSULIN LISPRO

SOLUTION;IV (INFUSION), SUBCUTANEOUS

ADMELOG			
+ SANOFI-AVENTIS US	1000 UNITS/10ML (100 UNITS/ML)	N209196 001	Dec 11, 2017
ADMELOG SOLOSTAR			
+ SANOFI-AVENTIS US	300 UNITS/3ML (100 UNITS/ML)	N209196 002	Dec 11, 2017

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG MIX 50/50			
+! LILLY	50 UNITS/ML;50 UNITS/ML	N021018 001	Dec 22, 1999
HUMALOG MIX 50/50 KWIKPEN			
+! LILLY	50 UNITS/ML;50 UNITS/ML	N021018 002	Sep 06, 2007
HUMALOG MIX 75/25			
+! LILLY	75 UNITS/ML;25 UNITS/ML	N021017 001	Dec 22, 1999
HUMALOG MIX 75/25 KWIKPEN			
+! LILLY	75 UNITS/ML;25 UNITS/ML	N021017 002	Sep 06, 2007

INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION

HUMALOG			
+! LILLY	100 UNITS/ML	N020563 001	Jun 14, 1996
HUMALOG KWIKPEN			
+! LILLY	100 UNITS/ML	N020563 003	Sep 06, 2007
SOLUTION;SUBCUTANEOUS			
HUMALOG KWIKPEN			
+! ELI LILLY AND CO	200 UNITS/ML	N205747 001	May 26, 2015

INSULIN RECOMBINANT HUMAN

POWDER;INHALATION

AFREZZA			
+ MANNKIND	4 UNITS/INH	N022472 001	Jun 27, 2014
+!	8 UNITS/INH	N022472 002	Jun 27, 2014
+	12 UNITS/INH	N022472 003	Apr 17, 2015

IOBENGUANE SULFATE I-123

SOLUTION;INTRAVENOUS

ADREVIEW			
+! GE HEALTHCARE	10mCi/5ML (2mCi/ML)	N022290 001	Sep 19, 2008

IODIXANOL

INJECTABLE;INJECTION

VISIPAQUE 270			
+! GE HEALTHCARE	55%	N020351 001	Mar 22, 1996
VISIPAQUE 320			
+! GE HEALTHCARE	65.2%	N020351 002	Mar 22, 1996
	65.2%	N020808 002	Aug 29, 1997

IOFLUPANE I-123

SOLUTION;INTRAVENOUS

DATSCAN			
+! GE HLTHCARE INC	5mCi/2.5ML (2mCi/ML)	N022454 001	Jan 14, 2011

TOHEXOL

FOR SOLUTION;ORAL

ORALTAG			
INTERPHARMA PRAHA	9.7GM/BOT	N205383 001	Mar 26, 2015
AS			

INJECTABLE;INJECTION

OMNIPAQ 140			
+! GE HEALTHCARE	30.2%	N018956 005	Nov 30, 1988

SOLUTION;INJECTION, ORAL

OMNIPAQ 350			
+! GE HEALTHCARE	75.5%	N018956 004	Dec 26, 1985
	75.5%	N020608 003	Oct 24, 1995

SOLUTION;INJECTION, ORAL, RECTAL

OMNIPAQ 180			
+! GE HEALTHCARE	38.8%	N018956 001	Dec 26, 1985

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-231 (of 436)

IOTHEXOL

SOLUTION; INJECTION, ORAL, RECTAL	
OMNIPAQUE 240	
+! GE HEALTHCARE	51.8%
	51.8%
OMNIPAQUE 300	
+! GE HEALTHCARE	64.7%
	64.7%

N018956 002	Dec 26, 1985
N020608 001	Oct 24, 1995
N018956 003	Dec 26, 1985
N020608 002	Oct 24, 1995

IOPAMIDOL

INJECTABLE; INJECTION

<u>ISOVUE-300</u>	
<b>AP</b> +! BRACCO	<b>61%</b>
<u>ISOVUE-370</u>	
<b>AP</b> +! BRACCO	<b>76%</b>
<u>SCANLUX-300</u>	
<b>AP</b> SANOCHEMIA CORP USA	<b>61%</b>
<u>SCANLUX-370</u>	
<b>AP</b> SANOCHEMIA CORP USA	<b>76%</b>
ISOVUE-200	
+! BRACCO	41%
ISOVUE-250	
+! BRACCO	51%
+!	51%
ISOVUE-300	
+! BRACCO	61%
ISOVUE-370	
+! BRACCO	76%
ISOVUE-M 200	
+! BRACCO	41%
ISOVUE-M 300	
+! BRACCO	61%

<b>N018735 002</b>	Dec 31, 1985
<b>N018735 003</b>	Dec 31, 1985
<b>A090394 001</b>	Jun 18, 2010
<b>A090394 002</b>	Jun 18, 2010
N018735 006	Jul 07, 1987
N018735 007	Jul 06, 1992
N020327 002	Oct 12, 1994
N020327 003	Oct 12, 1994
N020327 004	Oct 12, 1994
N018735 001	Dec 31, 1985
N018735 004	Dec 31, 1985

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)	
+! BAYER HLTHCARE	49.9%
+!	62.3%
+!	76.9%
ULTRAVIST 240	
+! BAYER HLTHCARE	49.9%
ULTRAVIST 300	
+! BAYER HLTHCARE	62.3%
ULTRAVIST 370	
+! BAYER HLTHCARE	76.9%

N021425 003	Mar 12, 2004
N021425 001	Sep 20, 2002
N021425 002	Sep 20, 2002
N020220 003	May 10, 1995
N020220 002	May 10, 1995
N020220 001	May 10, 1995

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY	
+! LIEBEL-FLARSHEIM	60%
CONRAY 30	
+! LIEBEL-FLARSHEIM	30%
CONRAY 43	
+! LIEBEL-FLARSHEIM	43%
SOLUTION; INTRAVESICAL	
CYSTO-CONRAY II	
LIEBEL-FLARSHEIM	17.2%

N013295 001	
N016983 001	
N013295 002	
N017057 002	

IOTHALAMATE SODIUM I-125

INJECTABLE; INJECTION

GLOFIL-125	
ISOTEX	250-300uCi/ML

N017279 001	
-------------	--

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 240	
+! LIEBEL-FLARSHEIM	51%
OPTIRAY 300	
+! LIEBEL-FLARSHEIM	64%
+!	64%
OPTIRAY 320	
+! LIEBEL-FLARSHEIM	68%
+!	68%

N019710 002	Dec 30, 1988
N019710 004	Jan 22, 1992
N020923 004	May 13, 1999
N019710 001	Dec 30, 1988
N020923 002	May 29, 1998

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-232 (of 436)

TOVERSOL

INJECTABLE; INJECTION

OPTIRAY 350

+!	LIEBEL-FLARSHEIM	74%
+!		74%

N019710	005	Jan 22, 1992
N020923	003	May 28, 1998

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT HFA

+!	BOEHRINGER	0.021MG/INH
	INGELHEIM	

N021527 001 Nov 27, 2004

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

<u>AN</u>	AUROBINDO PHARMA LTD	<u>0.02%</u>
<u>AN</u>	LANDELA PHARM	<u>0.02%</u>
<u>AN</u>	NEPHRON	<u>0.02%</u>
<u>AN</u>	RITEDOSE CORP	<u>0.02%</u>
<u>AN</u>	SUN PHARMA GLOBAL	<u>0.02%</u>
<u>AN</u>	WATSON LABS	<u>0.02%</u>

<u>A206543</u>	<u>001</u>	Oct 27, 2016
<u>A077072</u>	<u>001</u>	Jul 19, 2005
<u>A075562</u>	<u>001</u>	Sep 27, 2001
<u>A075693</u>	<u>001</u>	Jan 26, 2001
<u>A207903</u>	<u>001</u>	Jan 03, 2017
<u>A076291</u>	<u>001</u>	May 09, 2005

SPRAY, METERED; NASAL

ATROVENT

<u>AB</u>	+!	BOEHRINGER	<u>0.021MG/SPRAY</u>
		INGELHEIM	

<u>N020393</u>	<u>001</u>	Oct 20, 1995
<u>N020394</u>	<u>001</u>	Oct 20, 1995

IPRATROPIUM BROMIDE

<u>AB</u>	BAUSCH AND LOMB	<u>0.021MG/SPRAY</u>
<u>AB</u>		<u>0.042MG/SPRAY</u>
<u>AB</u>	MYLAN SPECIALITY LP	<u>0.021MG/SPRAY</u>
<u>AB</u>		<u>0.042MG/SPRAY</u>
<u>AB</u>	WEST-WARD PHARMS INT	<u>0.021MG/SPRAY</u>
<u>AB</u>		<u>0.042MG/SPRAY</u>

<u>A076025</u>	<u>001</u>	Mar 31, 2003
<u>A076103</u>	<u>001</u>	Mar 31, 2003
<u>A075552</u>	<u>001</u>	Mar 31, 2003
<u>A075553</u>	<u>001</u>	Mar 31, 2003
<u>A076664</u>	<u>001</u>	Nov 05, 2003
<u>A076598</u>	<u>001</u>	Nov 05, 2003

IRBESARTAN

TABLET; ORAL

AVAPRO

<u>AB</u>	+	SANOFI AVENTIS US	<u>75MG</u>
<u>AB</u>	+		<u>150MG</u>
<u>AB</u>	+!		<u>300MG</u>

<u>N020757</u>	<u>001</u>	Sep 30, 1997
<u>N020757</u>	<u>002</u>	Sep 30, 1997
<u>N020757</u>	<u>003</u>	Sep 30, 1997

IRBESARTAN

<u>AB</u>	ALEMBIC PHARMS LTD	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>
<u>AB</u>	APOTEX INC	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>
<u>AB</u>	AUROBINDO PHARMA LTD	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>

<u>A091236</u>	<u>001</u>	Oct 15, 2012
<u>A091236</u>	<u>002</u>	Oct 15, 2012
<u>A091236</u>	<u>003</u>	Oct 15, 2012
<u>A200832</u>	<u>001</u>	Oct 15, 2012
<u>A200832</u>	<u>002</u>	Oct 15, 2012
<u>A200832</u>	<u>003</u>	Oct 15, 2012
<u>A203081</u>	<u>001</u>	Sep 27, 2012
<u>A203081</u>	<u>002</u>	Sep 27, 2012
<u>A203081</u>	<u>003</u>	Sep 27, 2012

<u>AB</u>	CIPLA LTD	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>
<u>AB</u>	DR REDDYS LABS LTD	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>

<u>A077205</u>	<u>001</u>	Nov 14, 2012
<u>A077205</u>	<u>002</u>	Nov 14, 2012
<u>A077205</u>	<u>003</u>	Nov 14, 2012
<u>A203161</u>	<u>001</u>	Sep 27, 2012
<u>A203161</u>	<u>002</u>	Sep 27, 2012
<u>A203161</u>	<u>003</u>	Sep 27, 2012

<u>AB</u>	HETERO LABS LTD V	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>
<u>AB</u>	HISUN PHARM HANGZHOU	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>

<u>A202910</u>	<u>001</u>	Sep 27, 2012
<u>A202910</u>	<u>002</u>	Sep 27, 2012
<u>A202910</u>	<u>003</u>	Sep 27, 2012
<u>A206194</u>	<u>001</u>	Jun 14, 2016
<u>A206194</u>	<u>002</u>	Jun 14, 2016
<u>A206194</u>	<u>003</u>	Jun 14, 2016

<u>AB</u>	JUBILANT GENERICS	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>
<u>AB</u>	LUPIN LTD	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>

<u>A203534</u>	<u>001</u>	Feb 23, 2015
<u>A203534</u>	<u>002</u>	Feb 23, 2015
<u>A203534</u>	<u>003</u>	Feb 23, 2015
<u>A201531</u>	<u>001</u>	Oct 15, 2012
<u>A201531</u>	<u>002</u>	Oct 15, 2012
<u>A201531</u>	<u>003</u>	Oct 15, 2012

<u>AB</u>	MACLEODS PHARMS LTD	<u>75MG</u>
<u>AB</u>		<u>150MG</u>
<u>AB</u>		<u>300MG</u>
<u>AB</u>	PRINSTON INC	<u>75MG</u>
<u>AB</u>		<u>150MG</u>

<u>A202254</u>	<u>001</u>	Oct 03, 2012
<u>A202254</u>	<u>002</u>	Oct 03, 2012
<u>A202254</u>	<u>003</u>	Oct 03, 2012
<u>A203071</u>	<u>001</u>	Sep 27, 2012
<u>A203071</u>	<u>002</u>	Sep 27, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-233 (of 436)

IRBESARTAN

TABLET; ORAL

IRBESARTAN

<u>AB</u>		<u>300MG</u>	<u>A203071</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>	SANDOZ	<u>75MG</u>	<u>A077466</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A077466</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A077466</u>	<u>003</u>	Sep 27, 2012
<u>AB</u>	SCIEGEN PHARMS INC	<u>75MG</u>	<u>A204774</u>	<u>001</u>	Dec 07, 2015
<u>AB</u>		<u>150MG</u>	<u>A204774</u>	<u>002</u>	Dec 07, 2015
<u>AB</u>		<u>300MG</u>	<u>A204774</u>	<u>003</u>	Dec 07, 2015
<u>AB</u>	TEVA PHARMS	<u>75MG</u>	<u>A077159</u>	<u>001</u>	Mar 30, 2012
<u>AB</u>		<u>150MG</u>	<u>A077159</u>	<u>002</u>	Mar 30, 2012
<u>AB</u>		<u>300MG</u>	<u>A077159</u>	<u>003</u>	Mar 30, 2012
<u>AB</u>	UNICHEM LABS LTD	<u>75MG</u>	<u>A203020</u>	<u>001</u>	Dec 07, 2015
<u>AB</u>		<u>150MG</u>	<u>A203020</u>	<u>002</u>	Dec 07, 2015
<u>AB</u>		<u>300MG</u>	<u>A203020</u>	<u>003</u>	Dec 07, 2015
<u>AB</u>	WEST-WARD PHARMS INT	<u>75MG</u>	<u>A090201</u>	<u>001</u>	Oct 15, 2012
<u>AB</u>		<u>150MG</u>	<u>A090201</u>	<u>002</u>	Oct 15, 2012
<u>AB</u>		<u>300MG</u>	<u>A090201</u>	<u>003</u>	Oct 15, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>75MG</u>	<u>A079213</u>	<u>001</u>	Sep 27, 2012
<u>AB</u>		<u>150MG</u>	<u>A079213</u>	<u>002</u>	Sep 27, 2012
<u>AB</u>		<u>300MG</u>	<u>A079213</u>	<u>003</u>	Sep 27, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

<u>AP</u>	+!	PFIZER INC	<u>40MG/2ML (20MG/ML)</u>	<u>N020571</u>	<u>001</u>	Jun 14, 1996
<u>AP</u>	+!		<u>100MG/5ML (20MG/ML)</u>	<u>N020571</u>	<u>002</u>	Jun 14, 1996
<u>AP</u>	+!		<u>300MG/15ML (20MG/ML)</u>	<u>N020571</u>	<u>003</u>	Aug 05, 2010

IRINOTECAN HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>40MG/2ML (20MG/ML)</u>	<u>A079068</u>	<u>001</u>	Nov 21, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A079068</u>	<u>002</u>	Nov 21, 2008
<u>AP</u>	ACTAVIS TOTOWA	<u>40MG/2ML (20MG/ML)</u>	<u>A078589</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078589</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>		<u>500MG/25ML (20MG/ML)</u>	<u>A078589</u>	<u>003</u>	Nov 18, 2015
<u>AP</u>	AKORN	<u>40MG/2ML (20MG/ML)</u>	<u>A090726</u>	<u>001</u>	Sep 16, 2009
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090726</u>	<u>002</u>	Sep 16, 2009
<u>AP</u>	CIPLA LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A077219</u>	<u>001</u>	Feb 20, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077219</u>	<u>002</u>	Feb 20, 2008
<u>AP</u>	DR REDDYS LABS LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A078953</u>	<u>001</u>	Apr 15, 2010
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078953</u>	<u>002</u>	Apr 15, 2010
<u>AP</u>	EMCURE PHARMS LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A200771</u>	<u>001</u>	Feb 14, 2012
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A200771</u>	<u>002</u>	Feb 14, 2012
<u>AP</u>	FRESENIUS KABI ONCOL	<u>40MG/2ML (20MG/ML)</u>	<u>A078188</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078188</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>	FRESENIUS KABI USA	<u>40MG/2ML (20MG/ML)</u>	<u>A077776</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077776</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>	HIKMA FARMACEUTICA	<u>40MG/2ML (20MG/ML)</u>	<u>A091032</u>	<u>001</u>	Dec 20, 2010
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A091032</u>	<u>002</u>	Dec 20, 2010
<u>AP</u>	HISUN PHARM HANGZHOU	<u>40MG/2ML (20MG/ML)</u>	<u>A090016</u>	<u>001</u>	Jan 28, 2009
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090016</u>	<u>002</u>	Jan 28, 2009
<u>AP</u>	HOSPIRA	<u>40MG/2ML (20MG/ML)</u>	<u>A077915</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A077915</u>	<u>002</u>	Feb 27, 2008
<u>AP</u>	!	<u>500MG/25ML (20MG/ML)</u>	<u>A078796</u>	<u>001</u>	Feb 27, 2008
<u>AP</u>	INGENUS PHARMS LLC	<u>40MG/2ML (20MG/ML)</u>	<u>A206935</u>	<u>001</u>	May 26, 2017
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A206935</u>	<u>002</u>	May 26, 2017
<u>AP</u>	INTAS PHARMS USA	<u>40MG/2ML (20MG/ML)</u>	<u>A203054</u>	<u>001</u>	Aug 31, 2017
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A203054</u>	<u>002</u>	Aug 31, 2017
<u>AP</u>	JIANGSU HENGRI MED	<u>40MG/2ML (20MG/ML)</u>	<u>A090675</u>	<u>002</u>	Dec 16, 2011
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090675</u>	<u>001</u>	Dec 16, 2011
<u>AP</u>	MUSTAFA NEVZAT ILAC	<u>40MG/2ML (20MG/ML)</u>	<u>A090393</u>	<u>002</u>	May 13, 2011
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A090393</u>	<u>003</u>	May 13, 2011
<u>AP</u>	PLIVA LACHEMA	<u>40MG/2ML (20MG/ML)</u>	<u>A078122</u>	<u>001</u>	Oct 31, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078122</u>	<u>002</u>	Oct 31, 2008
<u>AP</u>	QILU PHARM CO LTD	<u>40MG/2ML (20MG/ML)</u>	<u>A203380</u>	<u>001</u>	May 03, 2016
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A203380</u>	<u>002</u>	May 03, 2016
<u>AP</u>		<u>300MG/15ML (20MG/ML)</u>	<u>A203380</u>	<u>003</u>	May 03, 2016
<u>AP</u>	SUN PHARMA GLOBAL	<u>40MG/2ML (20MG/ML)</u>	<u>A078805</u>	<u>001</u>	Apr 21, 2008
<u>AP</u>		<u>100MG/5ML (20MG/ML)</u>	<u>A078805</u>	<u>002</u>	Apr 21, 2008
<u>AP</u>	TEVA PHARMS USA	<u>40MG/2ML (20MG/ML)</u>	<u>A090101</u>	<u>002</u>	Feb 27, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-234 (of 436)

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

<u>AP</u>	<u>100MG/5ML (20MG/ML)</u>	<u>A090101 003</u>	Feb 27, 2008
<u>AP</u>	<u>500MG/25ML (20MG/ML)</u>	<u>A090101 001</u>	Nov 26, 2008
<u>AP</u>	<u>WEST-WARD PHARMS INT</u>	<u>A078753 001</u>	Dec 24, 2008
<u>AP</u>	<u>100MG/5ML (20MG/ML)</u>	<u>A078753 002</u>	Dec 24, 2008
	INJECTABLE, LIPOSOMAL; IV (INFUSION) ONIVYDE +! IPSEN INC	EQ 43MG BASE/10ML (EQ 4.3MG BASE/ML)	N207793 001 Oct 22, 2015

IRON DEXTRAN

INJECTABLE; INJECTION  
DEXFERRUM

BP	LUITPOLD INFED	EQ 50MG IRON/ML	N040024 001 Feb 23, 1996
BP	+! ALLERGAN SALES LLC PROFERDEX	EQ 50MG IRON/ML	N017441 001
BP	NEW RIVER	EQ 50MG IRON/ML	N017807 001

IRON SUCROSE

INJECTABLE; INTRAVENOUS  
VENOFER

+ LUITPOLD	EQ 50MG BASE/2.5ML (EQ 20MG BASE/ML)	N021135 002 Mar 20, 2005
+!	EQ 100MG BASE/5ML (EQ 20MG BASE/ML)	N021135 001 Nov 06, 2000
+	EQ 200MG BASE/10ML (EQ 20MG BASE/ML)	N021135 004 Feb 09, 2007

ISAVUCONAZONIUM SULFATE

CAPSULE; ORAL  
CRESEMBA

+! ASTELLAS	186MG	N207500 001 Mar 06, 2015
CRESEMBA	372MG	N207501 001 Mar 06, 2015

ISOCARBOXAZID

TABLET; ORAL  
MARPLAN

+! VALIDUS PHARMS INC	10MG	N011961 001
-----------------------	------	-------------

ISOFLURANE

LIQUID; INHALATION

FORANE

<u>AN</u> +! BAXTER HLTHCARE	<u>99.9%</u>	<u>N017624 001</u>
<u>AN</u> HALOCARBON PRODS	<u>99.9%</u>	<u>A075225 001</u> Oct 20, 1999
<u>AN</u> HOSPIRA	<u>99.9%</u>	<u>A074097 001</u> Jan 25, 1993
<u>AN</u> PIRAMAL CRITICAL	<u>99.9%</u>	<u>A074416 001</u> Sep 30, 1994
<u>AN</u> PIRAMAL ENT	<u>99.9%</u>	<u>A074502 001</u> Jun 27, 1995

ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

! SANDOZ INC	100MG/ML	A040648 001 Jul 05, 2005
SYRUP; ORAL ISONIAZID	50MG/5ML	A088235 001 Nov 10, 1983

TABLET; ORAL

ISONIAZID

<u>AA</u> BARR	<u>100MG</u>	<u>A080936 001</u>
	<u>300MG</u>	<u>A080937 002</u>
<u>AA</u> MIKART	<u>100MG</u>	<u>A040090 001</u> Jun 26, 1997
	<u>300MG</u>	<u>A040090 002</u> Jun 26, 1997
<u>AA</u> +! SANDOZ	<u>100MG</u>	<u>N008678 002</u>
<u>AA</u> +!	<u>300MG</u>	<u>N008678 003</u>
<u>AA</u> THEPHARMANETWORK LLC	<u>100MG</u>	<u>A202610 001</u> Oct 29, 2014
	<u>300MG</u>	<u>A202610 002</u> Oct 29, 2014
<u>AA</u> LANNEOTT	<u>300MG</u>	<u>A089776 001</u> Jun 13, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-235 (of 436)

ISONIAZID; PYRAZINAMIDE; RIFAMPIN

TABLET;ORAL

RIFATER

+! SANOFI AVENTIS US 50MG;300MG;120MG

N050705 001 May 31, 1994

ISONIAZID; RIFAMPIN

CAPSULE;ORAL

RIFAMATE

! SANOFI AVENTIS US 150MG;300MG

A061884 001

ISOPROTERENOL HYDROCHLORIDE

INJECTABLE;INJECTION

ISOPROTERENOL HYDROCHLORIDE

AP NEXUS PHARMS 0.2MG/ML

A206961 001 Aug 02, 2017

ISUPREL

AP +! HOSPIRA 0.2MG/ML

N010515 001

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE;ORAL

DILATRATE-SR

+! AUXILIUM PHARMS LLC 40MG

N019790 001 Sep 02, 1988

TABLET;ORAL

ISORDIL

AB + VALEANT PHARMS 5MG

N012093 007 Jul 29, 1988

NORTH

ISOSORBIDE DINITRATE

AB HIKMA INTL PHARMS 5MG

A086067 001 Oct 29, 1987

AB 10MG

A086066 001 Oct 29, 1987

AB 20MG

A088088 001 Nov 02, 1987

AB 30MG

A040591 001 Jan 10, 2007

AB PAR PHARM 5MG

A086923 001 Mar 12, 1987

AB 10MG

A086925 001 Mar 12, 1987

AB 20MG

A087537 001 Oct 02, 1987

AB ! 30MG

A087946 001 Jan 12, 1988

AB SANDOZ 5MG

A086221 001 Jan 07, 1988

AB 10MG

A086223 001 Jan 07, 1988

AB 20MG

A089367 001 Apr 07, 1988

ISORDIL

+! VALEANT PHARMS 40MG

N012093 001 Jul 29, 1988

NORTH

TABLET, EXTENDED RELEASE;ORAL

ISOSORBIDE DINITRATE

! SUN PHARM INDs INC 40MG

A040009 001 Dec 30, 1998

ISOSORBIDE MONONITRATE

TABLET;ORAL

ISOSORBIDE MONONITRATE

AB ACTAVIS ELIZABETH 10MG

A075037 002 Oct 30, 1998

AB 20MG

A075037 001 Oct 30, 1998

AB ANI PHARMS INC 20MG

A075147 001 Nov 27, 1998

AB HIKMA PHARMS 20MG

A075361 001 Oct 05, 2000

MONOKET

AB + LANNETT CO INC 10MG

N020215 002 Jun 30, 1993

AB +! 20MG

N020215 001 Jun 30, 1993

TABLET, EXTENDED RELEASE;ORAL

ISOSORBIDE MONONITRATE

AB ACCORD HLTHCARE 30MG

A209684 001 Oct 24, 2017

AB 60MG

A209684 002 Oct 24, 2017

AB 120MG

A209684 003 Oct 24, 2017

AB DEXCEL LTD 30MG

A075522 002 Sep 20, 2016

AB 60MG

A075522 001 Apr 17, 2000

AB HIKMA PHARMS 30MG

A076813 002 Mar 30, 2006

AB 60MG

A076813 001 Jan 07, 2005

AB KREMERS URBAN 30MG

A075155 002 Jan 13, 2000

AB 60MG

A075155 001 Oct 30, 1998

AB ! 120MG

A075155 003 Aug 04, 2000

AB NESHER PHARMS 30MG

A075395 001 Mar 16, 2000

AB 60MG

A075395 002 Mar 16, 2000

AB 120MG

A075395 003 Mar 16, 2000

AB TORRENT PHARMS 30MG

A200270 001 Jun 03, 2011

AB 60MG

A200495 001 Jun 03, 2011

AB 120MG

A200495 002 Jun 03, 2011

AB VINTAGE PHARMS 30MG

A090598 001 Aug 11, 2010

AB 60MG

A090598 002 Aug 11, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-236 (of 436)

ISOSORBIDE MONONITRATE

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE MONONITRATE

<u>AB</u>	<u>120MG</u>	<u>A090598 003</u> Aug 11, 2010
-----------	--------------	---------------------------------

ISOSULFAN BLUE

INJECTABLE; INJECTION

ISOSULFAN BLUE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>1%</u>	<u>A206831 001</u> Feb 02, 2016
<u>AP</u>	MYLAN INSTITUTIONAL	<u>1%</u>	<u>A090874 001</u> Jul 20, 2010

ISOTRETINOIN

CAPSULE; ORAL

AMNESTEEM

<u>AB</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A075945 001</u> Nov 08, 2002
<u>AB</u>		<u>20MG</u>	<u>A075945 002</u> Nov 08, 2002
<u>AB</u>		<u>40MG</u>	<u>A075945 003</u> Nov 08, 2002

CLARAVIS

<u>AB</u>	TEVA PHARMS USA	<u>10MG</u>	<u>A076356 001</u> Apr 11, 2003
<u>AB</u>		<u>20MG</u>	<u>A076135 002</u> Apr 11, 2003
<u>AB</u>		<u>30MG</u>	<u>A076135 003</u> May 11, 2006
<u>AB</u>	!	<u>40MG</u>	<u>A076135 001</u> Apr 11, 2003

ISOTRETINOIN

<u>AB</u>	AMNEAL PHARMS NY	<u>10MG</u>	<u>A207792 001</u> Sep 29, 2017
<u>AB</u>		<u>20MG</u>	<u>A207792 002</u> Sep 29, 2017
<u>AB</u>		<u>30MG</u>	<u>A207792 003</u> Sep 29, 2017
<u>AB</u>		<u>40MG</u>	<u>A207792 004</u> Sep 29, 2017

MYORISAN

<u>AB</u>	DOUGLAS PHARMS	<u>10MG</u>	<u>A076485 001</u> Jan 19, 2012
<u>AB</u>		<u>20MG</u>	<u>A076485 002</u> Jan 19, 2012
<u>AB</u>		<u>30MG</u>	<u>A076485 004</u> Aug 25, 2015
<u>AB</u>		<u>40MG</u>	<u>A076485 003</u> Jan 19, 2012

ZENATANE

<u>AB</u>	DR REDDYS LABS LTD	<u>10MG</u>	<u>A202099 001</u> Mar 25, 2013
<u>AB</u>		<u>20MG</u>	<u>A202099 002</u> Mar 25, 2013
<u>AB</u>		<u>30MG</u>	<u>A202099 004</u> Feb 23, 2015
<u>AB</u>		<u>40MG</u>	<u>A202099 003</u> Mar 25, 2013

ABSORICA

BX	+	SUN PHARM INDNS INC	<u>10MG</u>	N021951 001	May 25, 2012
BX	+		<u>20MG</u>	N021951 002	May 25, 2012
BX	+		<u>30MG</u>	N021951 003	May 25, 2012
BX	!+		<u>40MG</u>	N021951 004	May 25, 2012
	+		<u>25MG</u>	N021951 005	Aug 15, 2014
	+		<u>35MG</u>	N021951 006	Aug 15, 2014

ISRADIPINE

CAPSULE; ORAL

ISRADIPINE

<u>AB</u>	ELITE LABS INC	<u>2.5MG</u>	<u>A077169 001</u> Apr 24, 2006
<u>AB</u>		<u>5MG</u>	<u>A077169 002</u> Apr 24, 2006
<u>AB</u>	WATSON LABS TEVA	<u>2.5MG</u>	<u>A077317 001</u> Jan 05, 2006
<u>AB</u>	!	<u>5MG</u>	<u>A077317 002</u> Jan 05, 2006

ITRACONAZOLE

CAPSULE; ORAL

ITRACONAZOLE

<u>AB</u>	ACCORD HLTHCARE	<u>100MG</u>	<u>A205991 001</u> May 26, 2016
<u>AB</u>	ALEMBIC PHARMS LTD	<u>100MG</u>	<u>A206741 001</u> Dec 13, 2016
<u>AB</u>	ALKEM LABS LTD	<u>100MG</u>	<u>A208591 001</u> Jun 12, 2017
<u>AB</u>	AMNEAL PHARMS	<u>100MG</u>	<u>A205080 001</u> Sep 26, 2016
<u>AB</u>	JUBILANT GENERICS	<u>100MG</u>	<u>A203445 001</u> Feb 23, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>100MG</u>	<u>A200463 001</u> Jul 20, 2012
<u>AB</u>	PAR PHARM INC	<u>100MG</u>	<u>A205724 001</u> Dec 13, 2016
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A076104 001</u> May 28, 2004
<u>AB</u>	ZYDUS PHARMS USA INC	<u>100MG</u>	<u>A204672 001</u> Sep 19, 2017

SPORANOX

<u>AB</u>	+!	JANSSEN PHARMS	<u>100MG</u>	<u>N020083 001</u> Sep 11, 1992
-----------	----	----------------	--------------	---------------------------------

SOLUTION; ORAL

SPORANOX

+!	JANSSEN PHARMS	10MG/ML	N020657 001	Feb 21, 1997
----	----------------	---------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-237 (of 436)

ITRACONAZOLE

TABLET;ORAL

ONMEL

+! SEBELA IRELAND LTD 200MG

N022484 001 Apr 29, 2010

IVABRADINE HYDROCHLORIDE

TABLET;ORAL

CORLANOR

+ AMGEN INC

EQ 5MG BASE

+!

EQ 7.5MG BASE

N206143 001 Apr 15, 2015

N206143 002 Apr 15, 2015

IVACAFTOR

GRANULE;ORAL

KALYDECO

+ VERTEX PHARMS INC

50MG/PACKET

+!

75MG/PACKET

N207925 001 Mar 17, 2015

N207925 002 Mar 17, 2015

TABLET;ORAL

KALYDECO

+! VERTEX PHARMS

150MG

N203188 001 Jan 31, 2012

IVACAFTOR; LUMACAFTOR

TABLET;ORAL

ORKAMBI

+ VERTEX PHARMS INC

125MG;100MG

+!

125MG;200MG

N206038 002 Sep 28, 2016

N206038 001 Jul 02, 2015

IVERMECTIN

CREAM;TOPICAL

SOOLANTRA

+! GALDERMA LABS LP

1%

N206255 001 Dec 19, 2014

LOTION;TOPICAL

SKLICE

+! ARBOR PHARMS LLC

0.5%

N202736 001 Feb 07, 2012

TABLET;ORAL

IVERMECTIN

AB EDENBRIDGE PHARMS

3MG

A204154 001 Oct 24, 2014

STROMECTOL

AB +! MERCK SHARP DOHME

3MG

N050742 002 Oct 08, 1998

IXABEPILONE

INJECTABLE;IV (INFUSION)

IXEMPRA KIT

+! R-PHARM US LLC

15MG/VIAL

+

45MG/VIAL

N022065 001 Oct 16, 2007

N022065 002 Oct 16, 2007

IXAZOMIB CITRATE

CAPSULE;ORAL

NINLARO

+ MILLENNIUM PHARMS

EQ 2.3MG BASE

+

EQ 3MG BASE

+

EQ 4MG BASE

N208462 001 Nov 20, 2015

N208462 002 Nov 20, 2015

N208462 003 Nov 20, 2015

KANAMYCIN SULFATE

INJECTABLE;INJECTION

KANAMYCIN SULFATE

FRESENIUS KABI USA

EQ 500MG BASE/2ML

!

EQ 1GM BASE/3ML

A065111 001 Dec 17, 2002

A065111 002 Dec 17, 2002

KETAMINE HYDROCHLORIDE

INJECTABLE;INJECTION

KETALAR

AP +! PAR STERILE PRODUCTS

EQ 10MG BASE/ML

N016812 001

AP +!

EQ 50MG BASE/ML

N016812 002

AP +!

EQ 100MG BASE/ML

N016812 003

KETAMINE HYDROCHLORIDE

AP HOSPIRA

EQ 50MG BASE/ML

A074549 001 Jun 27, 1996

AP

EQ 100MG BASE/ML

A074549 002 Jun 27, 1996

AP MYLAN INSTITUTIONAL

EQ 10MG BASE/ML

A076092 001 Sep 30, 2008

AP

EQ 50MG BASE/ML

A076092 002 Dec 28, 2001

AP

EQ 100MG BASE/ML

A076092 003 Oct 25, 2002

AP WEST-WARD PHARMS INT

EQ 50MG BASE/ML

A074524 001 Mar 22, 1996

AP

EQ 100MG BASE/ML

A074524 002 Mar 22, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-238 (of 436)

KETOCONAZOLE

AEROSOL, FOAM;TOPICAL

EXTINA

AT +! MYLAN PHARMS INC 2%

N021738 001 Jun 12, 2007

KETOCONAZOLE

AT PERRIGO ISRAEL 2%

A091550 001 Aug 25, 2011

CREAM;TOPICAL

KETOCONAZOLE

AB FOUGERA PHARMS 2%

A076294 001 Apr 28, 2004

AB ! TEVA 2%

A075581 001 Apr 25, 2000

KETOZOLE

AB TARO 2%

A075638 001 Dec 18, 2002

GEL;TOPICAL

XOLEGEL

+! AQUA PHARMS

2%

N021946 001 Jul 28, 2006

SHAMPOO;TOPICAL

KETOCONAZOLE

AB PERRIGO NEW YORK 2%

A076419 001 Jan 07, 2004

AB TOLMAR 2%

A076942 001 Apr 11, 2005

NIZORAL

AB +! JANSSEN PHARMS 2%

N019927 001 Aug 31, 1990

TABLET;ORAL

KETOCONAZOLE

AB MYLAN 200MG

A075597 001 Dec 23, 1999

AB TARO 200MG

A075319 001 Jun 15, 1999

AB ! TEVA 200MG

A075273 001 Jun 15, 1999

KETOPROFEN

CAPSULE;ORAL

KETOPROFEN

AB HERITAGE PHARMS INC 50MG

A074014 002 Jan 29, 1993

AB 75MG

A074014 003 Jan 29, 1993

AB MYLAN 50MG

A074035 002 Dec 31, 1996

AB 75MG

A074035 003 Dec 31, 1996

AB TEVA 50MG

A073516 001 Dec 22, 1992

AB ! 75MG

A073517 001 Dec 22, 1992

HERITAGE PHARMS INC 25MG

A074014 001 Jan 29, 1993

CAPSULE, EXTENDED RELEASE;ORAL

KETOPROFEN

AB ACTAVIS LABS FL INC 100MG

A075270 002 Mar 24, 1999

AB 150MG

A075270 003 Mar 24, 1999

AB 200MG

A075270 001 Mar 24, 1999

AB MYLAN 100MG

A075679 003 Feb 20, 2002

AB 150MG

A075679 002 Feb 20, 2002

AB ! 200MG

A075679 001 Feb 20, 2002

KETOROLAC TROMETHAMINE

INJECTABLE;INJECTION

KETOROLAC TROMETHAMINE

AP AMPHASTAR PHARM 15MG/ML

A076209 001 Jul 21, 2004

AP 30MG/ML

A076209 002 Jul 21, 2004

AP CYCLE PHARMS LTD 30MG/ML

A209900 001 Sep 15, 2017

AP FRESENIUS KABI USA 15MG/ML

A075784 001 Jan 11, 2002

AP 15MG/ML

A203242 001 Oct 07, 2015

AP 30MG/ML

A075784 002 Jan 11, 2002

AP 30MG/ML

A203242 002 Oct 07, 2015

AP GLAND PHARMA LTD 15MG/ML

A204216 001 Nov 01, 2016

AP 30MG/ML

A204216 002 Nov 01, 2016

AP ! HOSPIRA 15MG/ML

A074802 001 Jun 05, 1997

AP 15MG/ML

A074993 001 Jan 27, 1999

AP ! 30MG/ML

A074802 002 Jun 05, 1997

AP 30MG/ML

A074993 002 Jan 27, 1999

AP SAGENT PHARMS 15MG/ML

A091065 001 Nov 27, 2013

AP 30MG/ML

A091065 002 Nov 27, 2013

AP SANDOZ INC 30MG/ML

A076271 002 Oct 06, 2004

AP WOCKHARDT 15MG/ML

A077942 001 Mar 27, 2007

AP 30MG/ML

A077942 002 Mar 27, 2007

SOLUTION/DROPS;OPHTHALMIC

ACULAR

AT +! ALLERGAN 0.5%

N019700 001 Nov 09, 1992

ACULAR LS

AT +! ALLERGAN 0.4%

N021528 001 May 30, 2003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-239 (of 436)

KETOROLAC TROMETHAMINE

SOLUTION/DROPS;OPHTHALMIC

KETOROLAC TROMETHAMINE

<u>AT</u>	AKORN	<u>0.4%</u>	<u>A078399</u> <u>001</u>	Nov 05, 2009
<u>AT</u>		<u>0.5%</u>	<u>A078434</u> <u>001</u>	Nov 05, 2009
<u>AT</u>	ALCON PHARMS LTD	<u>0.4%</u>	<u>A078721</u> <u>001</u>	Nov 05, 2009
<u>AT</u>	APOTEX INC	<u>0.4%</u>	<u>A077308</u> <u>001</u>	Nov 05, 2009
<u>AT</u>		<u>0.5%</u>	<u>A076109</u> <u>001</u>	Nov 05, 2009
<u>AT</u>	SANDOZ INC	<u>0.5%</u>	<u>A076583</u> <u>001</u>	Nov 05, 2009
<u>AT</u>	SUN PHARMA GLOBAL	<u>0.5%</u>	<u>A090017</u> <u>001</u>	Nov 05, 2009

ACUVAIL

+! ALLERGAN 0.45%

N022427 001 Jul 22, 2009

SPRAY, METERED;NASAL

SPRIX

+! EGALET US INC 15.75MG/SPRAY

N022382 001 May 14, 2010

TABLET;ORAL

KETOROLAC TROMETHAMINE

<u>AB</u> !	MYLAN	<u>10MG</u>	<u>A074761</u> <u>001</u>	May 16, 1997
<u>AB</u>	PLIVA	<u>10MG</u>	<u>A075284</u> <u>001</u>	Jun 23, 1999
<u>AB</u>	TEVA	<u>10MG</u>	<u>A074754</u> <u>001</u>	May 16, 1997

KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION;IRRIGATION

OMIDRIA

+! OMEROS EQ 0.3% BASE;EQ 1% BASE

N205388 001 May 30, 2014

L-GLUTAMINE

FOR SOLUTION;ORAL

ENDARI

+! EMMAUS MEDCL 5GM/PACKET

N208587 001 Jul 07, 2017

NUTRESTORE

+! EMMAUS MEDCL 5GM/PACKET

N021667 001 Jun 10, 2004

LABETALOL HYDROCHLORIDE

INJECTABLE;INJECTION

LABETALOL HYDROCHLORIDE

<u>AP</u>	AKORN INC	<u>5MG/ML</u>	<u>A075431</u> <u>001</u>	Nov 29, 1999
<u>AP</u>		<u>5MG/ML</u>	<u>A075524</u> <u>001</u>	Nov 29, 1999
<u>AP</u>	GLAND PHARMA LTD	<u>5MG/ML</u>	<u>A090699</u> <u>001</u>	Apr 03, 2012
<u>AP</u> !	HOSPIRA	<u>5MG/ML</u>	<u>A075239</u> <u>001</u>	Nov 29, 1999
<u>AP</u> !		<u>5MG/ML</u>	<u>A075240</u> <u>001</u>	Nov 29, 1999
<u>AP</u>	SAGENT AGILA LLC	<u>5MG/ML</u>	<u>A079134</u> <u>001</u>	Feb 03, 2010
<u>AP</u>	WEST-WARD PHARMS INT	<u>5MG/ML</u>	<u>A075303</u> <u>001</u>	May 28, 1999

TABLET;ORAL

LABETALOL HYDROCHLORIDE

<u>AB</u>	INNOGENIX	<u>100MG</u>	<u>A075215</u> <u>001</u>	Jul 29, 1999
<u>AB</u>		<u>200MG</u>	<u>A075215</u> <u>002</u>	Jul 29, 1999
<u>AB</u>		<u>300MG</u>	<u>A075215</u> <u>003</u>	Jul 29, 1999
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>100MG</u>	<u>A074787</u> <u>001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>A074787</u> <u>002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>A074787</u> <u>003</u>	Aug 03, 1998
<u>AB</u>	PAR FORM	<u>100MG</u>	<u>A200908</u> <u>001</u>	Jul 10, 2012
<u>AB</u>		<u>200MG</u>	<u>A200908</u> <u>002</u>	Jul 10, 2012
<u>AB</u>		<u>300MG</u>	<u>A200908</u> <u>003</u>	Jul 10, 2012
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>A075113</u> <u>001</u>	Aug 04, 1998
<u>AB</u> !		<u>200MG</u>	<u>A075113</u> <u>002</u>	Aug 04, 1998
<u>AB</u>		<u>300MG</u>	<u>A075113</u> <u>003</u>	Aug 04, 1998
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>A075133</u> <u>001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>A075133</u> <u>002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>A075133</u> <u>003</u>	Aug 03, 1998
<u>AB</u>	ZYDUS PHARMS USA INC	<u>100MG</u>	<u>A207743</u> <u>001</u>	Sep 19, 2017
<u>AB</u>		<u>200MG</u>	<u>A207743</u> <u>002</u>	Sep 19, 2017
<u>AB</u>		<u>300MG</u>	<u>A207743</u> <u>003</u>	Sep 19, 2017

TRANDATE

<u>AB</u>	CNTY LINE PHARMS	<u>100MG</u>	<u>N018716</u> <u>001</u>	May 24, 1985
<u>AB</u>		<u>200MG</u>	<u>N018716</u> <u>002</u>	Aug 01, 1984
<u>AB</u>		<u>300MG</u>	<u>N018716</u> <u>003</u>	Aug 01, 1984

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-240 (of 436)

LACOSAMIDE

SOLUTION; INTRAVENOUS

VIMPAT

+! UCB INC

200MG/20ML (10MG/ML)

N022254 001 Oct 28, 2008

SOLUTION; ORAL

VIMPAT

+! UCB INC

10MG/ML

N022255 001 Apr 20, 2010

TABLET; ORAL

VIMPAT

+ UCB INC

50MG

N022253 001 Oct 28, 2008

+ 100MG

N022253 002 Oct 28, 2008

+ 150MG

N022253 003 Oct 28, 2008

+! 200MG

N022253 004 Oct 28, 2008

LACTULOSE

FOR SOLUTION; ORAL

LACTULOSE

! CUMBERLAND PHARMS

10GM/PACKET

A074712 001 Dec 10, 1997

!

20GM/PACKET

A074712 002 Dec 10, 1997

SOLUTION; ORAL

CONSTILAC

AA ALRA

10GM/15ML

A071054 001 Jul 26, 1988

LACTULOSE

AA ANI PHARMS

10GM/15ML

A078430 001 Nov 28, 2007

AA FRESENIUS KABI

10GM/15ML

A090503 001 Jan 25, 2012

AA ! HI TECH PHARMA

10GM/15ML

A074076 001 Jul 03, 1995

AA PHARM ASSOC

10GM/15ML

A074623 001 Jul 30, 1996

AA VINTAGE PHARMS

10GM/15ML

A075993 001 Jul 26, 2001

AA VISTAPHARM

10GM/15ML

A074138 001 Sep 30, 1992

AA WEST-WARD PHARMS

10GM/15ML

A073591 001 May 29, 1992

INT

AA WOCKHARDT BIO AG

10GM/15ML

A074602 001 Nov 14, 1996

SOLUTION; ORAL, RECTAL

CHOLAC

AA ALRA

10GM/15ML

A071331 001 Jul 26, 1988

ENULOSE

AA ! ACTAVIS MID

10GM/15ML

A071548 001 Aug 15, 1988

ATLANTIC

GENERLAC

AA WOCKHARDT BIO AG

10GM/15ML

A074603 001 Oct 31, 1996

LACTULOSE

AA ANI PHARMS

10GM/15ML

A090426 001 Nov 21, 2008

AA BIO-PHARM INC

10GM/15ML

A203762 001 Mar 27, 2015

AA FRESENIUS KABI

10GM/15ML

A090502 001 Jan 25, 2012

AA HI TECH PHARMA

10GM/15ML

A074077 001 Jul 03, 1995

LAMIVUDINE

SOLUTION; ORAL

EPIVIR

AA +! VIIIV HLTHCARE

10MG/ML

N020596 001 Nov 17, 1995

LAMIVUDINE

AA AUROBINDO PHARMA LTD

10MG/ML

A077695 001 Nov 21, 2016

AA SILARX PHARMS INC

10MG/ML

A203564 001 Oct 31, 2014

EPIVIR-HBV

+! GLAXOSMITHKLINE

5MG/ML

N021004 001 Dec 08, 1998

TABLET; ORAL

EPIVIR

AB + VIIIV HLTHCARE

150MG

N020564 001 Nov 17, 1995

AB +!

300MG

N020564 003 Jun 24, 2002

EPIVIR-HBV

AB +! GLAXOSMITHKLINE

100MG

N021003 001 Dec 08, 1998

LAMIVUDINE

AB APOTEX

150MG

A091606 001 Dec 02, 2011

AB

300MG

A091606 002 Dec 02, 2011

AB APOTEX INC

100MG

A202941 001 Jan 02, 2014

AB APPCO PHARMA LLC

150MG

A206974 001 Nov 21, 2016

AB

300MG

A206974 002 Nov 21, 2016

AB AUROBINDO PHARMA LTD

150MG

A077464 001 Nov 21, 2016

AB

150MG

A202032 001 Nov 17, 2011

AB

300MG

A077464 002 Nov 21, 2016

AB

300MG

A202032 002 Nov 17, 2011

AB CIPLA LTD

150MG

A077221 001 Mar 03, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-241 (of 436)

LAMIVUDINE

TABLET;ORAL

LAMIVUDINE

<u>AB</u>		<u>300MG</u>	<u>A077221</u> <u>002</u>	Mar 03, 2017
<u>AB</u>	ECI PHARMS LLC	<u>150MG</u>	<u>A203586</u> <u>001</u>	Nov 21, 2016
<u>AB</u>	HETERO LABS LTD V	<u>100MG</u>	<u>A203260</u> <u>001</u>	Jan 02, 2014
<u>AB</u>		<u>150MG</u>	<u>A203277</u> <u>001</u>	Jan 06, 2014
<u>AB</u>		<u>300MG</u>	<u>A203277</u> <u>002</u>	Jan 06, 2014
<u>AB</u>	LUPIN LTD	<u>150MG</u>	<u>A205217</u> <u>001</u>	Dec 18, 2014
<u>AB</u>		<u>300MG</u>	<u>A205217</u> <u>002</u>	Dec 18, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>100MG</u>	<u>A204002</u> <u>001</u>	Dec 31, 2014
<u>AB</u>		<u>150MG</u>	<u>A204528</u> <u>001</u>	Mar 04, 2016
<u>AB</u>		<u>300MG</u>	<u>A204528</u> <u>002</u>	Mar 04, 2016

LAMIVUDINE; ZIDOVUDINE

TABLET;ORAL

COMBIVIR

<u>AB</u>	+!	VIVI HLTHCARE	<u>150MG;300MG</u>	<u>N020857</u> <u>001</u>	Sep 26, 1997
-----------	----	---------------	--------------------	---------------------------	--------------

LAMIVUDINE AND ZIDOVUDINE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>150MG;300MG</u>	<u>A077558</u> <u>001</u>	May 05, 2017
<u>AB</u>		<u>150MG;300MG</u>	<u>A202418</u> <u>001</u>	May 15, 2012
<u>AB</u>	HETERO LABS LTD III	<u>150MG;300MG</u>	<u>A079124</u> <u>001</u>	Sep 17, 2015
<u>AB</u>	HETERO LABS LTD V	<u>150MG;300MG</u>	<u>A203259</u> <u>001</u>	Feb 03, 2014
<u>AB</u>	LUPIN LTD	<u>150MG;300MG</u>	<u>A090246</u> <u>001</u>	May 15, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>150MG;300MG</u>	<u>A204005</u> <u>001</u>	Aug 28, 2014
<u>AB</u>	STRIDES PHARMA	<u>150MG;300MG</u>	<u>A079128</u> <u>001</u>	May 13, 2015
<u>AB</u>	TEVA PHARMS BX	<u>150MG;300MG</u>	<u>A079081</u> <u>001</u>	May 25, 2011
	PHEMACARE	150MG;300MG	N022018 001	Mar 17, 2017

LAMOTRIGINE

TABLET;ORAL

LAMICTAL

<u>AB</u>	+!	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N020241</u> <u>005</u>	Dec 27, 1994
<u>AB</u>	+		<u>100MG</u>	<u>N020241</u> <u>001</u>	Dec 27, 1994
<u>AB</u>	+		<u>150MG</u>	<u>N020241</u> <u>002</u>	Dec 27, 1994
<u>AB</u>	+		<u>200MG</u>	<u>N020241</u> <u>003</u>	Dec 27, 1994

LAMOTRIGINE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A090607</u> <u>001</u>	Jan 13, 2011
<u>AB</u>		<u>100MG</u>	<u>A090607</u> <u>002</u>	Jan 13, 2011
<u>AB</u>		<u>150MG</u>	<u>A090607</u> <u>003</u>	Jan 13, 2011
<u>AB</u>		<u>200MG</u>	<u>A090607</u> <u>004</u>	Jan 13, 2011
<u>AB</u>	ALKEM LABS LTD	<u>25MG</u>	<u>A200694</u> <u>001</u>	Jun 14, 2013
<u>AB</u>		<u>100MG</u>	<u>A200694</u> <u>002</u>	Jun 14, 2013
<u>AB</u>		<u>150MG</u>	<u>A200694</u> <u>003</u>	Jun 14, 2013
<u>AB</u>		<u>200MG</u>	<u>A200694</u> <u>004</u>	Jun 14, 2013
<u>AB</u>	APOTEX INC	<u>25MG</u>	<u>A078625</u> <u>001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078625</u> <u>002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A078625</u> <u>003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078625</u> <u>004</u>	Jan 27, 2009
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A078956</u> <u>001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078956</u> <u>002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A078956</u> <u>003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078956</u> <u>004</u>	Jan 27, 2009
<u>AB</u>	CIPLA LTD	<u>25MG</u>	<u>A077783</u> <u>001</u>	Nov 01, 2010
<u>AB</u>		<u>100MG</u>	<u>A077783</u> <u>002</u>	Nov 01, 2010
<u>AB</u>		<u>150MG</u>	<u>A077783</u> <u>003</u>	Nov 01, 2010
<u>AB</u>		<u>200MG</u>	<u>A077783</u> <u>004</u>	Nov 01, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>25MG</u>	<u>A076708</u> <u>001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A076708</u> <u>002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A076708</u> <u>003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A076708</u> <u>004</u>	Jan 27, 2009
<u>AB</u>	GLENMARK GENERICS	<u>25MG</u>	<u>A090169</u> <u>001</u>	May 12, 2012
<u>AB</u>		<u>100MG</u>	<u>A090169</u> <u>002</u>	May 12, 2012
<u>AB</u>		<u>150MG</u>	<u>A090169</u> <u>003</u>	May 12, 2012
<u>AB</u>		<u>200MG</u>	<u>A090169</u> <u>004</u>	May 12, 2012
<u>AB</u>	JUBILANT CADISTA	<u>25MG</u>	<u>A079132</u> <u>001</u>	Jan 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079132</u> <u>002</u>	Jan 27, 2009
<u>AB</u>		<u>150MG</u>	<u>A079132</u> <u>003</u>	Jan 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A079132</u> <u>004</u>	Jan 27, 2009
<u>AB</u>	LUPIN LTD	<u>25MG</u>	<u>A078691</u> <u>001</u>	Jun 01, 2010
<u>AB</u>		<u>100MG</u>	<u>A078691</u> <u>002</u>	Jun 01, 2010
<u>AB</u>		<u>150MG</u>	<u>A078691</u> <u>003</u>	Jun 01, 2010
<u>AB</u>		<u>200MG</u>	<u>A078691</u> <u>004</u>	Jun 01, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-242 (of 436)

**LAMOTRIGINE**

TABLET;ORAL

**LAMOTRIGINE**

<b>AB</b>	MYLAN	<b>25MG</b>	<b>A077420 001</b>	Jan 27, 2009
<b>AB</b>		<b>100MG</b>	<b>A077420 002</b>	Jan 27, 2009
<b>AB</b>		<b>150MG</b>	<b>A077420 003</b>	Jan 27, 2009
<b>AB</b>		<b>200MG</b>	<b>A077420 004</b>	Jan 27, 2009
<b>AB</b>	TARO PHARM INDS	<b>25MG</b>	<b>A078525 001</b>	Jan 27, 2009
<b>AB</b>		<b>100MG</b>	<b>A078525 002</b>	Jan 27, 2009
<b>AB</b>		<b>150MG</b>	<b>A078525 003</b>	Jan 27, 2009
<b>AB</b>		<b>200MG</b>	<b>A078525 004</b>	Jan 27, 2009
<b>AB</b>	TEVA	<b>25MG</b>	<b>A076388 001</b>	Aug 30, 2006
<b>AB</b>		<b>100MG</b>	<b>A076388 002</b>	Aug 30, 2006
<b>AB</b>		<b>150MG</b>	<b>A076388 003</b>	Aug 30, 2006
<b>AB</b>		<b>200MG</b>	<b>A076388 004</b>	Aug 30, 2006
<b>AB</b>	TORRENT PHARMS	<b>25MG</b>	<b>A078947 001</b>	Jan 27, 2009
<b>AB</b>		<b>100MG</b>	<b>A078947 002</b>	Jan 27, 2009
<b>AB</b>		<b>150MG</b>	<b>A078947 003</b>	Jan 27, 2009
<b>AB</b>		<b>200MG</b>	<b>A078947 004</b>	Jan 27, 2009
<b>AB</b>	UNICHEM LABS LTD	<b>25MG</b>	<b>A090170 001</b>	Oct 06, 2011
<b>AB</b>		<b>100MG</b>	<b>A090170 002</b>	Oct 06, 2011
<b>AB</b>		<b>150MG</b>	<b>A090170 003</b>	Oct 06, 2011
<b>AB</b>		<b>200MG</b>	<b>A090170 004</b>	Oct 06, 2011
<b>AB</b>	ZYDUS PHARMS USA	<b>25MG</b>	<b>A077633 001</b>	Jan 27, 2009
<b>AB</b>		<b>100MG</b>	<b>A077633 003</b>	Jan 27, 2009
<b>AB</b>		<b>150MG</b>	<b>A077633 004</b>	Jan 27, 2009
<b>AB</b>		<b>200MG</b>	<b>A077633 005</b>	Jan 27, 2009
		50MG	A077633 002	Jan 27, 2009
		250MG	A077633 006	Jan 27, 2009

TABLET, CHEWABLE;ORAL

**LAMICTAL CD**

<b>AB</b>	+	GLAXOSMITHKLINE LLC	<b>2MG</b>	<b>N020764 004</b>	Sep 08, 2000
<b>AB</b>	+		<b>5MG</b>	<b>N020764 001</b>	Aug 24, 1998
<b>AB</b>	+!		<b>25MG</b>	<b>N020764 002</b>	Aug 24, 1998

**LAMOTRIGINE**

<b>AB</b>	ALEMBIC PHARMS LTD	<b>5MG</b>	<b>A201168 001</b>	Jun 12, 2014
<b>AB</b>		<b>25MG</b>	<b>A201168 002</b>	Jun 12, 2014
<b>AB</b>	AUROBINDO PHARMA	<b>5MG</b>	<b>A090401 002</b>	Nov 04, 2009
<b>AB</b>		<b>25MG</b>	<b>A090401 003</b>	Nov 04, 2009
<b>AB</b>	DR REDDYS LABS LTD	<b>5MG</b>	<b>A076701 001</b>	Jan 22, 2009
<b>AB</b>		<b>25MG</b>	<b>A076701 002</b>	Jan 22, 2009
<b>AB</b>	GLENMARK PHARMS LTD	<b>5MG</b>	<b>A079099 001</b>	Feb 19, 2009
<b>AB</b>		<b>25MG</b>	<b>A079099 002</b>	Feb 19, 2009
<b>AB</b>	JUBILANT GENERICS	<b>5MG</b>	<b>A200220 001</b>	Feb 28, 2011
<b>AB</b>		<b>25MG</b>	<b>A200220 002</b>	Feb 28, 2011
<b>AB</b>	TARO	<b>5MG</b>	<b>A079204 001</b>	Feb 04, 2009
<b>AB</b>		<b>25MG</b>	<b>A079204 002</b>	Feb 04, 2009
<b>AB</b>	TEVA	<b>5MG</b>	<b>A076420 001</b>	Jun 21, 2006
<b>AB</b>		<b>25MG</b>	<b>A076420 002</b>	Jun 21, 2006
<b>AB</b>	WATSON LABS	<b>2MG</b>	<b>A076928 001</b>	Jan 22, 2009
<b>AB</b>		<b>5MG</b>	<b>A076928 002</b>	Jan 22, 2009
<b>AB</b>		<b>25MG</b>	<b>A076928 003</b>	Jan 22, 2009
<b>AB</b>	ZYDUS PHARMS USA	<b>5MG</b>	<b>A078009 002</b>	Jan 22, 2009
	INC			
<b>AB</b>		<b>25MG</b>	<b>A078009 003</b>	Jan 22, 2009

TABLET, EXTENDED RELEASE;ORAL

**LAMICTAL XR**

<b>AB</b>	+	GLAXOSMITHKLINE LLC	<b>25MG</b>	<b>N022115 001</b>	May 29, 2009
<b>AB</b>	+!		<b>50MG</b>	<b>N022115 002</b>	May 29, 2009
<b>AB</b>	+		<b>100MG</b>	<b>N022115 003</b>	May 29, 2009
<b>AB</b>	+!		<b>200MG</b>	<b>N022115 004</b>	May 29, 2009
<b>AB</b>	+		<b>250MG</b>	<b>N022115 006</b>	Jun 21, 2011
<b>AB</b>	+		<b>300MG</b>	<b>N022115 005</b>	Apr 14, 2010

**LAMOTRIGINE**

<b>AB</b>	ACTAVIS ELIZABETH	<b>100MG</b>	<b>A200672 003</b>	Oct 17, 2013
<b>AB</b>		<b>200MG</b>	<b>A200672 004</b>	Oct 17, 2013
<b>AB</b>		<b>25MG</b>	<b>A200672 001</b>	Oct 17, 2013
<b>AB</b>		<b>50MG</b>	<b>A200672 002</b>	Oct 17, 2013
<b>AB</b>		<b>250MG</b>	<b>A203733 001</b>	Nov 13, 2013
<b>AB</b>		<b>300MG</b>	<b>A200672 005</b>	Oct 17, 2013
<b>AB</b>	ANCHEM PHARMS	<b>25MG</b>	<b>A201374 001</b>	Dec 26, 2012
<b>AB</b>		<b>50MG</b>	<b>A201374 002</b>	Dec 26, 2012
<b>AB</b>		<b>100MG</b>	<b>A201374 003</b>	Dec 26, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-243 (of 436)

LAMOTRIGINE

TABLET, EXTENDED RELEASE;ORAL

LAMOTRIGINE

<u>AB</u>		<u>200MG</u>	<u>A201374 004</u>	Dec 26, 2012
<u>AB</u>		<u>250MG</u>	<u>A201374 005</u>	Dec 26, 2012
<u>AB</u>		<u>300MG</u>	<u>A201374 006</u>	Dec 26, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>25MG</u>	<u>A202383 001</u>	Jun 19, 2013
<u>AB</u>		<u>50MG</u>	<u>A202383 002</u>	Jun 19, 2013
<u>AB</u>		<u>100MG</u>	<u>A202383 003</u>	Jun 19, 2013
<u>AB</u>		<u>200MG</u>	<u>A202383 004</u>	Jun 19, 2013
<u>AB</u>		<u>300MG</u>	<u>A202383 005</u>	Jun 19, 2013
<u>AB</u>	HANDA PHARMS LLC	<u>25MG</u>	<u>A202887 001</u>	Jun 17, 2013
<u>AB</u>		<u>50MG</u>	<u>A202887 002</u>	Jun 17, 2013
<u>AB</u>		<u>100MG</u>	<u>A202887 003</u>	Jun 17, 2013
<u>AB</u>		<u>200MG</u>	<u>A202887 004</u>	Jun 17, 2013
<u>AB</u>	PAR PHARM	<u>25MG</u>	<u>A201791 001</u>	Jan 18, 2013
<u>AB</u>		<u>50MG</u>	<u>A201791 002</u>	Jan 18, 2013
<u>AB</u>		<u>100MG</u>	<u>A201791 003</u>	Jan 18, 2013
<u>AB</u>		<u>200MG</u>	<u>A201791 004</u>	Jan 18, 2013
<u>AB</u>		<u>250MG</u>	<u>A201791 005</u>	Jan 18, 2013
<u>AB</u>		<u>300MG</u>	<u>A201791 006</u>	Jan 18, 2013
<u>AB</u>	TORRENT PHARMS LTD	<u>25MG</u>	<u>A203370 001</u>	Dec 23, 2013
<u>AB</u>		<u>50MG</u>	<u>A203370 002</u>	Dec 23, 2013
<u>AB</u>		<u>100MG</u>	<u>A203370 003</u>	Dec 23, 2013
<u>AB</u>		<u>200MG</u>	<u>A203370 004</u>	Dec 23, 2013
<u>AB</u>	WOCKHARDT LTD	<u>25MG</u>	<u>A202498 001</u>	Jan 04, 2013
<u>AB</u>		<u>50MG</u>	<u>A202498 002</u>	Jan 04, 2013
<u>AB</u>		<u>100MG</u>	<u>A202498 003</u>	Jan 04, 2013
<u>AB</u>		<u>200MG</u>	<u>A202498 004</u>	Jan 04, 2013
<u>AB</u>		<u>300MG</u>	<u>A202498 005</u>	Jan 04, 2013

TABLET, ORALLY DISINTEGRATING;ORAL

LAMICTAL ODT

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>25MG</u>	<u>N022251 001</u>	May 08, 2009
<u>AB</u>	+!		<u>50MG</u>	<u>N022251 002</u>	May 08, 2009
<u>AB</u>	+		<u>100MG</u>	<u>N022251 003</u>	May 08, 2009
<u>AB</u>	+		<u>200MG</u>	<u>N022251 004</u>	May 08, 2009

LAMOTRIGINE

<u>AB</u>	IMPAX LABS INC	<u>25MG</u>	<u>A200828 001</u>	Jul 15, 2013
<u>AB</u>		<u>50MG</u>	<u>A200828 002</u>	Jul 15, 2013
<u>AB</u>		<u>100MG</u>	<u>A200828 003</u>	Jul 15, 2013
<u>AB</u>		<u>200MG</u>	<u>A200828 004</u>	Jul 15, 2013
<u>AB</u>	PAR PHARM	<u>25MG</u>	<u>A204158 001</u>	Oct 27, 2015
<u>AB</u>		<u>50MG</u>	<u>A204158 002</u>	Oct 27, 2015
<u>AB</u>		<u>100MG</u>	<u>A204158 003</u>	Oct 27, 2015
<u>AB</u>		<u>200MG</u>	<u>A204158 004</u>	Oct 27, 2015
<u>AB</u>	SCIEGEN PHARMS INC	<u>25MG</u>	<u>A206382 001</u>	Jun 17, 2016
<u>AB</u>		<u>50MG</u>	<u>A206382 002</u>	Jun 17, 2016
<u>AB</u>		<u>100MG</u>	<u>A206382 003</u>	Jun 17, 2016
<u>AB</u>		<u>200MG</u>	<u>A206382 004</u>	Jun 17, 2016

LANREOTIDE ACETATE

SOLUTION;SUBCUTANEOUS

SOMATULINE DEPOT

+!	IPSEN PHARMA	EQ 60MG BASE/0.2ML (EQ 60MG BASE/0.2ML)	N022074 001	Aug 30, 2007
+!		EQ 90MG BASE/0.3ML (EQ 90MG BASE/0.3ML)	N022074 002	Aug 30, 2007
+!		EQ 120MG BASE/0.5ML (EQ 120MG BASE/0.5ML)	N022074 003	Aug 30, 2007

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

<u>AB</u>	AJANTA PHARMA LTD	<u>15MG</u>	<u>A203957 001</u>	Oct 14, 2016
<u>AB</u>		<u>30MG</u>	<u>A203957 002</u>	Oct 14, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>15MG</u>	<u>A091269 001</u>	Oct 15, 2010
<u>AB</u>		<u>30MG</u>	<u>A091269 002</u>	Oct 15, 2010
<u>AB</u>	INVENTIA HLTHCARE	<u>15MG</u>	<u>A205868 001</u>	Aug 30, 2017
<u>AB</u>		<u>30MG</u>	<u>A205868 002</u>	Aug 30, 2017
<u>AB</u>	KREMERS URBAN PHARMS	<u>15MG</u>	<u>A207156 001</u>	Sep 28, 2017
<u>AB</u>		<u>30MG</u>	<u>A207156 002</u>	Sep 28, 2017
<u>AB</u>	LABS LICONSA	<u>15MG</u>	<u>A203203 001</u>	Jul 25, 2016
<u>AB</u>		<u>30MG</u>	<u>A203203 002</u>	Jul 25, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>15MG</u>	<u>A090763 001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A090763 002</u>	Nov 10, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-244 (of 436)

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

LANSOPRAZOLE

<u>AB</u>	NATCO PHARMA LTD	<u>15MG</u>	<u>A201921</u> <u>001</u>	Dec 18, 2012
<u>AB</u>		<u>30MG</u>	<u>A201921</u> <u>002</u>	Dec 18, 2012
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>A090331</u> <u>001</u>	Apr 23, 2010
<u>AB</u>		<u>30MG</u>	<u>A090331</u> <u>002</u>	Apr 23, 2010
<u>AB</u>	SUN PHARM INDs LTD	<u>15MG</u>	<u>A202637</u> <u>001</u>	Sep 13, 2013
<u>AB</u>		<u>30MG</u>	<u>A091509</u> <u>001</u>	Sep 13, 2013
<u>AB</u>	TEVA PHARMS	<u>15MG</u>	<u>A077255</u> <u>001</u>	Nov 10, 2009
<u>AB</u>		<u>30MG</u>	<u>A077255</u> <u>002</u>	Nov 10, 2009
<u>AB</u>	WOCKHARDT USA	<u>15MG</u>	<u>A202176</u> <u>001</u>	Sep 14, 2012
<u>AB</u>		<u>30MG</u>	<u>A202176</u> <u>002</u>	Sep 14, 2012
<u>AB</u>	ZYDUS HLTHCARE	<u>15MG</u>	<u>A202366</u> <u>001</u>	Aug 19, 2013
<u>AB</u>		<u>30MG</u>	<u>A202366</u> <u>002</u>	Aug 19, 2013

LANSOPRAZOLE

<u>AB</u>	KRKA TOVARNA ZDRAVIL	<u>15MG</u>	<u>A091212</u> <u>001</u>	Sep 16, 2013
<u>AB</u>		<u>30MG</u>	<u>A091212</u> <u>002</u>	Sep 16, 2013
<u>AB</u>	+ TAKEDA PHARMS USA	<u>15MG</u>	<u>N020406</u> <u>001</u>	May 10, 1995
<u>AB</u>	+!	<u>30MG</u>	<u>N020406</u> <u>002</u>	May 10, 1995

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

<u>AB</u>	TEVA PHARMS USA	<u>15MG</u>	<u>A208784</u> <u>001</u>	Sep 21, 2017
<u>AB</u>		<u>30MG</u>	<u>A208784</u> <u>002</u>	Sep 21, 2017
<u>AB</u>	+ TAKEDA PHARMS USA	<u>15MG</u>	<u>N021428</u> <u>001</u>	Aug 30, 2002
<u>AB</u>	+!	<u>30MG</u>	<u>N021428</u> <u>002</u>	Aug 30, 2002

LANTHANUM CARBONATE

POWDER;ORAL

FOSRENOL

+	SHIRE DEV LLC	EQ 750MG BASE	N204734	001	Sep 24, 2014
+		EQ 1GM BASE	N204734	002	Sep 24, 2014

TABLET, CHEWABLE;ORAL

FOSRENOL

<u>AB</u>	+ SHIRE LLC	<u>EQ 500MG BASE</u>	<u>N021468</u> <u>002</u>	Oct 26, 2004
<u>AB</u>	+	<u>EQ 750MG BASE</u>	<u>N021468</u> <u>003</u>	Nov 23, 2005
<u>AB</u>	+!	<u>EQ 1GM BASE</u>	<u>N021468</u> <u>004</u>	Nov 23, 2005

LANTHANUM CARBONATE

<u>AB</u>	NATCO PHARMA LTD	<u>EQ 500MG BASE</u>	<u>A090978</u> <u>001</u>	Aug 11, 2017
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A090978</u> <u>002</u>	Aug 11, 2017
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090978</u> <u>003</u>	Aug 11, 2017

LAPATINIB DITOSYLATE

TABLET;ORAL

TYKERB

+	NOVARTIS PHARMS CORP	EQ 250MG BASE	N022059	001	Mar 13, 2007
---	----------------------	---------------	---------	-----	--------------

LATANOPROST

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

<u>AT</u>	AKORN	<u>0.005%</u>	<u>A090887</u> <u>001</u>	Jul 19, 2011
<u>AT</u>	AMRING PHARMS	<u>0.005%</u>	<u>A200925</u> <u>001</u>	Mar 22, 2011
<u>AT</u>	BAUSCH AND LOMB	<u>0.005%</u>	<u>A201006</u> <u>001</u>	Mar 22, 2011
<u>AT</u>	DR REDDYS LABS LTD	<u>0.005%</u>	<u>A202077</u> <u>001</u>	Feb 11, 2013
<u>AT</u>	FDC LTD	<u>0.005%</u>	<u>A202442</u> <u>001</u>	Apr 22, 2016
<u>AT</u>	MYLAN	<u>0.005%</u>	<u>A201786</u> <u>001</u>	Mar 22, 2011
<u>AT</u>	SANDOZ INC	<u>0.005%</u>	<u>A091449</u> <u>001</u>	Mar 22, 2011

XALATAN

<u>AT</u>	+! PHARMACIA AND UPJOHN	<u>0.005%</u>	<u>N020597</u> <u>001</u>	Jun 05, 1996
-----------	-------------------------	---------------	---------------------------	--------------

LATANOPROSTENE BUNOD

SOLUTION/DROPS;OPHTHALMIC

VYZULTA

+	BAUSCH AND LOMB	0.024%	N207795	001	Nov 02, 2017
---	-----------------	--------	---------	-----	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-245 (of 436)

LEDIPASVIR; SOFOSBUVIR

TABLET;ORAL  
 HARVONI  
 +! GILEAD SCIENCES INC 90MG;400MG

N205834 001 Oct 10, 2014

LEFLUNOMIDE

TABLET;ORAL

ARAVA

AB + SANOFI AVENTIS US 10MG

AB +! 20MG

LEFLUNOMIDE

AB ALEMBIC PHARMS LTD 10MG

AB 20MG

AB APOTEX INC 10MG

AB 20MG

AB BARR 10MG

AB 20MG

AB HERITAGE PHARMS INC 10MG

AB 20MG

AB TEVA PHARMS 10MG

AB 20MG

N020905 001 Sep 10, 1998

N020905 002 Sep 10, 1998

ARAVA

+! SANOFI AVENTIS US 100MG

N020905 003 Sep 10, 1998

LENALIDOMIDE

CAPSULE;ORAL

REVLIMID

+ CELGENE 2.5MG

N021880 005 Dec 21, 2011

+ 5MG

N021880 001 Dec 27, 2005

+ 10MG

N021880 002 Dec 27, 2005

+ 15MG

N021880 003 Jun 29, 2006

+ 20MG

N021880 006 Jun 05, 2013

+! 25MG

N021880 004 Jun 29, 2006

LENVATINIB MESYLATE

CAPSULE;ORAL

LENVIMA

+ EISAI INC EQ 4MG BASE

N206947 001 Feb 13, 2015

+! EQ 10MG BASE

N206947 002 Feb 13, 2015

LESINURAD

TABLET;ORAL

ZURAMPIC

+! IRONWOOD PHARMS INC 200MG

N207988 001 Dec 22, 2015

LETHERMOVIR

SOLUTION;IV (INFUSION)

PREVYMIS

+! MERCK SHARP DOHME 240MG/12ML (20MG/ML)

N209940 001 Nov 08, 2017

+! 480MG/24ML (20MG/ML)

N209940 002 Nov 08, 2017

TABLET;ORAL

PREVYMIS

+ MERCK SHARP DOHME 240MG

N209939 001 Nov 08, 2017

+! 480MG

N209939 002 Nov 08, 2017

LETROZOLE

TABLET;ORAL

FEMARA

AB +! NOVARTIS PHARMS 2.5MG

N020726 001 Jul 25, 1997

LETROZOLE

AB ACCORD HLTHCARE 2.5MG

A090934 001 Jun 03, 2011

AB APOTEX INC 2.5MG

A091303 001 Apr 19, 2012

AB DR REDDYS LABS LTD 2.5MG

A091191 001 Jun 03, 2011

AB FRESENIUS KABI 2.5MG

A090491 001 Jun 03, 2011

ONCOL

AB HIKMA PHARMS 2.5MG

A203796 001 Jun 03, 2016

AB INDICUS PHARMA 2.5MG

A201804 001 Jun 03, 2011

AB JIANGSU HENGRUI MED 2.5MG

A202716 001 May 16, 2013

AB MYLAN 2.5MG

A078190 001 Dec 24, 2008

AB NATCO PHARMA LTD 2.5MG

A200161 001 Jun 03, 2011

AB TEVA PHARMS 2.5MG

A090289 001 Jun 03, 2011

AB VINTAGE PHARMS LLC 2.5MG

A090789 001 Jun 03, 2011

AB WEST-WARD PHARMS 2.5MG

A090838 001 Jun 03, 2011

INT

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PREScription DRUG PRODUCT LIST**

3-246 (of 436)

## LETROZOLE; RIBOCICLIB SUCCINATE

TABLET, TABLET;ORAL  
KISQALI FEMARA CO-PACK (COPACKAGED)  
+! NOVARTIS PHARMS 2.5MG,N/A;N/A,EQ 200MG BASE N209935 001 May 04, 2017  
CORP

## LEUCOVORIN CALCIUM

## INJECTABLE; INJECTION

## LEUCOVORIN CALCIUM

<u>AP</u>	TEVA PHARMS USA	<u>EQ 100MG BASE/VIAL</u>	<u>A081277 001</u>	Sep 28, 1993
<u>AP</u>		<u>EQ 350MG BASE/VIAL</u>	<u>A040174 001</u>	Jun 12, 1997
<u>AP</u>	! WEST-WARD PHARMS INT	<u>EQ 50MG BASE/VIAL</u>	<u>A089384 001</u>	Sep 14, 1987
<u>AP</u>	!	<u>EQ 100MG BASE/VIAL</u>	<u>A089717 001</u>	Mar 28, 1988
<u>LEUCOVORIN CALCIUM PRESERVATIVE FREE</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 200MG BASE/VIAL</u>	<u>A040258 001</u>	Feb 26, 1999
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A040286 001</u>	Feb 26, 1999
<u>AP</u>	MYLAN LABS LTD	<u>EQ 100MG BASE/VIAL</u>	<u>A203800 001</u>	May 19, 2017
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A203800 002</u>	May 19, 2017
<u>AP</u>		<u>EQ 350MG BASE/VIAL</u>	<u>A203800 003</u>	May 19, 2017
<u>AP</u>	SAGENT PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>A200753 001</u>	Sep 06, 2012
<u>AP</u>		<u>EQ 100MG BASE/VIAL</u>	<u>A200753 002</u>	Sep 06, 2012
<u>AP</u>		<u>EQ 200MG BASE/VIAL</u>	<u>A200753 003</u>	Sep 06, 2012
<u>AP</u>		<u>EQ 350MG BASE/VIAL</u>	<u>A200855 001</u>	Sep 06, 2012
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A209110 001</u>	Oct 26, 2017
<u>AP</u>	! WEST-WARD PHARMS INT	<u>EQ 200MG BASE/VIAL</u>	<u>A040056 001</u>	May 23, 1995
<u>AP</u>	!	<u>EQ 350MG BASE/VIAL</u>	<u>A040335 001</u>	Apr 20, 2000
		<u>EQ 10MG BASE/ML</u>	<u>A040347 001</u>	Apr 25, 2000

**TABLET; ORAL**

## LEUCOVORIN CALCIUM

<u>AB</u>	BARR	<u>EQ 5MG BASE</u>	<u>A071198 001</u>	Sep 24, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A071199 001</u>	Sep 24, 1987
<u>AB</u>	WEST-WARD PHARMS INT	<u>EQ 5MG BASE</u>	<u>A072733 001</u>	Feb 22, 1993
<u>AB</u>	!	<u>EQ 25MG BASE</u>	<u>A072736 001</u>	Feb 22, 1993
		EQ 10MG BASE	A072734 001	Feb 22, 1993
		EQ 15MG BASE	A072735 001	Feb 22, 1993

## LEUPROLIDE ACETATE

## INJECTABLE; INJECTION

## LEUPROLIDE ACETATE

<u>AP</u>	!	SANDOZ	<u>1MG/0.2ML</u>	<u>A074728</u>	<u>001</u>	Aug 04, 1998
<u>AP</u>		SUN PHARMA GLOBAL	<u>1MG/0.2ML</u>	<u>A078885</u>	<u>001</u>	Mar 09, 2009
<u>AP</u>		TEVA PHARMS USA	<u>1MG/0.2ML</u>	<u>A075471</u>	<u>001</u>	Oct 25, 2000
LUPRON DEPOT						
+!	ABBVIE ENDOCRINE	3.75MG		N020011	002	Oct 26, 1995
	INC					
+!		7.5MG/VIAL		N019732	001	Jan 26, 1989
+!		11.25MG/VIAL		N020708	001	Mar 07, 1997
+!		22.5MG/VIAL		N020517	001	Dec 22, 1995
+!		30MG/VIAL		N020517	002	May 30, 1997
+		45MG/VIAL		N020517	003	Jun 17, 2011
LUPRON DEPOT-PED						
+!	ABBVIE ENDOCRINE	7.5MG/VIAL		N020263	002	Apr 16, 1993
	INC					
+!		11.25MG/VIAL		N020263	005	Jan 21, 1994
+!		11.25MG/VIAL		N020263	007	Aug 15, 2011
+!		15MG/VIAL		N020263	006	Jan 21, 1994
+!		30MG/VIAL		N020263	008	Aug 15, 2011

#### **INJECTABLE; SUBCUTANEOUS**

ELIGARD

+!	TOLMAR THERAP	7.5MG/VIAL	N021343	001	Jan 23, 2002
+!		22.5MG/VIAL	N021379	001	Jul 24, 2002
+!		30MG/VIAL	N021488	001	Feb 13, 2003
+!		45MG/VIAL	N021731	001	Dec 14, 2004

## LEUPROLIDE ACETATE; NORETHINDRONE ACETATE

**INJECTABLE, TABLET; INTRAMUSCULAR, ORAL**

LUPANETA PACK

+ ABBVIE ENDOCRINE 3.75MG/VIAL,N/A;N/A,5MG N203696 001 Dec 14, 2012  
+! 11.25MG/VIAL,N/A;N/A,5MG N203696 002 Dec 14, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-247 (of 436)

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

LEVALBUTEROL HYDROCHLORIDE

<u>AN</u>	AUROBINDO PHARMA LTD	<u>EQ 0.25% BASE</u>	<u>A207628 001</u>	Jan 31, 2017
<u>AN</u>		<u>EQ 0.0103% BASE</u>	<u>A207625 001</u>	Dec 30, 2016
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A207625 002</u>	Dec 30, 2016
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A207625 003</u>	Dec 30, 2016
<u>AN</u>	CIPLA LTD	<u>EQ 0.021% BASE</u>	<u>A078171 002</u>	Dec 13, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A078171 003</u>	Dec 13, 2013
<u>AN</u>		<u>EQ 0.0103% BASE</u>	<u>A078171 001</u>	Dec 13, 2013
<u>AN</u>	IMPAX LABS INC	<u>EQ 0.0103% BASE</u>	<u>A077756 003</u>	Apr 09, 2008
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A077756 001</u>	Apr 09, 2008
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A077756 002</u>	Apr 09, 2008
<u>AN</u>	MYLAN SPECIALITY LP	<u>EQ 0.0103% BASE</u>	<u>A077800 001</u>	Mar 15, 2013
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A077800 002</u>	Mar 15, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A077800 003</u>	Mar 15, 2013
<u>AN</u>		<u>EQ 0.25% BASE</u>	<u>A078309 001</u>	Mar 20, 2009
<u>AN</u>	RITEDOSE CORP	<u>EQ 0.0103% BASE</u>	<u>A203653 001</u>	Mar 22, 2016
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A203653 002</u>	Mar 22, 2016
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A203653 003</u>	Mar 22, 2016
<u>AN</u>	TEVA PARENTERAL	<u>EQ 0.25% BASE</u>	<u>A200875 001</u>	Sep 11, 2014
<u>AN</u>	TEVA PHARMS USA	<u>EQ 0.0103% BASE</u>	<u>A090297 001</u>	Apr 26, 2013
<u>AN</u>		<u>EQ 0.021% BASE</u>	<u>A090297 002</u>	Apr 26, 2013
<u>AN</u>		<u>EQ 0.042% BASE</u>	<u>A090297 003</u>	Apr 26, 2013
<b>XOPENEX</b>				
<u>AN</u>	+! OAK PHARMS INC	<u>EQ 0.0103% BASE</u>	<u>N020837 003</u>	Jan 30, 2002
<u>AN</u>	+!	<u>EQ 0.021% BASE</u>	<u>N020837 001</u>	Mar 25, 1999
<u>AN</u>	+!	<u>EQ 0.042% BASE</u>	<u>N020837 002</u>	Mar 25, 1999
<u>AN</u>	+!	<u>EQ 0.25% BASE</u>	<u>N020837 004</u>	Jul 18, 2003

LEVALEBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+! SUNOVION

EQ 0.045MG BASE/INH

N021730 001 Mar 11, 2005

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

KEPPRA

<u>AP</u>	+! UCB INC	<u>500MG/5ML (100MG/ML)</u>	<u>N021872 001</u>	Jul 31, 2006
<b>LEVETIRACETAM</b>				
<u>AP</u>	AUROBINDO PHARMA LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A204312 001</u>	Feb 01, 2016
<u>AP</u>	FRESENIUS KABI USA	<u>500MG/5ML (100MG/ML)</u>	<u>A090813 001</u>	May 26, 2010
<u>AP</u>		<u>500MG/5ML (100MG/ML)</u>	<u>A090876 001</u>	Aug 13, 2015
<u>AP</u>	HIKMA FARMACEUTICA	<u>500MG/5ML (100MG/ML)</u>	<u>A090981 001</u>	Oct 13, 2011
<u>AP</u>	HOSPIRA INC	<u>500MG/5ML (100MG/ML)</u>	<u>A202869 001</u>	Apr 06, 2012
<u>AP</u>	JUBILANT GENERICS	<u>500MG/5ML (100MG/ML)</u>	<u>A206838 001</u>	Jun 02, 2016
<u>AP</u>	LUITPOLD	<u>500MG/5ML (100MG/ML)</u>	<u>A202143 001</u>	Jan 31, 2012
<u>AP</u>	MYLAN LABS LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A203308 001</u>	Sep 16, 2016
<u>AP</u>	SAGENT PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A091627 001</u>	Jun 26, 2013
<u>AP</u>	SUN PHARM INDs LTD	<u>500MG/5ML (100MG/ML)</u>	<u>A090754 001</u>	Jun 16, 2010
<u>AP</u>	X GEN PHARMS	<u>500MG/5ML (100MG/ML)</u>	<u>A091485 001</u>	Aug 05, 2011
<b>LEVETIRACETAM IN SODIUM CHLORIDE</b>				
<u>AP</u>	AUROBINDO PHARMA LTD	<u>500MG/100ML (5MG/ML)</u>	<u>A207160 001</u>	Jan 04, 2017
<u>AP</u>		<u>1000MG/100ML (10MG/ML)</u>	<u>A207160 002</u>	Jan 04, 2017
<u>AP</u>		<u>1500MG/100ML (15MG/ML)</u>	<u>A207160 003</u>	Jan 04, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>500MG/100ML (5MG/ML)</u>	<u>A206880 001</u>	Oct 25, 2017
<u>AP</u>		<u>1000MG/100ML (10MG/ML)</u>	<u>A206880 002</u>	Oct 25, 2017
<u>AP</u>		<u>1500MG/100ML (15MG/ML)</u>	<u>A206880 003</u>	Oct 25, 2017
<u>AP</u>	+! HQ SPECIALITY PHARMA	<u>500MG/100ML (5MG/ML)</u>	<u>N202543 001</u>	Nov 09, 2011
<u>AP</u>	+!	<u>1000MG/100ML (10MG/ML)</u>	<u>N202543 002</u>	Nov 09, 2011
<u>AP</u>	+!	<u>1500MG/100ML (15MG/ML)</u>	<u>N202543 003</u>	Nov 09, 2011

SOLUTION; ORAL

KEPPRA

<u>AA</u>	+! UCB INC	<u>100MG/ML</u>	<u>N021505 001</u>	Jul 15, 2003
<b>LEVETIRACETAM</b>				
<u>AA</u>	ACTAVIS MID ATLANTIC	<u>100MG/ML</u>	<u>A078976 001</u>	Jan 15, 2009
<u>AA</u>	ALLIED PHARMA INC	<u>100MG/ML</u>	<u>A078582 001</u>	Jan 15, 2009
<u>AA</u>	AMNEAL PHARMS	<u>100MG/ML</u>	<u>A090992 001</u>	Oct 27, 2009
<u>AA</u>	AUROBINDO PHARMA	<u>100MG/ML</u>	<u>A079063 001</u>	Jan 15, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-248 (of 436)

LEVETIRACETAM

SOLUTION;ORAL

LEVETIRACETAM

LTD

<u>AA</u>	BRECKENRIDGE PHARM	<u>100MG/ML</u>
<u>AA</u>	HETERO LABS LTD III	<u>100MG/ML</u>
<u>AA</u>	HI-TECH PHARMACAL	<u>100MG/ML</u>
<u>AA</u>	LUPIN LTD	<u>100MG/ML</u>
<u>AA</u>	ORIT LABS LLC	<u>100MG/ML</u>
<u>AA</u>	PHARM ASSOC	<u>100MG/ML</u>
<u>AA</u>	SILARX	<u>100MG/ML</u>
<u>AA</u>	TARO	<u>100MG/ML</u>
<u>AA</u>	TOLMAR	<u>100MG/ML</u>
<u>AA</u>	TRIS PHARMA INC	<u>100MG/ML</u>
<u>AA</u>	VINTAGE PHARMS	<u>100MG/ML</u>
<u>AA</u>	WOCKHARDT BIO AG	<u>100MG/ML</u>

TABLET;ORAL

KEPPRA

<u>AB</u>	+	UCB INC	<u>250MG</u>
<u>AB</u>	+		<u>500MG</u>
<u>AB</u>	+		<u>750MG</u>
<u>AB</u>	+	!	<u>1GM</u>

LEVETIRACETAM

<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	ACI HEALTHCARE LTD	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	ACIC PHARMS	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	AJANTA PHARMA	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	APOTEX INC	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	AUROBINDO PHARMA	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	BRECKENRIDGE PHARM	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	DR REDDYS LABS LTD	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	HETERO LABS LTD III	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	INVAGEN PHARMS	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>	LUPIN	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	MYLAN	<u>250MG</u>
<u>AB</u>		<u>500MG</u>
<u>AB</u>		<u>750MG</u>
<u>AB</u>		<u>1GM</u>
<u>AB</u>	ORCHID HLTHCARE	<u>250MG</u>
<u>AB</u>		<u>500MG</u>

<u>A079120</u>	<u>001</u>	Jan 16, 2009
<u>A203052</u>	<u>001</u>	Feb 28, 2013
<u>A090601</u>	<u>001</u>	Feb 28, 2012
<u>A090893</u>	<u>001</u>	Oct 17, 2011
<u>A203067</u>	<u>001</u>	May 09, 2013
<u>A201157</u>	<u>001</u>	Jun 04, 2015
<u>A090263</u>	<u>001</u>	Apr 03, 2009
<u>A078774</u>	<u>001</u>	Feb 10, 2009
<u>A079107</u>	<u>001</u>	Jan 15, 2009
<u>A090461</u>	<u>001</u>	Sep 30, 2010
<u>A090079</u>	<u>001</u>	Apr 11, 2012
<u>A090028</u>	<u>001</u>	Mar 03, 2010

<u>N021035</u>	<u>001</u>	Nov 30, 1999
<u>N021035</u>	<u>002</u>	Nov 30, 1999
<u>N021035</u>	<u>003</u>	Nov 30, 1999
<u>N021035</u>	<u>004</u>	Jan 06, 2006

<u>A090843</u>	<u>001</u>	Feb 14, 2011
<u>A090843</u>	<u>002</u>	Feb 14, 2011
<u>A090843</u>	<u>003</u>	Feb 14, 2011
<u>A090843</u>	<u>004</u>	Feb 14, 2011
<u>A078042</u>	<u>001</u>	Jan 15, 2009
<u>A078042</u>	<u>002</u>	Jan 15, 2009
<u>A078042</u>	<u>003</u>	Jan 15, 2009
<u>A078042</u>	<u>004</u>	Jan 15, 2009
<u>A090767</u>	<u>001</u>	Jul 28, 2010
<u>A090767</u>	<u>002</u>	Jul 28, 2010
<u>A090767</u>	<u>003</u>	Jul 28, 2010
<u>A090767</u>	<u>004</u>	Jul 28, 2010
<u>A201293</u>	<u>001</u>	Jun 14, 2011
<u>A201293</u>	<u>002</u>	Jun 14, 2011
<u>A201293</u>	<u>003</u>	Jun 14, 2011
<u>A201293</u>	<u>004</u>	Jun 14, 2011
<u>A078869</u>	<u>001</u>	Mar 13, 2009
<u>A078869</u>	<u>002</u>	Mar 13, 2009
<u>A078869</u>	<u>003</u>	Mar 13, 2009
<u>A078869</u>	<u>004</u>	Mar 13, 2009
<u>A078993</u>	<u>001</u>	Jan 15, 2009
<u>A078993</u>	<u>002</u>	Jan 15, 2009
<u>A078993</u>	<u>003</u>	Jan 15, 2009
<u>A078993</u>	<u>004</u>	Jan 15, 2009
<u>A090511</u>	<u>001</u>	Aug 18, 2011
<u>A090511</u>	<u>002</u>	Aug 18, 2011
<u>A090511</u>	<u>003</u>	Aug 18, 2011
<u>A090511</u>	<u>004</u>	Aug 18, 2011
<u>A076920</u>	<u>001</u>	Jan 15, 2009
<u>A076920</u>	<u>002</u>	Jan 15, 2009
<u>A076920</u>	<u>003</u>	Jan 15, 2009
<u>A078904</u>	<u>001</u>	Jan 15, 2009
<u>A090515</u>	<u>001</u>	Oct 08, 2010
<u>A090515</u>	<u>002</u>	Oct 08, 2010
<u>A090515</u>	<u>003</u>	Oct 08, 2010
<u>A090515</u>	<u>004</u>	Oct 08, 2010
<u>A078234</u>	<u>001</u>	Jan 15, 2009
<u>A078234</u>	<u>002</u>	Jan 15, 2009
<u>A078234</u>	<u>003</u>	Jan 15, 2009
<u>A078154</u>	<u>001</u>	Jan 15, 2009
<u>A078154</u>	<u>002</u>	Jan 15, 2009
<u>A078154</u>	<u>003</u>	Jan 15, 2009
<u>A090025</u>	<u>001</u>	Jan 15, 2009
<u>A076919</u>	<u>001</u>	Nov 04, 2008
<u>A076919</u>	<u>002</u>	Nov 04, 2008
<u>A076919</u>	<u>003</u>	Nov 04, 2008
<u>A090261</u>	<u>001</u>	Dec 08, 2009
<u>A078526</u>	<u>001</u>	Jan 15, 2009
<u>A078526</u>	<u>002</u>	Jan 15, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-249 (of 436)

LEVETIRACETAM

TABLET;ORAL

LEVETIRACETAM

<u>AB</u>		<u>750MG</u>	<u>A078526 003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A090484 001</u>	Aug 05, 2010
<u>AB</u>	PRINSTON INC	<u>250MG</u>	<u>A078106 001</u>	Feb 10, 2009
<u>AB</u>		<u>500MG</u>	<u>A078106 002</u>	Feb 10, 2009
<u>AB</u>		<u>750MG</u>	<u>A078106 003</u>	Feb 10, 2009
<u>AB</u>		<u>1GM</u>	<u>A078106 004</u>	Feb 10, 2009
<u>AB</u>	SECAN PHARMS	<u>500MG</u>	<u>A205102 004</u>	Dec 16, 2015
<u>AB</u>		<u>1GM</u>	<u>A205102 003</u>	Dec 16, 2015
<u>AB</u>	TARO	<u>250MG</u>	<u>A078960 004</u>	Feb 01, 2010
<u>AB</u>		<u>500MG</u>	<u>A078960 003</u>	Feb 01, 2010
<u>AB</u>		<u>750MG</u>	<u>A078960 002</u>	Feb 01, 2010
<u>AB</u>		<u>1GM</u>	<u>A078960 001</u>	Feb 01, 2010
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A078101 001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078101 002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078101 003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078101 004</u>	Jan 15, 2009
<u>AB</u>	TORRENT PHARMS	<u>250MG</u>	<u>A078858 001</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A078858 002</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A078858 003</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078858 004</u>	Jan 15, 2009
<u>AB</u>	VINTAGE PHARMS	<u>250MG</u>	<u>A077319 001</u>	Mar 20, 2009
<u>AB</u>		<u>250MG</u>	<u>A091491 001</u>	Dec 14, 2010
<u>AB</u>		<u>500MG</u>	<u>A077319 002</u>	Mar 20, 2009
<u>AB</u>		<u>500MG</u>	<u>A091491 002</u>	Dec 14, 2010
<u>AB</u>		<u>750MG</u>	<u>A091491 003</u>	Dec 14, 2010
<u>AB</u>		<u>750MG</u>	<u>A091491 004</u>	Dec 14, 2010
<u>AB</u>		<u>1GM</u>	<u>A079042 001</u>	Jan 15, 2009
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A079042 002</u>	Jan 15, 2009
<u>AB</u>		<u>500MG</u>	<u>A079042 003</u>	Jan 15, 2009
<u>AB</u>		<u>750MG</u>	<u>A079042 004</u>	Jan 15, 2009
<u>AB</u>		<u>1GM</u>	<u>A078918 001</u>	Apr 29, 2009
<u>AB</u>	ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A078918 002</u>	Apr 29, 2009
<u>AB</u>		<u>1GM</u>		
<u>ROWEEPRA</u>				
<u>AB</u>	LOTUS PHARM CO LTD	<u>250MG</u>	<u>A090906 002</u>	Oct 31, 2016
<u>AB</u>		<u>500MG</u>	<u>A090906 001</u>	Nov 05, 2010
<u>AB</u>		<u>750MG</u>	<u>A090906 003</u>	Oct 31, 2016
<u>AB</u>		<u>1GM</u>	<u>A090906 004</u>	Oct 31, 2016

TABLET, EXTENDED RELEASE;ORAL

KEPPRA XR

<u>AB</u>	+	UCB INC	<u>500MG</u>	<u>N022285 001</u>	Sep 12, 2008
<u>AB</u>	+!		<u>750MG</u>	<u>N022285 002</u>	Feb 12, 2009

LEVETIRACETAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>500MG</u>	<u>A091557 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091557 002</u>	Sep 12, 2011
<u>AB</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A091093 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091093 002</u>	Sep 12, 2011
<u>AB</u>	ANCHEN PHARMS	<u>500MG</u>	<u>A091360 001</u>	Oct 04, 2011
<u>AB</u>		<u>750MG</u>	<u>A091360 002</u>	Oct 04, 2011
<u>AB</u>	APOTEX INC	<u>500MG</u>	<u>A091261 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091261 002</u>	Sep 12, 2011
<u>AB</u>	DEXCEL PHARMA	<u>500MG</u>	<u>A202167 001</u>	Sep 04, 2015
<u>AB</u>		<u>750MG</u>	<u>A202167 002</u>	Sep 04, 2015
<u>AB</u>	ECI PHARMS LLC	<u>500MG</u>	<u>A204754 001</u>	Aug 26, 2016
<u>AB</u>		<u>750MG</u>	<u>A204754 002</u>	Aug 26, 2016
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>500MG</u>	<u>A204511 001</u>	Feb 23, 2016
<u>AB</u>		<u>750MG</u>	<u>A204511 002</u>	Feb 23, 2016
<u>AB</u>	LOTUS PHARM CO LTD	<u>500MG</u>	<u>A202095 002</u>	Jun 06, 2016
<u>AB</u>		<u>750MG</u>	<u>A202095 001</u>	Jun 06, 2016
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A091399 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091399 002</u>	Sep 12, 2011
<u>AB</u>	PHARMADAX INC	<u>500MG</u>	<u>A201464 001</u>	May 25, 2012
<u>AB</u>		<u>750MG</u>	<u>A201464 002</u>	May 25, 2012
<u>AB</u>	PHARMATAK INC	<u>500MG</u>	<u>A207175 001</u>	Sep 28, 2017
<u>AB</u>		<u>750MG</u>	<u>A207175 002</u>	Sep 28, 2017
<u>AB</u>	PRINSTON INC	<u>500MG</u>	<u>A202533 001</u>	Jul 20, 2012
<u>AB</u>		<u>500MG</u>	<u>A203468 001</u>	May 21, 2015
<u>AB</u>		<u>750MG</u>	<u>A202533 002</u>	Jul 20, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-250 (of 436)

LEVETIRACETAM

TABLET, EXTENDED RELEASE;ORAL

LEVETIRACETAM

<u>AB</u>		<u>750MG</u>	<u>A203468 002</u>	May 21, 2015
<u>AB</u>	ROUSES POINT PHARMS	<u>500MG</u>	<u>A202524 001</u>	Aug 27, 2012
<u>AB</u>		<u>750MG</u>	<u>A202524 002</u>	Aug 27, 2012
<u>AB</u>	SUN PHARM INDUSTRIES	<u>500MG</u>	<u>A091285 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091285 002</u>	Sep 12, 2011
<u>AB</u>	SUN PHARMA GLOBAL	<u>500MG</u>	<u>A203059 001</u>	Sep 09, 2013
<u>AB</u>		<u>750MG</u>	<u>A203059 002</u>	Sep 09, 2013
<u>AB</u>	TEVA PHARMS	<u>500MG</u>	<u>A091430 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091430 002</u>	Sep 12, 2011
<u>AB</u>	TORRENT PHARMS LTD	<u>500MG</u>	<u>A091338 001</u>	May 29, 2012
<u>AB</u>		<u>750MG</u>	<u>A091338 002</u>	May 29, 2012
<u>AB</u>	VIRTUS PHARMS	<u>500MG</u>	<u>A091291 001</u>	Sep 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A091291 002</u>	Sep 12, 2011
	APOTEX INC	1GM	A202958 001	Feb 25, 2015
TABLET, FOR SUSPENSION;ORAL				
SPRITAM				
+ APRECIA PHARMS	250MG		N207958 001	Jul 31, 2015
+	500MG		N207958 002	Jul 31, 2015
+	750MG		N207958 003	Jul 31, 2015
++!	1GM		N207958 004	Jul 31, 2015

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

<u>AKBETA</u>				
<u>AT</u>	AKORN	<u>0.25%</u>	<u>A074779 001</u>	Oct 29, 1996
<u>AT</u>		<u>0.5%</u>	<u>A074780 001</u>	Oct 29, 1996
<u>BETAGAN</u>				
<u>AT</u>	+! ALLERGAN	<u>0.25%</u>	<u>N019814 001</u>	Jun 28, 1989
<u>AT</u>	+!	<u>0.5%</u>	<u>N019219 002</u>	Dec 19, 1985
<u>LEVOBUNOLOL HYDROCHLORIDE</u>				
<u>AT</u>	BAUSCH AND LOMB	<u>0.5%</u>	<u>A074326 001</u>	Mar 04, 1994
<u>AT</u>	SANDOZ INC	<u>0.5%</u>	<u>A074850 001</u>	Oct 28, 1996

LEVOCARNITINE

INJECTABLE; INJECTION

<u>CARNITOR</u>				
<u>AP</u>	+! LEADIANT BIOSCI INC	<u>200MG/ML</u>	<u>N020182 001</u>	Dec 16, 1992
<u>LEVOCARNITINE</u>				
<u>AP</u>	LUITPOLD	<u>200MG/ML</u>	<u>A075861 001</u>	Jun 22, 2001
<u>AP</u>	WEST-WARD PHARMS INT	<u>200MG/ML</u>	<u>A075567 001</u>	Mar 29, 2001
SOLUTION;ORAL				
<u>CARNITOR</u>				
<u>AA</u>	+! LEADIANT BIOSCI INC	<u>1GM/10ML</u>	<u>N019257 001</u>	Apr 10, 1986
<u>CARNITOR SF</u>				
<u>AA</u>	+ LEADIANT BIOSCI INC	<u>1GM/10ML</u>	<u>N019257 002</u>	Mar 28, 2007
<u>LEVOCARNITINE</u>				
<u>AA</u>	HI TECH PHARMA	<u>1GM/10ML</u>	<u>A077399 001</u>	Oct 25, 2007
<u>AA</u>	LYNE	<u>1GM/10ML</u>	<u>A076851 001</u>	Aug 10, 2004
TABLET;ORAL				
<u>CARNITOR</u>				
<u>AB</u>	+! LEADIANT BIOSCI INC	<u>330MG</u>	<u>N018948 001</u>	Dec 27, 1985
<u>LEVOCARNITINE</u>				
<u>AB</u>	RISING PHARMS INC	<u>330MG</u>	<u>A076858 001</u>	Sep 20, 2004

LEVOSETIRIZINE DIHYDROCHLORIDE

SOLUTION;ORAL

<u>LEVOSETIRIZINE DIHYDROCHLORIDE</u>				
<u>AA</u>	APOTEX INC	<u>2.5MG/5ML</u>	<u>A202915 001</u>	Aug 21, 2014
<u>AA</u>	L PERRIGO CO	<u>2.5MG/5ML</u>	<u>A091263 001</u>	Nov 07, 2011
<u>AA</u>	SILARX PHARMS INC	<u>2.5MG/5ML</u>	<u>A204599 001</u>	May 15, 2017
<u>AA</u>	TARO PHARM INDNS	<u>2.5MG/5ML</u>	<u>A202673 001</u>	Jul 26, 2013
<u>XYZAL</u>				
<u>AA</u>	+! SANOFI-AVENTIS US	<u>2.5MG/5ML</u>	<u>N0222157 001</u>	Jan 28, 2008
TABLET;ORAL				
<u>LEVOSETIRIZINE DIHYDROCHLORIDE</u>				
<u>AB</u>	ALLIED PHARMA INC	<u>5MG</u>	<u>A204323 001</u>	Dec 20, 2016
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A203027 001</u>	Feb 13, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A090392 001</u>	Feb 24, 2011
<u>AB</u>	GLENMARK GENERICS	<u>5MG</u>	<u>A090385 001</u>	Feb 24, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-251 (of 436)

LEVOCECETIRIZINE DIHYDROCHLORIDE

TABLET;ORAL

LEVOCECETIRIZINE DIHYDROCHLORIDE

<u>AB</u>	HETERO LABS LTD III	<u>5MG</u>	<u>A091264</u> <u>001</u>	Jun 29, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A205564</u> <u>001</u>	Jan 11, 2016
<u>AB</u>	MICRO LABS LTD INDIA	<u>5MG</u>	<u>A202046</u> <u>001</u>	Sep 17, 2013
<u>AB</u>	SCIEGEN PHARMS INC	<u>5MG</u>	<u>A203646</u> <u>001</u>	Sep 09, 2014
<u>AB</u>	SUN PHARM INDs LTD	<u>5MG</u>	<u>A201653</u> <u>001</u>	Jun 26, 2015
<u>AB</u>	SUN PHARMA GLOBAL	<u>5MG</u>	<u>A090362</u> <u>001</u>	Jan 31, 2013
<u>AB</u>	SYNTHON PHARMS	<u>5MG</u>	<u>A090229</u> <u>001</u>	Nov 26, 2010
<u>AB</u>	TEVA PHARMS	<u>5MG</u>	<u>A090199</u> <u>001</u>	Aug 22, 2011
<b><u>XYZAL</u></b>				
<u>AB</u>	+! SANOFI-AVENTIS US	<u>5MG</u>	<u>N022064</u> <u>001</u>	May 25, 2007

LEVOFLOXACIN

INJECTABLE;INJECTION

LEVAQUIN

<u>AP</u>	+! JANSSEN PHARMS	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>N020635</u> <u>001</u>	Dec 20, 1996
<u>AP</u>	+!	<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>N020635</u> <u>004</u>	Dec 20, 1996
<b><u>LEVOFLOXACIN</u></b>				
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A202328</u> <u>001</u>	Jan 24, 2013
<u>AP</u>		<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A202328</u> <u>002</u>	Jan 24, 2013
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A091436</u> <u>001</u>	Jun 05, 2013
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A202590</u> <u>001</u>	Jan 24, 2013
<u>AP</u>		<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A202590</u> <u>002</u>	Jan 24, 2013
<u>AP</u>	SAGENT AGILA LLC	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A200560</u> <u>001</u>	Jun 20, 2011
<u>AP</u>		<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A200560</u> <u>002</u>	Jun 20, 2011
<u>AP</u>	ZYDUS PHARMS USA INC	<u>EQ 500MG/20ML (EQ 25MG/ML)</u>	<u>A205968</u> <u>001</u>	Jun 01, 2017
<u>AP</u>		<u>EQ 750MG/30ML (EQ 25MG/ML)</u>	<u>A205968</u> <u>002</u>	Jun 01, 2017
<b><u>LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u></b>				
<u>AP</u>	! ACS DOBFAR INFO SA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A090343</u> <u>001</u>	Jul 07, 2011
<u>AP</u>	!	<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A090343</u> <u>002</u>	Jul 07, 2011
<u>AP</u>	!	<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A090343</u> <u>003</u>	Jul 07, 2011
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A206919</u> <u>001</u>	Feb 10, 2016
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A206919</u> <u>002</u>	Feb 10, 2016
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A206919</u> <u>003</u>	Feb 10, 2016
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091397</u> <u>001</u>	Aug 08, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091397</u> <u>002</u>	Aug 08, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091397</u> <u>003</u>	Aug 08, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A200674</u> <u>001</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A200674</u> <u>002</u>	Jun 19, 2013
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A200674</u> <u>003</u>	Jun 19, 2013
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A091375</u> <u>001</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A091375</u> <u>002</u>	Sep 16, 2011
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A091375</u> <u>003</u>	Sep 16, 2011
<u>AP</u>	HOSPIRA INC	<u>EQ 250MG/50ML (EQ 5MG/ML)</u>	<u>A078579</u> <u>001</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 500MG/100ML (EQ 5MG/ML)</u>	<u>A078579</u> <u>002</u>	Sep 03, 2015
<u>AP</u>		<u>EQ 750MG/150ML (EQ 5MG/ML)</u>	<u>A078579</u> <u>003</u>	Sep 03, 2015

SOLUTION;ORAL

LEVOFLOXACIN

! HI TECH PHARMA SOLUTION/DROPS;OPHTHALMIC

LEVOFLOXACIN

<u>AT</u>	AKORN	<u>0.5%</u>	<u>A090268</u> <u>001</u>	Dec 20, 2010
<u>AT</u>	MYLAN LABS LTD	<u>0.5%</u>	<u>A204899</u> <u>001</u>	Dec 08, 2017
<u>AT</u>	! RISING PHARMS INC	<u>0.5%</u>	<u>A077700</u> <u>001</u>	Dec 20, 2010
<u>AT</u>	WATSON LABS TEVA	<u>0.5%</u>	<u>A076826</u> <u>001</u>	Feb 10, 2011

TABLET;ORAL

LEVAQUIN

<u>AB</u>	+	JANSSEN PHARMS	<u>250MG</u>	<u>N020634</u> <u>001</u>	Dec 20, 1996
<u>AB</u>	+		<u>500MG</u>	<u>N020634</u> <u>002</u>	Dec 20, 1996
<u>AB</u>	!+		<u>750MG</u>	<u>N020634</u> <u>003</u>	Sep 08, 2000

LEVOFLOXACIN

<u>AB</u>	APOTEX INC	<u>250MG</u>	<u>A090787</u> <u>001</u>	Sep 29, 2011
<u>AB</u>		<u>500MG</u>	<u>A090787</u> <u>002</u>	Sep 29, 2011
<u>AB</u>		<u>750MG</u>	<u>A090787</u> <u>003</u>	Sep 29, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>250MG</u>	<u>A201043</u> <u>001</u>	Jun 20, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-252 (of 436)

LEVOFLOXACIN

TABLET;ORAL

LEVOFLOXACIN

<u>AB</u>		<u>500MG</u>	<u>A201043 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A201043 003</u>	Jun 20, 2011
<u>AB</u>	CIPLA LTD	<u>250MG</u>	<u>A076890 001</u>	Mar 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A076890 002</u>	Mar 30, 2012
<u>AB</u>		<u>750MG</u>	<u>A076890 003</u>	Mar 30, 2012
<u>AB</u>	DR REDDYS LABS INC	<u>250MG</u>	<u>A076710 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076710 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076710 003</u>	Jun 20, 2011
<u>AB</u>	GLENMARK GENERICS	<u>250MG</u>	<u>A200250 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A200250 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A200250 003</u>	Jun 20, 2011
<u>AB</u>	HETERO LABS LTD V	<u>250MG</u>	<u>A202801 001</u>	Jan 08, 2015
<u>AB</u>		<u>500MG</u>	<u>A202801 002</u>	Jan 08, 2015
<u>AB</u>		<u>750MG</u>	<u>A202801 003</u>	Jan 08, 2015
<u>AB</u>	JUBILANT GENERICS	<u>250MG</u>	<u>A203613 001</u>	Jun 19, 2015
<u>AB</u>		<u>500MG</u>	<u>A203613 002</u>	Jun 19, 2015
<u>AB</u>	LUPIN	<u>250MG</u>	<u>A078424 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A078424 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A078424 003</u>	Jun 20, 2011
<u>AB</u>	MACLEODS PHARMS LTD	<u>250MG</u>	<u>A200839 001</u>	Mar 22, 2012
<u>AB</u>		<u>500MG</u>	<u>A200839 002</u>	Mar 22, 2012
<u>AB</u>		<u>750MG</u>	<u>A200839 003</u>	Mar 22, 2012
<u>AB</u>	ORCHID HLTHCARE	<u>250MG</u>	<u>A202200 001</u>	Jan 30, 2012
<u>AB</u>		<u>500MG</u>	<u>A202200 002</u>	Jan 30, 2012
<u>AB</u>		<u>750MG</u>	<u>A202200 003</u>	Jan 30, 2012
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A077438 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A077438 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A077438 003</u>	Jun 20, 2011
<u>AB</u>	TEVA	<u>250MG</u>	<u>A076361 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A076361 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A076361 003</u>	Jun 20, 2011
<u>AB</u>	TORRENT PHARMS	<u>250MG</u>	<u>A090722 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A090722 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A090722 003</u>	Jun 20, 2011
<u>AB</u>	WOCKHARDT	<u>250MG</u>	<u>A090367 001</u>	Jun 20, 2011
<u>AB</u>		<u>500MG</u>	<u>A090367 002</u>	Jun 20, 2011
<u>AB</u>		<u>750MG</u>	<u>A090367 003</u>	Jun 20, 2011
<u>AB</u>	ZYDUS PHARMS USA INC	<u>250MG</u>	<u>A077652 001</u>	Sep 07, 2012
<u>AB</u>		<u>500MG</u>	<u>A077652 002</u>	Sep 07, 2012
<u>AB</u>		<u>750MG</u>	<u>A077652 003</u>	Sep 07, 2012

LEVOLEUCOVORIN CALCIUM

POWDER;IV (INFUSION)

FUSILEV

<u>AP</u>	+!	SPECTRUM PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>N020140 001</u>	Mar 07, 2008
		<u>LEVOLEUCOVORIN CALCIUM</u>			
<u>AP</u>		ACTAVIS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A206516 001</u>	Feb 13, 2017
<u>AP</u>		AMNEAL PHARMS CO	<u>EQ 50MG BASE/VIAL</u>	<u>A207547 001</u>	Feb 13, 2017
<u>AP</u>		WEST-WARD PHARMS INT	<u>EQ 50MG BASE/VIAL</u>	<u>A206263 001</u>	Jun 16, 2016
+!		ACTAVIS LLC SOLUTION;IV (INFUSION)	EQ 175MG BASE/VIAL	N208723 001	Sep 29, 2016
		<u>LEVOLEUCOVORIN CALCIUM</u>			
<u>AP</u>		AMNEAL PHARMS CO	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A207548 001</u>	Sep 08, 2017
<u>AP</u>		MYLAN TEORANTA	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A203576 001</u>	Oct 20, 2015
<u>AP</u>			<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A203576 002</u>	Oct 20, 2015
<u>AP</u>		SANDOZ INC	<u>EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML)</u>	<u>A203563 001</u>	Mar 09, 2015
<u>AP</u>	!		<u>EQ 250MG BASE/25ML (EQ 10MG BASE/ML)</u>	<u>A203563 002</u>	Mar 09, 2015

LEVOMILNACIPRAN HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

FETZIMA

+	FOREST LABS INC	EQ 20MG BASE	N204168 001	Jul 25, 2013
+		EQ 40MG BASE	N204168 002	Jul 25, 2013
+		EQ 80MG BASE	N204168 003	Jul 25, 2013
+		EQ 120MG BASE	N204168 004	Jul 25, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-253 (of 436)

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

SCANDONEST L

! DEPROCO

0.05MG/ML; 2%

A088388 001 Oct 10, 1984

LEVONORGESTREL

INTRAUTERINE DEVICE; INTRAUTERINE

KYLEENA

+! BAYER HLTHCARE 19.5MG

N208224 001 Sep 16, 2016

LILETTA

MEDICINES360 52MG

N206229 001 Feb 26, 2015

MIRENA

+! BAYER HLTHCARE 52MG

N021225 001 Dec 06, 2000

SKYLA

+! BAYER HLTHCARE 13.5MG

N203159 001 Jan 09, 2013

TABLET; ORAL

LEVONORGESTREL

<b>AB</b>	LOTUS PHARM CO LTD	<b>0.75MG</b>	<b>A202684 001</b>	Sep 02, 2016
<b>AB</b>	MYLAN LABS LTD	<b>0.75MG</b>	<b>A202740 001</b>	Sep 02, 2016
<b>AB</b> !	PERRIGO R AND D	<b>0.75MG</b>	<b>A090740 001</b>	Dec 30, 2010

LEVORPHANOL TARTRATE

TABLET; ORAL

LEVORPHANOL TARTRATE

! SENTYNL THERAPS INC 2MG

A074278 001 Mar 31, 2000

LEVOOTHYROXINE SODIUM

CAPSULE; ORAL

TIROSINT

+ INSTITUT

BIOCHIMIQUE

0.013MG

N021924 013 Aug 01, 2007

+ 0.025MG

N021924 002 Oct 13, 2006

+ 0.05MG

N021924 003 Oct 13, 2006

+ 0.075MG

N021924 004 Oct 13, 2006

+ 0.088MG

N021924 010 Oct 02, 2009

+ 0.1MG

N021924 005 Oct 13, 2006

+ 0.112MG

N021924 008 Oct 02, 2009

+ 0.125MG

N021924 006 Oct 13, 2006

+ 0.137MG

N021924 009 Oct 02, 2009

+! 0.15MG

N021924 007 Oct 13, 2006

+ 0.175MG

N021924 011 Apr 25, 2017

+ 0.200MG

N021924 012 Apr 25, 2017

POWDER; INTRAVENOUS

LEVOOTHYROXINE SODIUM

<b>AP</b>	FERA PHARMS LLC	<b>100MCG/VIAL</b>	<b>A206163 001</b>	Jun 29, 2016
<b>AP</b>		<b>500MCG/VIAL</b>	<b>A206163 002</b>	Jun 29, 2016
<b>AP</b> +!	FRESENIUS KABI USA	<b>100MCG/VIAL</b>	<b>N202231 001</b>	Jun 24, 2011
<b>AP</b> +!		<b>200MCG/VIAL</b>	<b>N202231 002</b>	Jun 24, 2011
<b>AP</b> +!		<b>500MCG/VIAL</b>	<b>N202231 003</b>	Jun 24, 2011
<b>AP</b>	PAR STERILE PRODUCTS	<b>200MCG/VIAL</b>	<b>A205366 001</b>	Dec 07, 2015

LEVOOTHYROXINE SODIUM \*\*

\*\*See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET; ORAL

SYNTHROID

--> + ABBVIE	--> <b>AB1, AB2</b>	<b>0.025MG</b>	N021402 001	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.05MG</b>	N021402 002	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.075MG</b>	N021402 003	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.088MG</b>	N021402 004	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.1MG</b>	N021402 005	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.112MG</b>	N021402 006	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.125MG</b>	N021402 007	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.137MG</b>	N021402 008	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.15MG</b>	N021402 009	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.175MG</b>	N021402 010	Jul 24, 2002
--> +	--> <b>AB1, AB2</b>	<b>0.2MG</b>	N021402 012	Jul 24, 2002
--> +!	--> <b>AB1, AB2</b>	<b>0.3MG</b>	N021402 011	Jul 24, 2002

LEVO-T

--> + CEDIPROF INC	--> <b>AB1, AB2, AB3</b>	<b>0.025MG</b>	N021342 001	Mar 01, 2002
--> +	--> <b>AB1, AB2, AB3</b>	<b>0.05MG</b>	N021342 002	Mar 01, 2002
--> +	--> <b>AB1, AB2, AB3</b>	<b>0.075MG</b>	N021342 003	Mar 01, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-254 (of 436)

LEVOTHYROXINE SODIUM \*\*

\*\*See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVO-T

--> +	--> <u>AB1,AB2,AB3</u> <u>0.088MG</u>	N021342 004 Mar 01, 2002
--> +	--> <u>AB1,AB2,AB3</u> <u>0.1MG</u>	N021342 005 Mar 01, 2002
--> +	--> <u>AB1,AB2,AB3</u> <u>0.112MG</u>	N021342 006 Mar 01, 2002
--> +	--> <u>AB1,AB2,AB3</u> <u>0.125MG</u>	N021342 007 Mar 01, 2002
--> +	--> <u>AB1,AB2,AB3</u> <u>0.137MG</u>	N021342 012 Dec 08, 2003
--> +	--> <u>AB1,AB2,AB3</u> <u>0.15MG</u>	N021342 008 Mar 01, 2002
--> +	--> <u>AB1,AB2,AB3</u> <u>0.175MG</u>	N021342 009 Mar 01, 2002
--> +	--> <u>AB1,AB2,AB3</u> <u>0.2MG</u>	N021342 010 Mar 01, 2002
--> +!	--> <u>AB1,AB2,AB3</u> <u>0.3MG</u>	N021342 011 Mar 01, 2002

UNITHROID

--> + STEVENS J	--> <u>AB1,AB2,AB3</u> <u>0.025MG</u>	N021210 001 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.05MG</u>	N021210 002 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.075MG</u>	N021210 003 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.088MG</u>	N021210 004 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.1MG</u>	N021210 005 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.112MG</u>	N021210 006 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.125MG</u>	N021210 007 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.137MG</u>	N021210 012 Feb 08, 2008
--> +	--> <u>AB1,AB2,AB3</u> <u>0.15MG</u>	N021210 008 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.175MG</u>	N021210 009 Aug 21, 2000
--> +	--> <u>AB1,AB2,AB3</u> <u>0.2MG</u>	N021210 010 Aug 21, 2000
--> +!	--> <u>AB1,AB2,AB3</u> <u>0.3MG</u>	N021210 011 Aug 21, 2000

LEVOTHYROXINE SODIUM

--> MYLAN	--> <u>AB1,AB2,AB3,AB4</u> <u>0.025MG</u>	A076187 001 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.05MG</u>	A076187 002 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.075MG</u>	A076187 003 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.088MG</u>	A076187 004 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.1MG</u>	A076187 005 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.112MG</u>	A076187 006 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.125MG</u>	A076187 007 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.137MG</u>	A076187 012 Dec 13, 2006
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.15MG</u>	A076187 008 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.175MG</u>	A076187 009 Jun 05, 2002
-->	--> <u>AB1,AB2,AB3,AB4</u> <u>0.2MG</u>	A076187 010 Jun 05, 2002
--> !	--> <u>AB1,AB2,AB3,AB4</u> <u>0.3MG</u>	A076187 011 Jun 05, 2002

LEVOXYL

--> + KING PHARMS	--> <u>AB1,AB3</u> <u>0.025MG</u>	N021301 001 May 25, 2001
--> +	--> <u>AB1,AB3</u> <u>0.05MG</u>	N021301 002 May 25, 2001
--> +	--> <u>AB1,AB3</u> <u>0.075MG</u>	N021301 003 May 25, 2001
--> +	--> <u>AB1,AB3</u> <u>0.088MG</u>	N021301 004 May 25, 2001
--> +	--> <u>AB1,AB3</u> <u>0.1MG</u>	N021301 005 May 25, 2001
--> +	--> <u>AB1,AB3</u> <u>0.112MG</u>	N021301 006 May 25, 2001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-255 (of 436)

LEVOTHYROXINE SODIUM \*\*

\*\*See current Annual Edition, 1.8 Description of Special Situations, Levothyroxine Sodium

TABLET;ORAL

LEVOXYL

--> +	--> AB1,AB3	0.125MG	N021301 007 May 25, 2001
--> +	--> AB1,AB3	0.137MG	N021301 008 May 25, 2001
--> +	--> AB1,AB3	0.15MG	N021301 009 May 25, 2001
--> +	--> AB1,AB3	0.175MG	N021301 010 May 25, 2001
--> +!	--> AB1,AB3	0.2MG	N021301 011 May 25, 2001

LIDOCAINE

OINTMENT;TOPICAL

LIDOCAINE

<u>AT</u>	ALKEM LABS LTD	<u>5%</u>	<u>A207810 001</u> Mar 10, 2017
<u>AT</u>	AMNEAL PHARMS	<u>5%</u>	<u>A206297 001</u> Aug 07, 2015
<u>AT</u> !	FOUGERA PHARMS INC	<u>5%</u>	<u>A080198 001</u>
<u>AT</u>	GLENMARK PHARMS LTD	<u>5%</u>	<u>A206498 001</u> Sep 09, 2016
<u>AT</u>	RICONPHARMA LLC	<u>5%</u>	<u>A208604 001</u> Sep 20, 2017
<u>AT</u>	SEPTODONT INC	<u>5%</u>	<u>A040911 001</u> May 23, 2011
<u>AT</u>	TARO	<u>5%</u>	<u>A086724 001</u>
<u>AT</u>	TELIGENT PHARMA INC	<u>5%</u>	<u>A205318 001</u> Feb 01, 2016
<u>AT</u>	VITRUVIAS THERAP	<u>5%</u>	<u>A208822 001</u> Sep 25, 2017

PATCH;TOPICAL

LIDOCAINE

<u>AB</u>	ACTAVIS LABS UT INC	<u>5%</u>	<u>A200675 001</u> Aug 23, 2012
<u>AB</u>	MYLAN TECHNOLOGIES	<u>5%</u>	<u>A202346 001</u> Aug 07, 2015

LIDODERM

<u>AB</u> +!	TEIKOKU PHARMA USA	<u>5%</u>
--------------	--------------------	-----------

N020612 001 Mar 19, 1999

LIDOCAINE HYDROCHLORIDE

GEL;OPHTHALMIC

AKTEN	+! AKORN	3.5%	N022221 001 Oct 07, 2008
-------	----------	------	--------------------------

INJECTABLE;INJECTION

LIDOCAINE HYDROCHLORIDE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>1%</u>	<u>A207182 001</u> Oct 30, 2017
<u>AP</u>		<u>2%</u>	<u>A207182 002</u> Oct 30, 2017
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088328 001</u> May 17, 1984
<u>AP</u>		<u>1%</u>	<u>A083158 001</u>
<u>AP</u>		<u>1%</u>	<u>A088329 001</u> May 17, 1984
<u>AP</u>		<u>2%</u>	<u>A040078 001</u> Jun 23, 1995
<u>AP</u>		<u>2%</u>	<u>A083158 002</u>
<u>AP</u>		<u>2%</u>	<u>A088294 001</u> May 17, 1984
<u>AP</u>		<u>20%</u>	<u>A083158 003</u>
<u>AP</u>	INTL MEDICATION	<u>1%</u>	<u>A083173 001</u>
<u>AP</u>		<u>2%</u>	<u>A083173 002</u>
<u>AP</u>	LUITPOLD	<u>1%</u>	<u>A080850 001</u>
<u>AP</u>	LUITPOLD PHARMS INC	<u>1%</u>	<u>A091564 001</u> Aug 14, 2015

LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>200MG/100ML</u>	<u>N019830 002</u> Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML</u>	<u>N018461 002</u>

LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>400MG/100ML</u>	<u>N019830 003</u> Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>400MG/100ML</u>	<u>N018461 003</u>

LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>800MG/100ML</u>	<u>N019830 004</u> Apr 08, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>800MG/100ML</u>	<u>N018461 004</u> Feb 22, 1982

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A088586 001</u> Jul 24, 1985
<u>AP</u>	HOSPIRA	<u>0.5%</u>	<u>A088325 001</u> Jul 31, 1984
<u>AP</u>		<u>1%</u>	<u>A088299 001</u> Jul 31, 1984
<u>AP</u>		<u>2%</u>	<u>A088327 001</u> Jul 31, 1984

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A203040 001</u> Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>A203082 001</u> Mar 14, 2013

<u>AP</u>	FRESENIUS KABI USA	<u>2%</u>	<u>A203040 002</u> Mar 14, 2013
<u>AP</u>		<u>2%</u>	<u>N017584 001</u>

<u>AP</u>	FRESENIUS KABI USA	<u>4%</u>	<u>N017584 002</u>
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>A080408 001</u>



38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-257 (of 436)

LIDOCAINE; TETRACAIN

CREAM;TOPICAL			
PLIAGLIS			
+!	TARO PHARMS	7%;7%	N021717 001 Jun 29, 2006
PATCH;TOPICAL			
SYNERA			
+!	GALEN SPECIALTY	70MG;70MG	N021623 001 Jun 23, 2005

LIFITEGRAST

SOLUTION/DROPS;OPHTHALMIC			
XXIDRA			
+!	SHIRE DEV LLC	5%	N208073 001 Jul 11, 2016

LINACLOTIDE

CAPSULE;ORAL			
LINZESS			
+ FOREST LABS LLC	72MCG	N202811 003 Jan 25, 2017	
+	145MCG	N202811 001 Aug 30, 2012	
+!	290MCG	N202811 002 Aug 30, 2012	

LINAGLIPTIN

TABLET;ORAL			
TRADJENTA			
+! BOEHRINGER	5MG	N201280 001 May 02, 2011	
INGELHEIM			

LINAGLIPTIN; METFORMIN HYDROCHLORIDE

TABLET;ORAL			
JENTADUETO			
+ BOEHRINGER	2.5MG;500MG	N201281 001 Jan 30, 2012	
INGELHEIM			
+	2.5MG;850MG	N201281 002 Jan 30, 2012	
+!	2.5MG;1GM	N201281 003 Jan 30, 2012	
TABLET, EXTENDED RELEASE;ORAL			
JENTADUETO XR			
+ BOEHRINGER	2.5MG;1GM	N208026 001 May 27, 2016	
INGELHEIM			
+!	5MG;1GM	N208026 002 May 27, 2016	

LINCOMYCIN HYDROCHLORIDE

INJECTABLE;INJECTION

LINCOCIN

<u>AP</u> +! PHARMACIA AND	<u>EQ 300MG BASE/ML</u>	<u>N050317 001</u>
UPJOHN		

LINCOMYCIN

<u>AP</u> X-GEN PHARMS INC	<u>EQ 300MG BASE/ML</u>	<u>A201746 001</u> Jun 04, 2015
----------------------------	-------------------------	---------------------------------

LINDANE

LOTION;TOPICAL

LINDANE

<u>AT</u> OLTA PHARMS	<u>1%</u>	<u>A087313 001</u>
<u>AT</u> ! WOCKHARDT	<u>1%</u>	<u>A088190 001</u> Aug 16, 1984
SHAMPOO;TOPICAL		

LINDANE

<u>AT</u> OLTA PHARMS	<u>1%</u>	<u>A087266 001</u>
<u>AT</u> ! WOCKHARDT BIO AG	<u>1%</u>	<u>A088191 001</u> Sep 18, 1984

LINEZOLID

FOR SUSPENSION;ORAL

LINEZOLID

<u>AB</u> WEST-WARD PHARMS	<u>100MG/5ML</u>	<u>A200068 001</u> Jun 03, 2015
INT		

ZYVOX

<u>AB</u> +! PHARMACIA AND	<u>100MG/5ML</u>	<u>N021132 001</u> Apr 18, 2000
UPJOHN		

SOLUTION;IV (INFUSION)

LINEZOLID

<u>AP</u> AUROBINDO PHARMA	<u>600MG/300ML (2MG/ML)</u>	<u>A206917 001</u> Aug 04, 2016
LTD		
<u>AP</u> FRESENIUS KABI USA	<u>600MG/300ML (2MG/ML)</u>	<u>A204764 001</u> Mar 15, 2016
	<u>600MG/300ML (2MG/ML)</u>	<u>A205442 001</u> Jul 07, 2015
<u>AP</u> HOSPIRA INC	<u>200MG/100ML (2MG/ML)</u>	<u>A207001 001</u> Jul 07, 2017
	<u>600MG/300ML (2MG/ML)</u>	<u>A207001 002</u> Jul 07, 2017
<u>AP</u> HQ SPCLT PHARMA	<u>200MG/100ML (2MG/ML)</u>	<u>A205154 001</u> Dec 06, 2017
	<u>600MG/300ML (2MG/ML)</u>	<u>A205154 002</u> Dec 06, 2017
<u>AP</u> MYLAN LABS LTD	<u>200MG/100ML (2MG/ML)</u>	<u>A207354 001</u> Dec 20, 2016
	<u>600MG/300ML (2MG/ML)</u>	<u>A207354 002</u> Dec 20, 2016
<u>AP</u> NANG KUANG PHARM CO	<u>200MG/100ML (2MG/ML)</u>	
	<u>600MG/300ML (2MG/ML)</u>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-258 (of 436)

LINEZOLID

SOLUTION;IV (INFUSION)

LINEZOLID

<u>AP</u>	SAGENT PHARMS	<u>200MG/100ML (2MG/ML)</u>	<u>A204696 001</u>	Mar 02, 2017
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A204696 002</u>	Mar 02, 2017
<u>AP</u>	SANDOZ INC	<u>200MG/100ML (2MG/ML)</u>	<u>A200904 001</u>	Jul 16, 2015
<u>AP</u>		<u>600MG/300ML (2MG/ML)</u>	<u>A200904 002</u>	Jul 16, 2015
<u>AP</u>	TEVA PHARMS	<u>600MG/300ML (2MG/ML)</u>	<u>A200222 001</u>	Jun 27, 2012
	<b>ZYVOX</b>			
<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>200MG/100ML (2MG/ML)</u>	<u>N021131 001</u>	Apr 18, 2000
<u>AP</u>	+! LINEZOLID IN SODIUM CHLORIDE	<u>600MG/300ML (2MG/ML)</u>	<u>N021131 003</u>	Apr 18, 2000
	+! HOSPIRA INC	0.9% IN PLASTIC CONTAINER		
		600MG/300ML (2MG/ML)		
	ZYVOX			
	+ PHARMACIA AND UPJOHN	400MG/200ML (2MG/ML)		
			N021131 002	Apr 18, 2000

TABLET;ORAL

LINEZOLID

<u>AB</u>	ALEMBIC PHARMS LTD	<u>600MG</u>	<u>A205233 001</u>	Dec 21, 2015
<u>AB</u>	ALKEM LABS LTD	<u>600MG</u>	<u>A205517 001</u>	Dec 21, 2015
<u>AB</u>	AMNEAL PHARMS	<u>600MG</u>	<u>A204536 001</u>	Dec 21, 2015
<u>AB</u>	GATE PHARMS	<u>600MG</u>	<u>A091210 001</u>	Feb 05, 2016
<u>AB</u>	GLENMARK PHARMS	<u>600MG</u>	<u>A078987 001</u>	Dec 21, 2015
<u>AB</u>	HETERO LABS LTD V	<u>600MG</u>	<u>A204239 001</u>	Dec 21, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>600MG</u>	<u>A078845 001</u>	Dec 21, 2015
<u>AB</u>	NOVEL LABS INC	<u>600MG</u>	<u>A207526 001</u>	Aug 22, 2016
<u>AB</u>	TEVA PHARMS USA	<u>600MG</u>	<u>A078061 001</u>	May 18, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>600MG</u>	<u>A206097 001</u>	Feb 22, 2017
	<b>ZYVOX</b>			
<u>AB</u>	+! PHARMACIA AND UPJOHN	<u>600MG</u>	<u>N021130 002</u>	Apr 18, 2000

LIOTHYRONINE SODIUM

INJECTABLE;INJECTION

LIOTHYRONINE SODIUM

<u>AP</u>	X GEN PHARMS	<u>EQ 0.01MG BASE/ML</u>	<u>A076923 001</u>	Aug 17, 2005
<u>AP</u>	+! PAR STERILE PRODUCTS	<u>EQ 0.01MG BASE/ML</u>	<u>N020105 001</u>	Dec 31, 1991

TABLET;ORAL

CYTOMEL

<u>AB</u>	+ KING PHARMS	<u>EQ 0.005MG BASE</u>	<u>N010379 001</u>	
<u>AB</u>	+	<u>EQ 0.025MG BASE</u>	<u>N010379 002</u>	
<u>AB</u>	+!	<u>EQ 0.05MG BASE</u>	<u>N010379 003</u>	

LIOTHYRONINE SODIUM

<u>AB</u>	MAYNE PHARMA INC	<u>EQ 0.005MG BASE</u>	<u>A090097 001</u>	Mar 20, 2009
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A090097 002</u>	Mar 20, 2009
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A090097 003</u>	Mar 20, 2009
<u>AB</u>	MYLAN	<u>EQ 0.005MG BASE</u>	<u>A090326 001</u>	Jul 14, 2009
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A090326 002</u>	Jul 14, 2009
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A090326 003</u>	Jul 14, 2009
<u>AB</u>	SIGMAPHARM LABS LLC	<u>EQ 0.005MG BASE</u>	<u>A200295 001</u>	Nov 29, 2012
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A200295 002</u>	Nov 29, 2012
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A200295 003</u>	Nov 29, 2012
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 0.005MG BASE</u>	<u>A091382 001</u>	Apr 20, 2016
<u>AB</u>		<u>EQ 0.025MG BASE</u>	<u>A091382 002</u>	Apr 20, 2016
<u>AB</u>		<u>EQ 0.05MG BASE</u>	<u>A091382 003</u>	Apr 20, 2016

LIOTRIX (T4;T3)

TABLET;ORAL

THYROLAR-0.25

+ FOREST LABS 0.0125MG;0.0031MG N016807 001

THYROLAR-0.5

+ FOREST LABS 0.025MG;0.0063MG N016807 005

THYROLAR-1

+ FOREST LABS 0.05MG;0.0125MG N016807 004

THYROLAR-2

+ FOREST LABS 0.1MG;0.025MG N016807 002

THYROLAR-3

+! FOREST LABS 0.15MG;0.0375MG N016807 003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-259 (of 436)

**LIRAGLUTIDE RECOMBINANT**

SOLUTION;SUBCUTANEOUS

SAXENDA

+!	NOVO NORDISK INC	18MG/3ML (6MG/ML)	N206321 001 Dec 23, 2014
+!	NOVO NORDISK INC	18MG/3ML (6MG/ML)	N022341 001 Jan 25, 2010

**LISDEXAMFETAMINE Dimesylate**

CAPSULE;ORAL

VYVANSE

+	SHIRE DEVELOPMENT	10MG	N021977 007 Oct 30, 2014
+		20MG	N021977 004 Dec 10, 2007
+		30MG	N021977 001 Feb 23, 2007
+		40MG	N021977 005 Dec 10, 2007
+		50MG	N021977 002 Feb 23, 2007
+		60MG	N021977 006 Dec 10, 2007
+!		70MG	N021977 003 Feb 23, 2007

TABLET, CHEWABLE;ORAL

VYVANSE

+	SHIRE DEV LLC	10MG	N208510 001 Jan 28, 2017
+		20MG	N208510 002 Jan 28, 2017
+		30MG	N208510 003 Jan 28, 2017
+		40MG	N208510 004 Jan 28, 2017
+		50MG	N208510 005 Jan 28, 2017
+!		60MG	N208510 006 Jan 28, 2017

**LISINOPRIL**

SOLUTION;ORAL

QBRELIS

+!	SILVERGATE PHARMS	1MG/ML	N208401 001 Jul 29, 2016
----	-------------------	--------	--------------------------

TABLET;ORAL

**LISINOPRIL**

<b><u>AB</u></b>	ACCORD HLTHCARE	<b><u>2.5MG</u></b>	<b><u>A202554 001</u></b> Jul 30, 2013
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A202554 002</u></b> Jul 30, 2013
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A202554 003</u></b> Jul 30, 2013
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A202554 004</u></b> Jul 30, 2013
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A202554 005</u></b> Jul 30, 2013
<b><u>AB</u></b>		<b><u>40MG</u></b>	<b><u>A202554 006</u></b> Jul 30, 2013
<b><u>AB</u></b>	APOTEX INC	<b><u>2.5MG</u></b>	<b><u>A076102 001</u></b> Sep 30, 2002
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A076102 002</u></b> Sep 30, 2002
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A076102 003</u></b> Sep 30, 2002
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A076102 004</u></b> Sep 30, 2002
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A076102 005</u></b> Sep 30, 2002
<b><u>AB</u></b>		<b><u>40MG</u></b>	<b><u>A076102 006</u></b> Sep 30, 2002
<b><u>AB</u></b>	AUROBINDO	<b><u>2.5MG</u></b>	<b><u>A077622 001</u></b> Feb 22, 2006
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A077622 002</u></b> Feb 22, 2006
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A077622 003</u></b> Feb 22, 2006
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A077622 004</u></b> Feb 22, 2006
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A077622 005</u></b> Feb 22, 2006
<b><u>AB</u></b>		<b><u>40MG</u></b>	<b><u>A077622 006</u></b> Feb 22, 2006
<b><u>AB</u></b>	HIKMA INTL PHARMS	<b><u>2.5MG</u></b>	<b><u>A076063 001</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A076063 002</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A076063 003</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A076063 004</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A076063 006</u></b> Jun 27, 2003
<b><u>AB</u></b>		<b><u>40MG</u></b>	<b><u>A076063 005</u></b> Jul 01, 2002
<b><u>AB</u></b>	INVAGEN PHARMS	<b><u>2.5MG</u></b>	<b><u>A203508 001</u></b> Oct 29, 2013
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A203508 002</u></b> Oct 29, 2013
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A203508 003</u></b> Oct 29, 2013
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A203508 004</u></b> Oct 29, 2013
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A203508 005</u></b> Oct 29, 2013
<b><u>AB</u></b>		<b><u>40MG</u></b>	<b><u>A203508 006</u></b> Oct 29, 2013
<b><u>AB</u></b>	IVAX SUB TEVA PHARMS	<b><u>2.5MG</u></b>	<b><u>A075752 001</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A075752 002</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A075752 003</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A075752 004</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A075752 005</u></b> Jul 01, 2002
<b><u>AB</u></b>		<b><u>40MG</u></b>	<b><u>A075752 006</u></b> Jul 01, 2002
<b><u>AB</u></b>	LUPIN	<b><u>2.5MG</u></b>	<b><u>A077321 001</u></b> Sep 09, 2005
<b><u>AB</u></b>		<b><u>5MG</u></b>	<b><u>A077321 002</u></b> Sep 09, 2005
<b><u>AB</u></b>		<b><u>10MG</u></b>	<b><u>A077321 003</u></b> Sep 09, 2005
<b><u>AB</u></b>		<b><u>20MG</u></b>	<b><u>A077321 004</u></b> Sep 09, 2005
<b><u>AB</u></b>		<b><u>30MG</u></b>	<b><u>A077321 005</u></b> Sep 09, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-260 (of 436)

LISINOPRIL

CAPSULE; ORAL

LISINOPRIL

<u>AB</u>		<u>40MG</u>	<u>A077321</u>	<u>006</u>	Sep 09, 2005
<u>AB</u>	MYLAN	<u>2 . 5MG</u>	<u>A076071</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076071</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076071</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076071</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076071</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A076071</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	PRINSTON INC	<u>2 . 5MG</u>	<u>A075743</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>2 . 5MG</u>	<u>A076180</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075743</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076180</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075743</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076180</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075743</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076164</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075743</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076164</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075743</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A076164</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>2 . 5MG</u>	<u>A075994</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075994</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075994</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075994</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075994</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A075994</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	SUN PHARM INDs LTD	<u>2 . 5MG</u>	<u>A075944</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A075944</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A075944</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A075944</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A075944</u>	<u>006</u>	Feb 11, 2003
<u>AB</u>		<u>40MG</u>	<u>A075944</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>2 . 5MG</u>	<u>A076059</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>A076059</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>A076059</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>A076059</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>A076059</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>A076059</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	WOCKHARDT	<u>2 . 5MG</u>	<u>A078402</u>	<u>001</u>	Apr 19, 2007
<u>AB</u>		<u>5MG</u>	<u>A078402</u>	<u>002</u>	Apr 19, 2007
<u>AB</u>		<u>10MG</u>	<u>A078402</u>	<u>003</u>	Apr 19, 2007
<u>AB</u>		<u>20MG</u>	<u>A078402</u>	<u>004</u>	Apr 19, 2007
<u>AB</u>		<u>30MG</u>	<u>A078402</u>	<u>005</u>	Apr 19, 2007
<u>AB</u>		<u>40MG</u>	<u>A078402</u>	<u>006</u>	Apr 19, 2007
<b>PRINIVIL</b>					
<u>AB</u>	MERCK	<u>5MG</u>	<u>N019558</u>	<u>001</u>	Dec 29, 1987
<u>AB</u>		<u>10MG</u>	<u>N019558</u>	<u>002</u>	Dec 29, 1987
<u>AB</u>		<u>20MG</u>	<u>N019558</u>	<u>003</u>	Dec 29, 1987
<u>AB</u>		<u>40MG</u>	<u>N019558</u>	<u>004</u>	Oct 25, 1988
<b>ZESTRIL</b>					
<u>AB</u>	+ ALVOGEN MALTA	<u>2 . 5MG</u>	<u>N019777</u>	<u>005</u>	Apr 29, 1993
<u>AB</u>	+	<u>5MG</u>	<u>N019777</u>	<u>001</u>	May 19, 1988
<u>AB</u>	+	<u>10MG</u>	<u>N019777</u>	<u>002</u>	May 19, 1988
<u>AB</u>	+	<u>20MG</u>	<u>N019777</u>	<u>003</u>	May 19, 1988
<u>AB</u>	+	<u>30MG</u>	<u>N019777</u>	<u>006</u>	Jan 20, 1999
<u>AB</u>	!+	<u>40MG</u>	<u>N019777</u>	<u>004</u>	May 19, 1988
<b>LITHIUM CARBONATE</b>					
CAPSULE; ORAL					
<b>LITHIUM CARBONATE</b>					
<u>AB</u>	ALEMBIC LTD	<u>150MG</u>	<u>A079159</u>	<u>001</u>	Jan 12, 2009
<u>AB</u>		<u>300MG</u>	<u>A079159</u>	<u>002</u>	Jan 12, 2009
<u>AB</u>		<u>600MG</u>	<u>A079159</u>	<u>003</u>	Jan 12, 2009
<u>AB</u>	GLENMARK GENERICS	<u>150MG</u>	<u>A079139</u>	<u>001</u>	Feb 03, 2009
<u>AB</u>		<u>300MG</u>	<u>A079139</u>	<u>002</u>	Feb 03, 2009
<u>AB</u>		<u>600MG</u>	<u>A079139</u>	<u>003</u>	Feb 03, 2009
<u>AB</u>	HETERO LABS LTD III	<u>150MG</u>	<u>A090702</u>	<u>001</u>	Sep 25, 2009
<u>AB</u>		<u>300MG</u>	<u>A090702</u>	<u>002</u>	Sep 25, 2009
<u>AB</u>		<u>600MG</u>	<u>A090702</u>	<u>003</u>	Sep 25, 2009
<u>AB</u>	MYLAN PHARMS INC	<u>150MG</u>	<u>A076243</u>	<u>002</u>	Feb 24, 2003
<u>AB</u>		<u>300MG</u>	<u>A076243</u>	<u>001</u>	Jun 27, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-261 (of 436)

LITHIUM CARBONATE

CAPSULE;ORAL

LITHIUM CARBONATE

<u>AB</u>		<u>600MG</u>	<u>A078763 001</u>	Apr 15, 2008
<u>AB</u>	+ WEST-WARD PHARMS INT	<u>150MG</u>	<u>N017812 002</u>	Jan 28, 1987
<u>AB</u>	+	<u>300MG</u>	<u>N017812 001</u>	
<u>AB</u>	+!	<u>600MG</u>	<u>N017812 003</u>	Jan 28, 1987

TABLET;ORAL

LITHIUM CARBONATE

<u>AB</u>	SUN PHARM INDs INC	<u>300MG</u>	<u>A091027 001</u>	Jun 24, 2010
<u>AB</u>	+! WEST-WARD PHARMS INT	<u>300MG</u>	<u>N018558 001</u>	Jan 29, 1982

TABLET, EXTENDED RELEASE;ORAL

LITHIUM CARBONATE

<u>AB</u>	ALEMBIC PHARMS LTD	<u>300MG</u>	<u>A204445 001</u>	Jun 10, 2015
<u>AB</u>	GLENMARK GENERICS	<u>450MG</u>	<u>A091616 001</u>	Feb 14, 2011
<u>AB</u>	GLENMARK PHARMS INC	<u>300MG</u>	<u>A091544 001</u>	Dec 27, 2010
<u>AB</u>	HERITAGE PHARMA	<u>300MG</u>	<u>A205532 001</u>	Sep 29, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>300MG</u>	<u>A202288 001</u>	Jun 29, 2012
<u>AB</u>		<u>450MG</u>	<u>A202219 001</u>	Aug 08, 2012
<u>AB</u>	UNIQUE PHARM LABS	<u>300MG</u>	<u>A204779 001</u>	Jul 27, 2015
<u>AB</u>		<u>450MG</u>	<u>A205663 001</u>	Jun 05, 2017
<u>AB</u>	WEST-WARD PHARMS INT	<u>300MG</u>	<u>A076832 001</u>	Oct 28, 2004
<u>AB</u>	!	<u>450MG</u>	<u>A076691 001</u>	Jan 05, 2004

LITHOBID

<u>AB</u>	+! ANI PHARMS INC	<u>300MG</u>	<u>N018027 001</u>
-----------	-------------------	--------------	--------------------

LITHIUM CITRATE

SYRUP;ORAL

LITHIUM CITRATE

<u>AA</u>	+! WEST-WARD PHARMS INT	<u>EQ 300MG CARBONATE/5ML</u>	<u>N018421 001</u>
<u>AA</u>	WOCKHARDT BIO AG	<u>EQ 300MG CARBONATE/5ML</u>	<u>A070755 001</u> May 21, 1986

LIXISENATIDE

SOLUTION;SUBCUTANEOUS

ADLYXIN

+!	SANOFI-AVENTIS US	0.15MG/3ML (0.05MG/ML)
+!		0.3MG/3ML (0.1MG/ML)

N208471 001 Jul 27, 2016  
 N208471 002 Jul 27, 2016

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS;OPHTHALMIC ALOMIDE

+!	NOVARTIS PHARMS CORP	EQ 0.1% BASE	N020191 001 Sep 23, 1993
----	----------------------	--------------	--------------------------

LOMITAPIDE MESYLATE

CAPSULE;ORAL

JUXTAPID

+	AEGERION	EQ 5MG BASE	N203858 001 Dec 21, 2012
+		EQ 10MG BASE	N203858 002 Dec 21, 2012
+		EQ 20MG BASE	N203858 003 Dec 21, 2012
+		EQ 30MG BASE	N203858 004 Apr 23, 2015
+		EQ 40MG BASE	N203858 005 Apr 23, 2015
+		EQ 60MG BASE	N203858 006 Apr 23, 2015

LOMUSTINE

CAPSULE;ORAL

GLEOSTINE

+	CORDEN PHARMA	5MG	N017588 004 Dec 19, 2014
+		10MG	N017588 001
+		40MG	N017588 002
+		100MG	N017588 003

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL

LOPERAMIDE HYDROCHLORIDE

<u>AB</u>	! MYLAN	<u>2MG</u>	<u>A072741 001</u>	Sep 18, 1991
<u>AB</u>	TEVA	<u>2MG</u>	<u>A073192 001</u>	Apr 30, 1992

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-262 (of 436)

LOPINAVIR; RITONAVIR

SOLUTION;ORAL

KALETRA

<u>AA</u>	+!	ABBVIE	<u>80MG/ML;20MG/ML</u>	<u>N021251</u>	<u>001</u>	Sep 15, 2000
<u>AA</u>		<u>LOPINAVIR AND RITONAVIR</u>				
<u>AA</u>		SILARX PHARMS INC	<u>80MG/ML;20MG/ML</u>	<u>A207407</u>	<u>001</u>	Dec 27, 2016
		TABLET;ORAL				
		KALETRA				
	+	ABBVIE	100MG;25MG	N021906	002	Nov 09, 2007
	+		200MG;50MG	N021906	001	Oct 28, 2005

LORAZEPAM

CONCENTRATE;ORAL

LORAZEPAM

<u>AA</u>		AMNEAL PHARMS	<u>2MG/ML</u>	<u>A091383</u>	<u>001</u>	Dec 23, 2009
<u>AA</u>		HI-TECH PHARMA CO	<u>2MG/ML</u>	<u>A200169</u>	<u>001</u>	Jan 30, 2012
<u>AA</u>		LUPIN LTD	<u>2MG/ML</u>	<u>A091407</u>	<u>001</u>	Feb 19, 2013
<u>AA</u>		PHARM ASSOC	<u>2MG/ML</u>	<u>A090260</u>	<u>001</u>	Jun 15, 2010
<u>AA</u>		SAPITALIS PHARMS	<u>2MG/ML</u>	<u>A079244</u>	<u>001</u>	Apr 28, 2009

LORAZEPAM INTENSOL

<u>AA</u>	!	WEST-WARD PHARMS INT	<u>2MG/ML</u>	<u>A072755</u>	<u>001</u>	Jun 28, 1991
-----------	---	----------------------	---------------	----------------	------------	--------------

INJECTABLE;INJECTION

ATIVAN

<u>AP</u>	+	WEST-WARD PHARMS INT	<u>2MG/ML</u>	<u>N018140</u>	<u>001</u>	
<u>AP</u>	+		<u>4MG/ML</u>	<u>N018140</u>	<u>002</u>	

LORAZEPAM

<u>AP</u>		AKORN	<u>2MG/ML</u>	<u>A075025</u>	<u>001</u>	Jul 23, 1998
<u>AP</u>		HOSPIRA	<u>2MG/ML</u>	<u>A074243</u>	<u>001</u>	Apr 12, 1994
<u>AP</u>			<u>2MG/ML</u>	<u>A074282</u>	<u>001</u>	May 27, 1994
<u>AP</u>			<u>4MG/ML</u>	<u>A074243</u>	<u>002</u>	Apr 12, 1994
<u>AP</u>			<u>4MG/ML</u>	<u>A074282</u>	<u>002</u>	May 27, 1994
<u>AP</u>		INTL MEDICATION SYS	<u>2MG/ML</u>	<u>A076150</u>	<u>001</u>	Nov 15, 2004

LORAZEPAM PRESERVATIVE FREE

<u>AP</u>		BEDFORD LABS	<u>2MG/ML</u>	<u>A077074</u>	<u>001</u>	Jul 13, 2005
<u>AP</u>			<u>4MG/ML</u>	<u>A077074</u>	<u>002</u>	Jul 13, 2005

TABLET;ORAL

ATIVAN

<u>AB</u>	+	VALEANT INTL	<u>0.5MG</u>	<u>N017794</u>	<u>001</u>	
<u>AB</u>	+		<u>1MG</u>	<u>N017794</u>	<u>002</u>	
<u>AB</u>	+		<u>2MG</u>	<u>N017794</u>	<u>003</u>	

LORAZEPAM

<u>AB</u>		AMNEAL PHARMS	<u>0.5MG</u>	<u>A078826</u>	<u>001</u>	Jun 23, 2010
<u>AB</u>			<u>1MG</u>	<u>A078826</u>	<u>002</u>	Jun 23, 2010
<u>AB</u>			<u>2MG</u>	<u>A078826</u>	<u>003</u>	Jun 23, 2010
<u>AB</u>		ANI PHARMS INC	<u>0.5MG</u>	<u>A077396</u>	<u>001</u>	Dec 13, 2006
<u>AB</u>			<u>1MG</u>	<u>A077396</u>	<u>002</u>	Dec 13, 2006
<u>AB</u>			<u>2MG</u>	<u>A077396</u>	<u>003</u>	Dec 13, 2006
<u>AB</u>		AUROLIFE PHARMA LLC	<u>0.5MG</u>	<u>A203572</u>	<u>001</u>	Dec 22, 2017
<u>AB</u>			<u>1MG</u>	<u>A203572</u>	<u>002</u>	Dec 22, 2017
<u>AB</u>			<u>2MG</u>	<u>A203572</u>	<u>003</u>	Dec 22, 2017

LEADING PHARMA LLC

<u>AB</u>		LEADING PHARMA LLC	<u>0.5MG</u>	<u>A078203</u>	<u>001</u>	Jul 30, 2007
<u>AB</u>			<u>1MG</u>	<u>A078203</u>	<u>002</u>	Jul 30, 2007
<u>AB</u>			<u>2MG</u>	<u>A078203</u>	<u>003</u>	Jul 30, 2007
<u>AB</u>		MYLAN	<u>0.5MG</u>	<u>A071589</u>	<u>001</u>	Oct 13, 1987
<u>AB</u>			<u>0.5MG</u>	<u>A077657</u>	<u>001</u>	Mar 16, 2006

SANDOZ

<u>AB</u>		SANDOZ	<u>0.5MG</u>	<u>A071141</u>	<u>002</u>	Apr 21, 1987
<u>AB</u>			<u>1MG</u>	<u>A071141</u>	<u>003</u>	Apr 21, 1987
<u>AB</u>			<u>2MG</u>	<u>A071141</u>	<u>001</u>	Apr 21, 1987
<u>AB</u>		SUN PHARM INDs LTD	<u>0.5MG</u>	<u>A076045</u>	<u>001</u>	Aug 29, 2001
<u>AB</u>			<u>1MG</u>	<u>A076045</u>	<u>002</u>	Aug 29, 2001

VINTAGE PHARMS

<u>AB</u>		VINTAGE PHARMS	<u>0.5MG</u>	<u>A077754</u>	<u>001</u>	May 10, 2006
<u>AB</u>			<u>1MG</u>	<u>A077754</u>	<u>002</u>	May 10, 2006
<u>AB</u>			<u>2MG</u>	<u>A077754</u>	<u>003</u>	May 10, 2006
<u>AB</u>		WATSON LABS	<u>0.5MG</u>	<u>A072926</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>			<u>1MG</u>	<u>A072927</u>	<u>001</u>	Oct 31, 1991

WATSON LABS

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-263 (of 436)

LORCASERIN HYDROCHLORIDE

TABLET;ORAL

BELVIQ

+! EISAI INC 10MG

N022529 001 Jun 27, 2012

TABLET, EXTENDED RELEASE;ORAL

BELVIQ XR

+! EISAI INC 20MG

N208524 001 Jul 15, 2016

LOSARTAN POTASSIUM

TABLET;ORAL

COZAAR

<u>AB</u>	+	MERCK SHARP DOHME	<u>25MG</u>
<u>AB</u>	+		<u>50MG</u>
<u>AB</u>	+		<u>100MG</u>

<u>N020386</u>	<u>001</u>	Apr 14, 1995
<u>N020386</u>	<u>002</u>	Apr 14, 1995
<u>N020386</u>	<u>003</u>	Oct 13, 1998

LOSARTAN POTASSIUM

<u>AB</u>		ALEMBIC PHARMS LTD	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A090428</u>	<u>001</u>	Oct 06, 2010
<u>A090428</u>	<u>002</u>	Oct 06, 2010
<u>A090428</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		APOTEX CORP	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A090790</u>	<u>001</u>	Oct 06, 2010
<u>A090790</u>	<u>002</u>	Oct 06, 2010
<u>A090790</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A090083</u>	<u>001</u>	Oct 06, 2010
<u>A090083</u>	<u>002</u>	Oct 06, 2010
<u>A090083</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		CADISTA PHARMS	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A201170</u>	<u>001</u>	Sep 18, 2012
<u>A201170</u>	<u>002</u>	Sep 18, 2012
<u>A201170</u>	<u>003</u>	Sep 18, 2012

<u>AB</u>		HETERO LABS LTD V	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A203835</u>	<u>001</u>	Aug 12, 2015
<u>A203835</u>	<u>002</u>	Aug 12, 2015
<u>A203835</u>	<u>003</u>	Aug 12, 2015

<u>AB</u>		IPCA LABS LTD	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A200290</u>	<u>001</u>	Aug 30, 2013
<u>A200290</u>	<u>002</u>	Aug 30, 2013
<u>A200290</u>	<u>003</u>	Aug 30, 2013

<u>AB</u>		LUPIN LTD	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A078232</u>	<u>001</u>	Oct 06, 2010
<u>A078232</u>	<u>002</u>	Oct 06, 2010
<u>A078232</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		MACLEODS PHARMS LTD	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A202230</u>	<u>001</u>	May 30, 2012
<u>A202230</u>	<u>002</u>	May 30, 2012
<u>A202230</u>	<u>003</u>	May 30, 2012

<u>AB</u>		MYLAN	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A091590</u>	<u>001</u>	Oct 06, 2010
<u>A091590</u>	<u>002</u>	Oct 06, 2010
<u>A091590</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		PRINSTON INC	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A091497</u>	<u>001</u>	Jun 06, 2011
<u>A091497</u>	<u>002</u>	Jun 06, 2011
<u>A091497</u>	<u>003</u>	Jun 06, 2011

<u>AB</u>		SANDOZ	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A077424</u>	<u>001</u>	Oct 06, 2010
<u>A077424</u>	<u>002</u>	Oct 06, 2010
<u>A077424</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		TEVA	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A076958</u>	<u>001</u>	Apr 06, 2010
<u>A076958</u>	<u>002</u>	Apr 06, 2010
<u>A076958</u>	<u>003</u>	Apr 06, 2010

<u>AB</u>		TORRENT PHARMS	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A090467</u>	<u>001</u>	Oct 06, 2010
<u>A090467</u>	<u>002</u>	Oct 06, 2010
<u>A090467</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		UNICHEM LABS LTD	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A203030</u>	<u>001</u>	Oct 14, 2015
<u>A203030</u>	<u>002</u>	Oct 14, 2015
<u>A203030</u>	<u>003</u>	Oct 14, 2015

<u>AB</u>		UPSHER-SMITH LABS	<u>25MG</u>
<u>AB</u>			<u>50MG</u>
<u>AB</u>			<u>100MG</u>

<u>A090544</u>	<u>001</u>	Oct 06, 2010
<u>A090544</u>	<u>002</u>	Oct 06, 2010
<u>A090544</u>	<u>003</u>	Oct 06, 2010

<u>AB</u>		VIVA HLTHCARE	<u>25MG</u>
<			

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-264 (of 436)

LOSARTAN POTASSIUM

TABLET;ORAL

LOSARTAN POTASSIUM

<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078243 001</u> Oct 06, 2010
<u>AB</u>		<u>50MG</u>	<u>A078243 002</u> Oct 06, 2010
<u>AB</u>		<u>100MG</u>	<u>A078243 003</u> Oct 06, 2010

LOTEPREDNOL ETABONATE

GEL;OPHTHALMIC

LOTEMAX

+! BAUSCH AND LOMB INC 0.5% N202872 001 Sep 28, 2012

OINTMENT;OPHTHALMIC

LOTEMAX

+! BAUSCH AND LOMB 0.5% N200738 001 Apr 15, 2011

SUSPENSION/DROPS;OPHTHALMIC

ALREX

+! BAUSCH AND LOMB 0.2% N020803 001 Mar 09, 1998

LOTEMAX

+! BAUSCH AND LOMB 0.5% N020583 001 Mar 09, 1998

LOTEPREDNOL ETABONATE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC

ZYLET

+! BAUSCH AND LOMB 0.5%;0.3% N050804 001 Dec 14, 2004

LOVASTATIN

TABLET;ORAL

LOVASTATIN

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A075828 001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075828 002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075828 003</u> Dec 17, 2001
<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A077748 001</u> Feb 28, 2007
<u>AB</u>		<u>20MG</u>	<u>A077748 002</u> Feb 28, 2007
<u>AB</u>		<u>40MG</u>	<u>A077748 003</u> Feb 28, 2007
<u>AB</u>	CARLSBAD	<u>10MG</u>	<u>A075991 001</u> Jun 05, 2002
<u>AB</u>		<u>20MG</u>	<u>A075991 002</u> Jun 05, 2002
<u>AB</u>	!	<u>40MG</u>	<u>A075991 003</u> Jun 05, 2002
<u>AB</u>	LUPIN	<u>10MG</u>	<u>A078296 001</u> Mar 14, 2008
<u>AB</u>		<u>20MG</u>	<u>A078296 002</u> Nov 01, 2007
<u>AB</u>		<u>40MG</u>	<u>A078296 003</u> Nov 01, 2007
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A075451 001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075451 002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075451 003</u> Dec 17, 2001
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A075300 001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075636 001</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075300 002</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075636 002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075300 003</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075636 003</u> Dec 17, 2001
<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A077520 001</u> Apr 14, 2006
<u>AB</u>		<u>20MG</u>	<u>A077520 002</u> Apr 14, 2006
<u>AB</u>		<u>40MG</u>	<u>A077520 003</u> Apr 14, 2006
<u>AB</u>	TEVA	<u>10MG</u>	<u>A075551 003</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>A075551 002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>A075551 001</u> Dec 17, 2001

TABLET, EXTENDED RELEASE;ORAL

ALTOPREV

+! COVIS PHARMA BV 20MG N021316 002 Jun 26, 2002

+! COVIS PHARMA BV 40MG N021316 003 Jun 26, 2002

+! COVIS PHARMA BV 60MG N021316 004 Jun 26, 2002

LOXAPINE

POWDER;INHALATION

ADASUVE

+! GALEN UK 10MG N022549 001 Dec 21, 2012

LOXAPINE SUCCINATE

CAPSULE;ORAL

LOXAPINE SUCCINATE

<u>AB</u>	ELITE LABS INC	<u>EQ 5MG BASE</u>	<u>A076868 001</u> Aug 04, 2005
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076868 002</u> Aug 04, 2005
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A076868 003</u> Aug 04, 2005
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076868 004</u> Aug 04, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-265 (of 436)

LOXAPINE SUCCINATE

CAPSULE;ORAL

**LOXAPINE SUCCINATE**

<b>AB</b>	LANNETT HOLDINGS INC	<b>EQ 5MG BASE</b>	<b>A090695 001</b>	Sep 26, 2011
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A090695 002</b>	Sep 26, 2011
<b>AB</b>		<b>EQ 25MG BASE</b>	<b>A090695 003</b>	Sep 26, 2011
<b>AB</b>		<b>EQ 50MG BASE</b>	<b>A090695 004</b>	Sep 26, 2011
<b>AB</b>	MYLAN	<b>EQ 5MG BASE</b>	<b>A076762 001</b>	Nov 01, 2004
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A076762 002</b>	Nov 01, 2004
<b>AB</b>		<b>EQ 25MG BASE</b>	<b>A076762 003</b>	Nov 01, 2004
<b>AB</b>		<b>EQ 50MG BASE</b>	<b>A076762 004</b>	Nov 01, 2004
<b>AB</b>	WATSON LABS	<b>EQ 5MG BASE</b>	<b>A072204 001</b>	Jun 15, 1988
<b>AB</b>		<b>EQ 10MG BASE</b>	<b>A072205 001</b>	Jun 15, 1988
<b>AB</b>	!	<b>EQ 25MG BASE</b>	<b>A072206 001</b>	Jun 15, 1988
<b>AB</b>		<b>EQ 50MG BASE</b>	<b>A072062 001</b>	Jun 15, 1988

LUBIPROSTONE

CAPSULE;ORAL

AMITIZA

+	SUCAMPO PHARMA LLC	8MCG	N021908 002	Apr 29, 2008
+!		24MCG	N021908 001	Jan 31, 2006

LULICONAZOLE

CREAM;TOPICAL

LUZU

+	MEDICIS	1%	N204153 001	Nov 14, 2013
---	---------	----	-------------	--------------

LURASIDONE HYDROCHLORIDE

TABLET;ORAL

LATUDA

+	SUNOVION PHARMS INC	20MG	N200603 003	Dec 07, 2011
+!		40MG	N200603 001	Oct 28, 2010
+		60MG	N200603 005	Jul 12, 2013
+		80MG	N200603 002	Oct 28, 2010
+		120MG	N200603 004	Apr 26, 2012

MACIMORELIN ACETATE

FOR SOLUTION;ORAL

MACRILEN

+	AETERNA ZENTARIS	EQ 60MG BASE/POUCH	N205598 001	Dec 21, 2017
---	------------------	--------------------	-------------	--------------

MACITENTAN

TABLET;ORAL

OPSUMIT

+	ACTELION PHARMS LTD	10MG	N204410 001	Oct 18, 2013
---	---------------------	------	-------------	--------------

MAFENIDE ACETATE

CREAM;TOPICAL

SULFAMYLYON

+	MYLAN INSTITUTIONAL	EQ 85MG BASE/GM	N016763 001
---	---------------------	-----------------	-------------

FOR SOLUTION;TOPICAL

**MAFENIDE ACETATE**

<b>AT</b>	NOVAST LABS LTD	<b>5%</b>	<b>A206716 001</b>	Jul 31, 2017
<b>AT</b>	PAR FORM	<b>5%</b>	<b>A201511 001</b>	Feb 12, 2013

**SULFAMYLYON**

<b>AT</b>	+! MYLAN INSTITUTIONAL	<b>5%</b>	<b>N019832 003</b>	Jun 05, 1998
-----------	------------------------	-----------	--------------------	--------------

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE;INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER				
+! B BRAUN	30MG/100ML;37MG/100ML;0.82MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML;12MG/100ML		N019696 001	Sep 29, 1989

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE;INJECTION

**PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER**

<b>AP</b>	BAXTER HLTHCARE	<b>30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML</b>	<b>N017378 001</b>	
-----------	-----------------	--	--------------------	--

**PLASMA-LYTE A IN PLASTIC CONTAINER**

<b>AP</b>	BAXTER HLTHCARE	<b>30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML</b>	<b>N017378 002</b>	Nov 22, 1982
-----------	-----------------	--	--------------------	--------------

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN	30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML	N019711 001	Sep 29, 1989
---------	---	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-266 (of 436)

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

NORMOSOL-R IN PLASTIC CONTAINER

ICU MEDICAL INC	30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG /100ML; 502MG/100ML	N017586 001
-----------------	--	-------------

SOLUTION; IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

B BRAUN	30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG /100ML; 500MG/100ML	N019024 001	Jun 08, 1984
---------	--	-------------	--------------

PHYSIOSOL IN PLASTIC CONTAINER

ICU MEDICAL INC	30MG/100ML; 37MG/100ML; 222MG/100ML; 526MG /100ML; 502MG/100ML	N017637 002	Jul 08, 1982
-----------------	--	-------------	--------------

MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; INJECTION

NORMOCARB HF 25

+! DIALYSIS SUPS	0.21GM/100ML; 2.8GM/100ML; 9.07GM/100ML	N021910 001	Jul 26, 2006
------------------	---	-------------	--------------

NORMOCARB HF 35

+! DIALYSIS SUPS	0.21GM/100ML; 3.97GM/100ML; 8.3GM/100ML	N021910 002	Jul 26, 2006
------------------	---	-------------	--------------

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	FRESENIUS KABI USA	<u>1GM/100ML</u>	<u>A206486 001</u>	Mar 07, 2016
<u>AP</u>	+! HOSPIRA	<u>1GM/100ML</u>	<u>N020488 001</u>	Jul 11, 1995
<u>AP</u>	HQ SPCLT PHARMA	<u>1GM/100ML</u>	<u>A207349 001</u>	Mar 02, 2016

MAGNESIUM SULFATE IN PLASTIC CONTAINER

<u>AP</u>	FRESENIUS KABI USA	<u>4GM/100ML (40MG/ML)</u>	<u>A206485 001</u>	Mar 15, 2016
<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A206485 002</u>	Mar 15, 2016
<u>AP</u>		<u>2GM/50ML (40MG/ML)</u>	<u>A206485 003</u>	Mar 15, 2016
<u>AP</u>		<u>20GM/500ML (40MG/ML)</u>	<u>A206485 004</u>	Mar 15, 2016
<u>AP</u>		<u>40GM/1000ML (40MG/ML)</u>	<u>A206485 005</u>	Mar 15, 2016
<u>AP</u>	+ HOSPIRA	<u>2GM/50ML (40MG/ML)</u>	<u>N020309 003</u>	Jan 26, 2007
<u>AP</u>	+!	<u>4GM/100ML (40MG/ML)</u>	<u>N020309 001</u>	Jun 24, 1994
<u>AP</u>	+!	<u>4GM/50ML (80MG/ML)</u>	<u>N020309 002</u>	Jun 24, 1994
<u>AP</u>	+	<u>20GM/500ML (40MG/ML)</u>	<u>N020309 004</u>	Jan 18, 1995
<u>AP</u>	+	<u>40GM/1000ML (40MG/ML)</u>	<u>N020309 005</u>	Jan 18, 1995
<u>AP</u>	HQ SPCLT PHARMA	<u>2GM/50ML (40MG/ML)</u>	<u>A207350 001</u>	Dec 06, 2017
<u>AP</u>		<u>4GM/100ML (40MG/ML)</u>	<u>A207350 002</u>	Dec 06, 2017
<u>AP</u>		<u>4GM/50ML (80MG/ML)</u>	<u>A207350 003</u>	Dec 06, 2017
<u>AP</u>		<u>20GM/500ML (40MG/ML)</u>	<u>A207350 004</u>	Dec 06, 2017
<u>AP</u>		<u>40GM/1000ML (40MG/ML)</u>	<u>A207350 005</u>	Dec 06, 2017

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! HOSPIRA	2GM/100ML
------------	-----------

N020488 002	Jul 11, 1995
-------------	--------------

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

<u>AP</u>	EXELA PHARMA SCS LLC	<u>5GM/10ML (500MG/ML)</u>	<u>A206039 001</u>	Dec 18, 2014
<u>AP</u>	+! FRESENIUS KABI USA	<u>5GM/10ML (500MG/ML)</u>	<u>N019316 001</u>	Sep 08, 1986
<u>AP</u>	! HOSPIRA	<u>5GM/10ML (500MG/ML)</u>	<u>A075151 001</u>	Apr 25, 2000
	+! FRESENIUS KABI USA	10GM/20ML (500MG/ML)	N019316 003	Jan 29, 2016
	+!	25GM/50ML (500MG/ML)	N019316 004	Jan 29, 2016
	HOSPIRA INC	10GM/20ML (500MG/ML)	A202411 001	May 14, 2015

MAGNESIUM SULFATE

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MAGNESIUM SULFATE

+! FRESENIUS KABI USA	1GM/2ML (500MG/ML)
-----------------------	--------------------

N019316 002	Sep 08, 1986
-------------	--------------

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

TIS-U-SOL

<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML</u>	<u>N018508 001</u>	Feb 19, 1982
-----------	-----------------	--	--------------------	--------------

TIS-U-SOL IN PLASTIC CONTAINER

<u>AT</u>	BAXTER HLTHCARE	<u>20MG/100ML; 40MG/100ML; 6.25MG/100ML; 800MG/100ML; 8.75MG/100ML</u>	<u>N018336 001</u>
-----------	-----------------	--	--------------------

MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

POWDER; ORAL

COLPREP KIT

+! GATOR PHARMS	1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT
-----------------	-----------------------------------

N204553 001	Dec 27, 2016
-------------	--------------

SOLUTION; ORAL

SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE

<u>AA</u>	NOVEL LABS INC	<u>1.6GM/BOT; 3.13GM/BOT; 17.5GM/BOT</u>	<u>A202511 001</u>	Feb 23, 2017
-----------	----------------	--	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-267 (of 436)

MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE

SOLUTION;ORAL

SUPREP BOWEL PREP KIT

AA +! BRAINTREE LABS 1.6GM/BOT;3.13GM/BOT;17.5GM/BOT N022372 001 Aug 05, 2010

MALATHION

LOTION;TOPICAL

MALATHION

AT SUVEN LIFE 0.5% A091559 001 May 23, 2012

OVIDE

AT +! TARO PHARM 0.5% N018613 001 Aug 02, 1982

MANGANESE CHLORIDE

INJECTABLE;INJECTION

MANGANESE CHLORIDE IN PLASTIC CONTAINER

HOSPIRA EQ 0.1MG MANGANESE/ML N018962 001 Jun 26, 1986

MANNITOL

INJECTABLE;INJECTION

MANNITOL 10% IN PLASTIC CONTAINER

AP B BRAUN 10GM/100ML N020006 002 Jul 26, 1993

AP ICU MEDICAL INC 10GM/100ML N019603 002 Jan 08, 1987

MANNITOL 15% IN PLASTIC CONTAINER

AP B BRAUN 15GM/100ML N020006 003 Jul 26, 1993

AP ICU MEDICAL INC 15GM/100ML N019603 003 Jan 08, 1990

MANNITOL 20% IN PLASTIC CONTAINER

AP B BRAUN 20GM/100ML N020006 004 Jul 26, 1993

AP ICU MEDICAL INC 20GM/100ML N019603 004 Jan 08, 1990

MANNITOL 25%

AP FRESENIUS KABI USA 12.5GM/50ML A080677 001

AP HOSPIRA 12.5GM/50ML N016269 006 Aug 25, 1994

AP INTL MEDICATION 12.5GM/50ML A083051 001

AP LUITPOLD 12.5GM/50ML A087409 001 Jan 21, 1982

MANNITOL 5% IN PLASTIC CONTAINER

AP B BRAUN 5GM/100ML N020006 001 Jul 26, 1993

AP ICU MEDICAL INC 5GM/100ML N019603 001 Jan 08, 1987

OSMITROL 10% IN WATER

AP BAXTER HLTHCARE 10GM/100ML N013684 002

OSMITROL 10% IN WATER IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 10GM/100ML N013684 006

OSMITROL 15% IN WATER

AP BAXTER HLTHCARE 15GM/100ML N013684 004

OSMITROL 15% IN WATER IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 15GM/100ML N013684 008

OSMITROL 20% IN WATER

AP BAXTER HLTHCARE 20GM/100ML N013684 003

OSMITROL 20% IN WATER IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 20GM/100ML N013684 007

OSMITROL 5% IN WATER

AP BAXTER HLTHCARE 5GM/100ML N013684 001

OSMITROL 5% IN WATER IN PLASTIC CONTAINER

AP BAXTER HLTHCARE 5GM/100ML N013684 005

SOLUTION;IRRIGATION

RESECTISOL IN PLASTIC CONTAINER

B BRAUN 5GM/100ML N016772 002

MANNITOL; SORBITOL

SOLUTION;IRRIGATION

SORBITOL-MANNITOL IN PLASTIC CONTAINER

ICU MEDICAL INC 540MG/100ML;2.7GM/100ML N018316 001

MAPROTILINE HYDROCHLORIDE

TABLET;ORAL

MAPROTILINE HYDROCHLORIDE

MYLAN 25MG A072285 002 Oct 03, 1988

! 50MG A072285 001 Oct 03, 1988

75MG A072285 003 Oct 03, 1988

MARAVIROC

SOLUTION;ORAL

SELZENTRY

+! VIIV HLTHCARE 20MG/ML N208984 001 Nov 04, 2016

TABLET;ORAL

SELZENTRY

+ VIIV HLTHCARE 25MG N022128 003 Nov 04, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-268 (of 436)

MARAVIROC

TABLET;ORAL			
SELZENTRY			
+	75MG	N022128 004	Nov 04, 2016
+	150MG	N022128 001	Aug 06, 2007
+!	300MG	N022128 002	Aug 06, 2007

MEBENDAZOLE

TABLET, CHEWABLE;ORAL			
EMVERM			
! IMPAX LABS INC	100MG	A073580 001	Jan 04, 1995
VERMOX			
+! JANSSEN PHARMS	500MG	N208398 001	Oct 19, 2016

MECAMYLAMINE HYDROCHLORIDE

TABLET;ORAL			
MECAMYLAMINE HYDROCHLORIDE			
! NEXGEN PHARMA	2.5MG	A204054 001	Mar 19, 2013

MECASERMIN RECOMBINANT

INJECTABLE;SUBCUTANEOUS			
INCRELEX			
+! IPSEN INC	40MG/4ML (10MG/ML)	N021839 001	Aug 30, 2005

MECHILORETHAMINE HYDROCHLORIDE

GEL;TOPICAL			
VALCHLOR			
+! ATELION PHARMS LTD	EQ 0.016% BASE	N202317 001	Aug 23, 2013
INJECTABLE;INJECTION			
MUSTARGEN			
+! RECORDATI RARE	10MG/VIAL	N006695 001	

MECLIZINE HYDROCHLORIDE

TABLET;ORAL			
<b>MECLIZINE HYDROCHLORIDE</b>			
<b>AA</b>	AMNEAL PHARMS	<b>12.5MG</b>	<b>A201451 001</b> Feb 23, 2011
<b>AA</b>		<b>25MG</b>	<b>A201451 002</b> Feb 23, 2011
<b>AA</b>	EPIC PHARMA LLC	<b>12.5MG</b>	<b>A200294 001</b> Apr 13, 2012
<b>AA</b>		<b>25MG</b>	<b>A200294 002</b> Apr 13, 2012
<b>AA</b>	JUBILANT CADISTA	<b>12.5MG</b>	<b>A040659 001</b> Jun 04, 2010
<b>AA</b>		<b>25MG</b>	<b>A040659 002</b> Jun 04, 2010
<b>AA</b>	MYLAN PHARMS INC	<b>12.5MG</b>	<b>A202640 001</b> Sep 17, 2012
<b>AA</b>		<b>25MG</b>	<b>A202640 002</b> Sep 17, 2012
<b>AA</b>	PAR PHARM	<b>12.5MG</b>	<b>A087127 001</b>
<b>AA</b>		<b>25MG</b>	<b>A087128 001</b>
<b>AA</b>	SANDOZ	<b>12.5MG</b>	<b>A084843 002</b> May 22, 1989
<b>AA</b>		<b>25MG</b>	<b>A084092 003</b> May 22, 1989

MECLOFENAMATE SODIUM

CAPSULE;ORAL			
MECLOFENAMATE SODIUM			
MYLAN	EQ 50MG BASE	A071081 002	Sep 03, 1986
!	EQ 100MG BASE	A071081 001	Sep 03, 1986

MEDROXYPROGESTERONE ACETATE

INJECTABLE;INJECTION			
<b>DEPO-PROVERA</b>			

<b>AB</b>	+! PHARMACIA AND UPJOHN	<b>150MG/ML</b>	<b>N020246 001</b> Oct 29, 1992
-----------	-------------------------	-----------------	---------------------------------

MEDROXYPROGESTERONE ACETATE

<b>AB</b>	AMPHASTAR PHARMS INC	<b>150MG/ML</b>	<b>A077235 001</b> Nov 28, 2017
-----------	----------------------	-----------------	---------------------------------

<b>AB</b>	TEVA PHARMS USA	<b>150MG/ML</b>	<b>A077334 001</b> Nov 28, 2017
<b>AB</b>	DEPO-PROVERA		<b>A076553 001</b> Jul 28, 2004

+! PHARMACIA AND UPJOHN

INJECTABLE;SUBCUTANEOUS			
DEPO-SUBQ PROVERA 104			
+! PHARMACIA AND UPJOHN	400MG/ML	N012541 003	

DEPO-SUBQ PROVERA 104			
+! PHARMACIA AND UPJOHN	400MG/ML	N012541 003	

INJECTABLE;SUBCUTANEOUS			
DEPO-SUBQ PROVERA 104			

+! PHARMACIA AND UPJOHN

INJECTABLE;SUBCUTANEOUS			
DEPO-SUBQ PROVERA 104			

+! PHARMACIA AND UPJOHN

INJECTABLE;SUBCUTANEOUS			
DEPO-SUBQ PROVERA 104			

+! PHARMACIA AND UPJOHN

MEDROXYPROGESTERONE ACETATE

<b>AB</b>	BARR	<b>2.5MG</b>	<b>A040159 001</b> Aug 09, 1996
<b>AB</b>		<b>5MG</b>	<b>A040159 002</b> Aug 09, 1996
<b>AB</b>		<b>10MG</b>	<b>A040159 003</b> Aug 09, 1996

## MEDROXYPROGESTERONE ACETATE

TABLET; ORAL

PROVERA

AB + PHARMACIA AND 2.5MG N011839 001  
UPJOHN  
AB + 5MG N011839 003  
AB +! 10MG N011839 004

## MEFENAMIC ACID

CAPSULE; ORAL

## **MEFENAMIC ACID**

<u><b>AB</b></u>	BRECKENRIDGE PHARM	<u><b>250MG</b></u>	<u><b>A090359 001</b></u>	Feb 05, 2013
<u><b>AB</b></u>	LUPIN LTD	<u><b>250MG</b></u>	<u><b>A091322 001</b></u>	Jul 22, 2011
<u><b>AB</b></u>	VINTAGE PHARMS LLC	<u><b>250MG</b></u>	<u><b>A091608 001</b></u>	Jun 02, 2014
<u><b>AB</b></u>	VIVA HEALTHCARE	<u><b>250MG</b></u>	<u><b>A090562 001</b></u>	Nov 19, 2010

BONSTELL

**AB** +! SHTONOGT INC 250MG N015034 003

## MEETQUINE HYDROCHLORIDE

**TABLET; ORAL**

#### MEETOUTLINE HYDROCHILOBE

AB ! BARR 250MG A076392 001 Dec 29, 2003  
AB WEST-WARD PHARMS 250MG A076523 001 Oct 01, 2004  
INT

MEGESTROL ACETATE

**SUSPENSION; ORAL**

MEGACE ES

<u>AB</u>	<u>+</u>	ENDO PHARMS INC	<u>125MG/ML</u>	<u>N021778</u>	<u>001</u>	Jul 05, 2005
<b><u>MEGESTROL ACETATE</u></b>						
<u>AB</u>		BRECKENRIDGE PHARM	<u>125MG/ML</u>	<u>A204688</u>	<u>001</u>	Dec 01, 2017
<u>AB</u>		HI-TECH PHARMACAL	<u>40MG/ML</u>	<u>A203960</u>	<u>001</u>	Jun 09, 2017
<u>AB</u>		PAR PHARM	<u>40MG/ML</u>	<u>A075671</u>	<u>001</u>	Jul 25, 2001
<u>AB</u>		TEVA PHARMS	<u>40MG/ML</u>	<u>A075681</u>	<u>001</u>	May 05, 2003
<u>AB</u>		TWI PHARMS INC	<u>125MG/ML</u>	<u>A203139</u>	<u>001</u>	Aug 27, 2014
<u>AB</u>	<u>!</u>	WEST-WARD PHARMS	<u>40MG/ML</u>	<u>A075997</u>	<u>001</u>	Feb 15, 2002
		INT				
<u>AB</u>		WOCKHARDT BIO AG	<u>40MG/ML</u>	<u>A076721</u>	<u>001</u>	Nov 01, 2004

TABLET; ORAL

## MEGESTROL ACETATE

<u>AB</u>	BARR	<u>20MG</u>	<u>A074621</u>	<u>002</u>	Aug 16, 1996
<u>AB</u>		<u>40MG</u>	<u>A074621</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	PAR PHARM	<u>20MG</u>	<u>A072422</u>	<u>001</u>	Aug 08, 1988
<u>AB</u>		<u>40MG</u>	<u>A072423</u>	<u>001</u>	Aug 08, 1988
<u>AB</u>	WEST-WARD PHARMS	<u>20MG</u>	<u>A074458</u>	<u>001</u>	Sep 29, 1995
	INT				
<u>AB</u>		<u>40MG</u>	<u>A074458</u>	<u>002</u>	Sep 29, 1995

## METOXTCAM

CAPSULE: OBAT

VIVIODEX

+ IROKO PHARMS LLC 5MG N207233 001 Oct 22, 2015  
+! 10MG N207233 002 Oct 22, 2015

**TABLET; ORAL**

## MELOXICAM

<b><u>AB</u></b>	APOTEX INC	<b><u>7.5MG</u></b>	<b><u>A077882_001</u></b>	Jul 20, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077882_002</u></b>	Jul 20, 2006
<b><u>AB</u></b>	AUROBINDO PHARMA	<b><u>7.5MG</u></b>	<b><u>A078008_001</u></b>	Oct 02, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A078008_002</u></b>	Oct 02, 2006
<b><u>AB</u></b>	BRECKENRIDGE PHARM	<b><u>7.5MG</u></b>	<b><u>A077920_001</u></b>	Jul 19, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077920_002</u></b>	Jul 19, 2006
<b><u>AB</u></b>	CIPLA LTD	<b><u>7.5MG</u></b>	<b><u>A077929_001</u></b>	Jul 19, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077929_002</u></b>	Jul 19, 2006
<b><u>AB</u></b>	DR REDDYS LABS INC	<b><u>7.5MG</u></b>	<b><u>A077931_001</u></b>	Jul 25, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077931_002</u></b>	Jul 25, 2006
<b><u>AB</u></b>	GLENMARK GENERICS	<b><u>7.5MG</u></b>	<b><u>A077932_001</u></b>	Jul 19, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077932_002</u></b>	Jul 19, 2006
<b><u>AB</u></b>	LUPIN PHARMS	<b><u>7.5MG</u></b>	<b><u>A077944_001</u></b>	Jul 19, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077944_002</u></b>	Jul 19, 2006
<b><u>AB</u></b>	MYLAN	<b><u>7.5MG</u></b>	<b><u>A077923_001</u></b>	Jul 19, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077923_002</u></b>	Jul 19, 2006
<b><u>AB</u></b>	PURACAP PHARM	<b><u>7.5MG</u></b>	<b><u>A077938_001</u></b>	Jul 19, 2006
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077938_002</u></b>	Jul 19, 2006
<b><u>AB</u></b>	STRIDES PHARMA	<b><u>7.5MG</u></b>	<b><u>A077928_001</u></b>	May 13, 2009
<b><u>AB</u></b>		<b><u>15MG</u></b>	<b><u>A077928_002</u></b>	May 13, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-270 (of 436)

MELOXICAM

TABLET;ORAL

MELOXICAM

<u>AB</u>	TARO	<u>7.5MG</u>	<u>A078102 001</u>	Nov 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A078102 002</u>	Nov 07, 2006
<u>AB</u>	TEVA PHARMS	<u>7.5MG</u>	<u>A077936 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077936 002</u>	Jul 19, 2006
<u>AB</u>	UNICHEM	<u>7.5MG</u>	<u>A077927 001</u>	Dec 20, 2006
<u>AB</u>		<u>15MG</u>	<u>A077927 002</u>	Dec 20, 2006
<u>AB</u>	YUNG SHIN PHARM	<u>7.5MG</u>	<u>A077918 001</u>	Dec 07, 2006
<u>AB</u>		<u>15MG</u>	<u>A077918 002</u>	Dec 07, 2006
<u>AB</u>	ZYDUS PHARMS USA	<u>7.5MG</u>	<u>A077921 001</u>	Jul 19, 2006
<u>AB</u>		<u>15MG</u>	<u>A077921 002</u>	Jul 19, 2006
<u>MOBIC</u>				
<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>7.5MG</u>	<u>N020938 001</u>	Apr 13, 2000
<u>AB</u>	+!	<u>15MG</u>	<u>N020938 002</u>	Aug 23, 2000

MELPHALAN

TABLET;ORAL

ALKERAN

<u>AB</u>	+! APOTEX INC	<u>2MG</u>	<u>N014691 002</u>
<u>AB</u>	MELPHALAN	<u>2MG</u>	<u>A207809 001</u> Mar 22, 2017

MELPHALAN HYDROCHLORIDE

INJECTABLE;INJECTION

MELPHALAN HYDROCHLORIDE

<u>AP</u>	ACTAVIS LLC	<u>EQ 50MG BASE/VIAL</u>	<u>A206018 001</u>	Dec 19, 2016
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 50MG BASE/VIAL</u>	<u>A203655 001</u>	Dec 08, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 50MG BASE/VIAL</u>	<u>A203393 001</u>	Dec 22, 2017
<u>AP</u>	! MYLAN INSTITUTIONAL	<u>EQ 50MG BASE/VIAL</u>	<u>A090270 001</u>	Jun 09, 2009
<u>AP</u>	PAR STERILE PRODUCTS	<u>EQ 50MG BASE/VIAL</u>	<u>A204773 001</u>	Aug 22, 2016
<u>AP</u>	SAGENT PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>A201379 001</u>	Feb 28, 2017
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 50MG BASE/VIAL</u>	<u>A090303 001</u>	Oct 28, 2010

POWDER;IV (INFUSION)

EVOMELA

+! SPECTRUM PHARMS	EQ 50MG BASE/VIAL	N207155 001	Mar 10, 2016
--------------------	-------------------	-------------	--------------

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>7MG</u>	<u>A205825 001</u>	Oct 12, 2016
<u>AB</u>		<u>14MG</u>	<u>A205825 002</u>	Oct 12, 2016
<u>AB</u>		<u>21MG</u>	<u>A205825 003</u>	Oct 12, 2016
<u>AB</u>		<u>28MG</u>	<u>A205825 004</u>	Oct 12, 2016
<u>AB</u>	ANCHEM PHARMS	<u>7MG</u>	<u>A205784 001</u>	Jun 09, 2017
<u>AB</u>		<u>14MG</u>	<u>A205784 002</u>	Jun 09, 2017
<u>AB</u>		<u>21MG</u>	<u>A205784 003</u>	Jun 09, 2017
<u>AB</u>		<u>28MG</u>	<u>A205784 004</u>	Jun 09, 2017
<u>AB</u>	APOTEX INC	<u>7MG</u>	<u>A206135 001</u>	Nov 22, 2016
<u>AB</u>		<u>14MG</u>	<u>A206135 002</u>	Nov 22, 2016
<u>AB</u>		<u>21MG</u>	<u>A206135 003</u>	Nov 22, 2016
<u>AB</u>		<u>28MG</u>	<u>A206135 004</u>	Nov 22, 2016
<u>AB</u>	LUPIN LTD	<u>7MG</u>	<u>A206028 001</u>	Sep 28, 2016
<u>AB</u>		<u>14MG</u>	<u>A206028 002</u>	Sep 28, 2016
<u>AB</u>		<u>21MG</u>	<u>A206028 003</u>	Sep 28, 2016
<u>AB</u>		<u>28MG</u>	<u>A206028 004</u>	Sep 28, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>7MG</u>	<u>A206032 001</u>	Sep 28, 2016
<u>AB</u>		<u>14MG</u>	<u>A206032 002</u>	Sep 28, 2016
<u>AB</u>		<u>21MG</u>	<u>A206032 003</u>	Sep 28, 2016
<u>AB</u>		<u>28MG</u>	<u>A206032 004</u>	Sep 28, 2016
<u>AB</u>	SUN PHARMA GLOBAL	<u>7MG</u>	<u>A205905 001</u>	Sep 28, 2016
<u>AB</u>		<u>14MG</u>	<u>A205905 002</u>	Sep 28, 2016
<u>AB</u>		<u>21MG</u>	<u>A205905 003</u>	Sep 28, 2016
<u>AB</u>		<u>28MG</u>	<u>A205905 004</u>	Sep 28, 2016
<u>AB</u>	ZYDUS PHARMS USA INC	<u>7MG</u>	<u>A203293 001</u>	Aug 03, 2017
<u>AB</u>		<u>14MG</u>	<u>A203293 002</u>	Aug 03, 2017
<u>AB</u>		<u>21MG</u>	<u>A203293 003</u>	Aug 03, 2017
<u>AB</u>		<u>28MG</u>	<u>A203293 004</u>	Aug 03, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-271 (of 436)

MEMANTINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

NAMENDA XR

<u>AB</u>	+	FOREST LABS LLC	<u>7MG</u>	<u>N022525</u> <u>001</u>	Jun 21, 2010
<u>AB</u>	+		<u>14MG</u>	<u>N022525</u> <u>002</u>	Jun 21, 2010
<u>AB</u>	+		<u>21MG</u>	<u>N022525</u> <u>003</u>	Jun 21, 2010
<u>AB</u>	+!		<u>28MG</u>	<u>N022525</u> <u>004</u>	Jun 21, 2010

SOLUTION; ORAL

MEMANTINE HYDROCHLORIDE

<u>AA</u>		BIO-PHARM INC	<u>2MG/ML</u>	<u>A205446</u> <u>001</u>	Dec 07, 2015
<u>AA</u>		MACLEODS PHARMS LTD	<u>2MG/ML</u>	<u>A202790</u> <u>001</u>	Oct 13, 2015
<u>AA</u>		SILARX PHARMS INC	<u>2MG/ML</u>	<u>A204033</u> <u>001</u>	Oct 13, 2015

NAMENDA

<u>AA</u>	+!	FOREST LABS LLC	<u>2MG/ML</u>	<u>N021627</u> <u>001</u>	Apr 18, 2005
-----------	----	-----------------	---------------	---------------------------	--------------

TABLET; ORAL

MEMANTINE HYDROCHLORIDE

<u>AB</u>		AJANTA PHARMA LTD	<u>5MG</u>	<u>A206528</u> <u>001</u>	Nov 30, 2015
<u>AB</u>			<u>10MG</u>	<u>A206528</u> <u>002</u>	Nov 30, 2015
<u>AB</u>		ALEMBIC PHARMS LTD	<u>5MG</u>	<u>A200891</u> <u>001</u>	Oct 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A200891</u> <u>002</u>	Oct 13, 2015
<u>AB</u>		AMNEAL PHARMS	<u>5MG</u>	<u>A090041</u> <u>001</u>	Apr 10, 2015
<u>AB</u>			<u>10MG</u>	<u>A090041</u> <u>002</u>	Apr 10, 2015
<u>AB</u>		AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203175</u> <u>001</u>	Oct 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A203175</u> <u>002</u>	Oct 13, 2015
<u>AB</u>		DR REDDYS LABS LTD	<u>5MG</u>	<u>A090048</u> <u>001</u>	Apr 14, 2010
<u>AB</u>			<u>10MG</u>	<u>A090048</u> <u>002</u>	Apr 14, 2010
<u>AB</u>		JUBILANT GENERICS	<u>5MG</u>	<u>A091585</u> <u>001</u>	Oct 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A091585</u> <u>002</u>	Oct 13, 2015
<u>AB</u>		LUPIN LTD	<u>5MG</u>	<u>A090051</u> <u>001</u>	Apr 10, 2015
<u>AB</u>			<u>10MG</u>	<u>A090051</u> <u>002</u>	Apr 10, 2015
<u>AB</u>		MACLEODS PHARMS LTD	<u>5MG</u>	<u>A202840</u> <u>001</u>	Oct 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A202840</u> <u>002</u>	Oct 13, 2015
<u>AB</u>		MYLAN PHARMS INC	<u>5MG</u>	<u>A079225</u> <u>001</u>	Jan 30, 2015
<u>AB</u>			<u>10MG</u>	<u>A079225</u> <u>002</u>	Jan 30, 2015
<u>AB</u>		PURACAP PHARM LLC	<u>5MG</u>	<u>A206855</u> <u>001</u>	Nov 17, 2015
<u>AB</u>			<u>10MG</u>	<u>A206855</u> <u>002</u>	Nov 17, 2015
<u>AB</u>		SILARX PHARMS INC	<u>5MG</u>	<u>A207236</u> <u>001</u>	Nov 10, 2016
<u>AB</u>			<u>10MG</u>	<u>A207236</u> <u>002</u>	Nov 10, 2016
<u>AB</u>		STRIDES PHARMA	<u>5MG</u>	<u>A202350</u> <u>001</u>	May 23, 2017
<u>AB</u>			<u>10MG</u>	<u>A202350</u> <u>002</u>	May 23, 2017
<u>AB</u>		SUN PHARMA GLOBAL	<u>5MG</u>	<u>A090058</u> <u>001</u>	May 05, 2010
<u>AB</u>			<u>10MG</u>	<u>A090058</u> <u>002</u>	May 05, 2010
<u>AB</u>		TEVA PHARMS	<u>5MG</u>	<u>A090052</u> <u>001</u>	Oct 25, 2011
<u>AB</u>			<u>10MG</u>	<u>A090052</u> <u>002</u>	Oct 25, 2011
<u>AB</u>		TORRENT PHARMS LTD	<u>5MG</u>	<u>A200155</u> <u>001</u>	Oct 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A200155</u> <u>002</u>	Oct 13, 2015
<u>AB</u>		UNICHEM LABS LTD	<u>5MG</u>	<u>A200022</u> <u>001</u>	Oct 13, 2015
<u>AB</u>			<u>10MG</u>	<u>A200022</u> <u>002</u>	Oct 13, 2015
<u>AB</u>		UPSHER-SMITH LABS	<u>5MG</u>	<u>A090043</u> <u>001</u>	Jul 31, 2015
<u>AB</u>			<u>10MG</u>	<u>A090043</u> <u>002</u>	Jul 31, 2015
<u>AB</u>		WOCKHARDT LTD	<u>5MG</u>	<u>A090073</u> <u>001</u>	Sep 04, 2015
<u>AB</u>			<u>10MG</u>	<u>A090073</u> <u>002</u>	Sep 04, 2015
<u>AB</u>		ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A090961</u> <u>001</u>	Jul 10, 2017
<u>AB</u>			<u>10MG</u>	<u>A090961</u> <u>002</u>	Jul 10, 2017
<u>NAMENDA</u>					
<u>AB</u>	+	FOREST LABS LLC	<u>5MG</u>	<u>N021487</u> <u>001</u>	Oct 16, 2003
<u>AB</u>	+!		<u>10MG</u>	<u>N021487</u> <u>002</u>	Oct 16, 2003

MENOTROPINS (FSH; LH)

INJECTABLE; SUBCUTANEOUS				
MENOPUR				
+!	FERRING		75 IU/VIAL; 75 IU/VIAL	N021663 001 Oct 29, 2004

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION				
<u>DEMEROL</u>				

<u>AP</u>	+!	HOSPIRA	<u>25MG/ML</u>	<u>N021171</u> <u>001</u>
<u>AP</u>	+!		<u>50MG/ML</u>	<u>N021171</u> <u>002</u>
<u>AP</u>	+!		<u>75MG/ML</u>	<u>N021171</u> <u>003</u>
<u>AP</u>	+!		<u>100MG/ML</u>	<u>N021171</u> <u>004</u>

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-272 (of 436)

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HYDROCHLORIDE

<u>AP</u>	WEST-WARD PHARMS INT	<u>25MG/ML</u>	<u>A080445 001</u>
<u>AP</u>		<u>25MG/ML</u>	<u>A080445 007</u>
<u>AP</u>		<u>50MG/ML</u>	<u>A080445 002</u>
<u>AP</u>		<u>50MG/ML</u>	<u>A080445 008</u>
<u>AP</u>		<u>75MG/ML</u>	<u>A080445 003</u>
<u>AP</u>		<u>75MG/ML</u>	<u>A080445 009</u>
<u>AP</u>		<u>100MG/ML</u>	<u>A080445 004</u>
<u>AP</u>		<u>100MG/ML</u>	<u>A080445 010</u>

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u> !	ICU MEDICAL INC	<u>10MG/ML</u>	<u>A088432 001</u> Aug 16, 1984
<u>AP</u>	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A081002 001</u> Jul 30, 1993

SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

!	WEST-WARD PHARMS INT	50MG/5ML
---	-------------------------	----------

A088744 001 Jan 30, 1985

TABLET; ORAL

DEMEROL

<u>AA</u> +!	US PHARM HOLDINGS	<u>50MG</u>	<u>N005010 001</u>
<u>AA</u> +!		<u>100MG</u>	<u>N005010 004</u>
<u>MEPERIDINE HYDROCHLORIDE</u>			
<u>AA</u>	BARR	<u>50MG</u>	<u>A088639 001</u> Jul 02, 1984
<u>AA</u>		<u>100MG</u>	<u>A088640 001</u> Sep 19, 1984
<u>AA</u>	EPIC PHARMA	<u>50MG</u>	<u>A040331 001</u> May 28, 1999
<u>AA</u>		<u>100MG</u>	<u>A040331 002</u> May 28, 1999
<u>AA</u>	MIKART	<u>50MG</u>	<u>A040893 001</u> Jun 24, 2009
<u>AA</u>		<u>100MG</u>	<u>A040893 003</u> Jun 24, 2009
<u>AA</u>	SPECGX LLC	<u>50MG</u>	<u>A040352 001</u> Jun 13, 2000
<u>AA</u>		<u>100MG</u>	<u>A040352 002</u> Jun 13, 2000
<u>AA</u>	SUN PHARM IND'S INC	<u>50MG</u>	<u>A040446 001</u> Aug 08, 2002
<u>AA</u>		<u>100MG</u>	<u>A040446 002</u> Aug 08, 2002
<u>AA</u>	VINTAGE PHARMS	<u>50MG</u>	<u>A040191 001</u> Dec 17, 1998
<u>AA</u>		<u>100MG</u>	<u>A040191 002</u> Dec 17, 1998
<u>AA</u>	WEST-WARD PHARMS INT	<u>50MG</u>	<u>A040110 001</u> Mar 12, 1997
<u>AA</u>		<u>100MG</u>	<u>A040110 002</u> Mar 12, 1997
	MIKART	75MG	A040893 002 Jun 24, 2009
		150MG	A040893 004 Jun 24, 2009

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

<u>AP</u> +!	HOSPIRA	<u>1%</u>	<u>N012250 001</u>
<u>AP</u> +!		<u>1.5%</u>	<u>N012250 005</u>
<u>AP</u> +!		<u>2%</u>	<u>N012250 002</u>

ISOCOCAINE HYDROCHLORIDE

<u>AP</u> !	SEPTODONT INC	<u>3%</u>	<u>A080925 001</u>
-------------	---------------	-----------	--------------------

MEPIVACAINE HYDROCHLORIDE

<u>AP</u>	HOSPIRA INC	<u>3%</u>	<u>A040806 001</u> Apr 28, 2008
-----------	-------------	-----------	---------------------------------

POLOCAINE

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A089407 001</u> Dec 01, 1986
<u>AP</u>		<u>2%</u>	<u>A089410 001</u> Dec 01, 1986

POLOCAINE-MPF

<u>AP</u>	FRESENIUS KABI USA	<u>1%</u>	<u>A089406 001</u> Dec 01, 1986
<u>AP</u>		<u>1.5%</u>	<u>A089408 001</u> Dec 01, 1986
<u>AP</u>		<u>2%</u>	<u>A089409 001</u> Dec 01, 1986

SCANDONEST PLAIN

<u>AP</u> !	DEFROCO	<u>3%</u>	<u>A088387 001</u> Oct 10, 1984
-------------	---------	-----------	---------------------------------

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

<u>AA</u>	ALEMBIC PHARMS LTD	<u>200MG</u>	<u>A090122 001</u> Feb 18, 2009
<u>AA</u>		<u>400MG</u>	<u>A090122 002</u> Feb 18, 2009
<u>AA</u>	INVAGEN PHARMS	<u>200MG</u>	<u>A040797 001</u> Feb 27, 2008
<u>AA</u>		<u>400MG</u>	<u>A040797 002</u> Feb 27, 2008
<u>AA</u> !	WATSON LABS	<u>200MG</u>	<u>A083304 001</u>
<u>AA</u> !		<u>400MG</u>	<u>A083308 001</u>

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-273 (of 436)

MERCAPTOPURINE

SUSPENSION;ORAL

PURIXAN

+! NOVA LABS LTD

20MG/ML

N205919 001 Apr 28, 2014

TABLET;ORAL

MERCAPTOPURINE

**AB** DR REDDYS LABS SA **50MG**

**A040461 001** Feb 11, 2004

**AB** MYLAN **50MG**

**A040594 001** Jul 01, 2005

**AB** ! WEST-WARD PHARMS INT

**A040528 001** Feb 13, 2004

PURINETHOL

**AB** + STASON PHARMS **50MG**

**N009053 002**

MEROPENEM

INJECTABLE;INJECTION

MEROPENEM

**AP** ACS DOBFAR **500MG/VIAL**

**A091404 001** Oct 26, 2011

**AP** **1GM/VIAL**

**A091404 002** Oct 26, 2011

**AP** AMNEAL PHARMS **500MG/VIAL**

**A205883 001** Apr 12, 2016

**AP** **1GM/VIAL**

**A205883 002** Apr 12, 2016

**AP** AUROBINDO PHARMA LTD **500MG/VIAL**

**A205835 001** Mar 27, 2017

**AP** **1GM/VIAL**

**A205835 002** Mar 27, 2017

**AP** DAEWOONG PHARM CO **500MG/VIAL**

**A204854 001** Dec 18, 2015

**AP** **1GM/VIAL**

**A204854 002** Dec 18, 2015

**AP** GLAND PHARMA LTD **500MG/VIAL**

**A206141 001** Jun 08, 2016

**AP** **1GM/VIAL**

**A206141 002** Jun 08, 2016

**AP** HOSPIRA INC **500MG/VIAL**

**A090940 001** Jun 22, 2010

**AP** **1GM/VIAL**

**A090940 002** Jun 22, 2010

**AP** PAR STERILE PRODUCTS **500MG/VIAL**

**A204139 001** Jun 09, 2016

**AP** **1GM/VIAL**

**A204139 002** Jun 09, 2016

**AP** SAVIOR LIFETEC CORP **500MG/VIAL**

**A206086 001** Apr 19, 2016

**AP** **1GM/VIAL**

**A206086 002** Apr 19, 2016

MERREM

**AP** +! PFIZER **500MG/VIAL**

**N050706 003** Jun 21, 1996

**AP** +! **1GM/VIAL**

**N050706 001** Jun 21, 1996

POWDER;IV (INFUSION)

MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER

B BRAUN MEDICAL INC 500MG/VIAL

N202106 001 Apr 30, 2015

1GM/VIAL

N202106 002 Apr 30, 2015

MEROPELEM; VABORBACTAM

POWDER;IV (INFUSION)

VABOMERE

+! REMPEX PHARMS

1GM/VIAL;1GM/VIAL

N209776 001 Aug 29, 2017

MEDCNS

MESALAMINE

CAPSULE, DELAYED RELEASE;ORAL

DELZICOL

+! APIL

400MG

N204412 001 Feb 01, 2013

CAPSULE, EXTENDED RELEASE;ORAL

APRISO

+! VALEANT PHARMS INTL

375MG

N022301 001 Oct 31, 2008

PENTASA

+! SHIRE

250MG

N020049 001 May 10, 1993

+!

500MG

N020049 002 Jul 08, 2004

ENEMA;RECTAL

MESALAMINE

**AB** G AND W LABS INC **4GM/60ML**

**A076841 001** Sep 30, 2004

**AB** PERRIGO ISRAEL **4GM/60ML**

**A076751 001** Sep 17, 2004

ROWASA

**AB** +! MYLAN SPECIALITY LP **4GM/60ML**

**N019618 001** Dec 24, 1987

SFROWASA

**AB** + MYLAN SPECIALITY LP **4GM/60ML**

**N019618 002** Jun 20, 2008

SUPPOSITORY;RECTAL

CANASA

**AB** +! FOREST LABS LLC **1GM**

**N021252 002** Nov 05, 2004

MESALAMINE

**AB** MYLAN PHARMS INC **1GM**

**A204354 001** Nov 24, 2015

TABLET, DELAYED RELEASE;ORAL

ASACOL HD

**AB** +! APIL **800MG**

**N021830 001** May 29, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-274 (of 436)

MESALAMINE

TABLET, DELAYED RELEASE;ORAL

LIALDA

<u>AB</u>	+!	SHIRE	<u>1.2GM</u>	<u>N022000 001</u>	Jan 16, 2007
<b><u>MESALAMINE</u></b>					
<u>AB</u>		ZYDUS PHARMS USA INC	<u>800MG</u>	<u>A203286 001</u>	Jul 21, 2017
<u>AB</u>			<u>1.2GM</u>	<u>A091640 001</u>	Jun 05, 2017

MESNA

INJECTABLE; INTRAVENOUS

MESNA

<u>AP</u>		FRESENIUS KABI USA	<u>100MG/ML</u>	<u>A075811 001</u>	Apr 26, 2001
<u>AP</u>		GLAND PHARMA LTD	<u>100MG/ML</u>	<u>A206992 001</u>	Dec 18, 2017
<u>AP</u>		MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<u>A076488 001</u>	Mar 08, 2012
<u>AP</u>		SAGENT PHARMS	<u>100MG/ML</u>	<u>A090913 001</u>	Apr 13, 2010
<u>AP</u>		TEVA PHARMS USA	<u>100MG/ML</u>	<u>A075764 001</u>	Apr 27, 2001
<u>AP</u>		WEST-WARD PHARMS INT	<u>100MG/ML</u>	<u>A075739 001</u>	Jan 09, 2004

MESNEX

<u>AP</u>	+!	BAXTER HLTHCARE	<u>100MG/ML</u>	<u>N019884 001</u>	Dec 30, 1988
TABLET;ORAL					
	MESNEX				

+

! BAXTER HLTHCARE

400MG

N020855 001 Mar 21, 2002

MESTRANOL; NORETHINDRONE

TABLET;ORAL-28

NORINYL 1+50 28-DAY

! ACTAVIS LABS UT INC 0.05MG;1MG

N016659 001

METAPROTERENOL SULFATE

SYRUP;ORAL

METAPROTERENOL SULFATE

! SILARX

10MG/5ML

A073632 001 Jul 22, 1992

TABLET;ORAL

METAPROTERENOL SULFATE

PAR PHARM

10MG

A072024 001 Jun 28, 1988

!

20MG

A072025 001 Jun 28, 1988

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

METARAMINOL BITARTRATE

!

FRESENIUS KABI USA

EQ 10MG BASE/ML

A080722 001

METAXALONE

TABLET;ORAL

METAXALONE

<u>AB</u>		ACTAVIS LABS FL INC	<u>800MG</u>	<u>A203695 001</u>	Jun 15, 2017
<u>AB</u>		AMNEAL PHARMS	<u>800MG</u>	<u>A203399 001</u>	Jun 21, 2013
<u>AB</u>		LANNETT HOLDINGS INC	<u>800MG</u>	<u>A204770 001</u>	Nov 22, 2016
<u>AB</u>		SANDOZ	<u>800MG</u>	<u>A040445 001</u>	Mar 31, 2010
<u>AB</u>		SCIEGEN PHARMS INC	<u>800MG</u>	<u>A207466 001</u>	Aug 31, 2017

SKELAXIN

<u>AB</u>	+!	KING PHARMS	<u>800MG</u>	<u>N013217 003</u>	Aug 30, 2002
METAXALONE					
		COREPHARMA	400MG	A040486 001	Feb 27, 2015

METFORMIN HYDROCHLORIDE

SOLUTION;ORAL

RIOMET

! SUN PHARM INDs LTD

500MG/5ML

N021591 001 Sep 11, 2003

TABLET;ORAL

GLUCOPHAGE

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N020357 001</u>	Mar 03, 1995
<u>AB</u>	+		<u>850MG</u>	<u>N020357 002</u>	Mar 03, 1995
<u>AB</u>	+		<u>1GM</u>	<u>N020357 005</u>	Nov 05, 1998

METFORMIN HYDROCHLORIDE

<u>AB</u>		ALKEM	<u>500MG</u>	<u>A091184 001</u>	Nov 01, 2010
<u>AB</u>			<u>850MG</u>	<u>A091184 002</u>	Nov 01, 2010
<u>AB</u>			<u>1GM</u>	<u>A091184 003</u>	Nov 01, 2010
<u>AB</u>		AMNEAL PHARMS NY	<u>500MG</u>	<u>A077880 001</u>	Jun 05, 2006
<u>AB</u>			<u>850MG</u>	<u>A077880 002</u>	Jun 05, 2006
<u>AB</u>			<u>1GM</u>	<u>A077880 003</u>	Jun 05, 2006
<u>AB</u>		APOTEX	<u>500MG</u>	<u>A075984 001</u>	Apr 23, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-275 (of 436)

METFORMIN HYDROCHLORIDE

TABLET;ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>		<u>500MG</u>	<u>A090666 001</u>	Dec 07, 2011
<u>AB</u>		<u>850MG</u>	<u>A075984 002</u>	Apr 23, 2002
<u>AB</u>		<u>850MG</u>	<u>A090666 002</u>	Dec 07, 2011
<u>AB</u>		<u>1GM</u>	<u>A075984 003</u>	Apr 23, 2002
<u>AB</u>		<u>1GM</u>	<u>A090666 003</u>	Dec 07, 2011
<u>AB</u>	ATLAS PHARMS LLC	<u>500MG</u>	<u>A076033 001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A076033 002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A076033 003</u>	Jan 24, 2002
<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>A077095 001</u>	Jan 14, 2005
<u>AB</u>		<u>850MG</u>	<u>A077095 002</u>	Jan 14, 2005
<u>AB</u>		<u>1GM</u>	<u>A077095 003</u>	Jan 14, 2005
<u>AB</u>	CHARTWELL LIFE SCI	<u>500MG</u>	<u>A075972 001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A075972 002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A075972 003</u>	Jan 24, 2002
<u>AB</u>	CSPC OUYI PHARM CO	<u>500MG</u>	<u>A205096 001</u>	Jul 11, 2016
<u>AB</u>		<u>850MG</u>	<u>A205096 002</u>	Jul 11, 2016
<u>AB</u>		<u>1GM</u>	<u>A205096 003</u>	Jul 11, 2016
<u>AB</u>	DR REDDYS LABS INC	<u>500MG</u>	<u>A077787 001</u>	Aug 23, 2006
<u>AB</u>		<u>850MG</u>	<u>A077787 002</u>	Aug 23, 2006
<u>AB</u>		<u>1GM</u>	<u>A077787 003</u>	Aug 23, 2006
<u>AB</u>	GLENMARK GENERICS	<u>500MG</u>	<u>A078170 001</u>	May 23, 2008
<u>AB</u>		<u>850MG</u>	<u>A078170 002</u>	May 23, 2008
<u>AB</u>		<u>1GM</u>	<u>A078170 003</u>	May 23, 2008
<u>AB</u>	GRANULES INDIA	<u>500MG</u>	<u>A090564 001</u>	Apr 22, 2010
<u>AB</u>		<u>850MG</u>	<u>A090564 002</u>	Apr 22, 2010
<u>AB</u>		<u>1GM</u>	<u>A090564 003</u>	Apr 22, 2010
<u>AB</u>	INDICUS PHARMA	<u>500MG</u>	<u>A079148 001</u>	Nov 25, 2008
<u>AB</u>		<u>850MG</u>	<u>A079148 002</u>	Nov 25, 2008
<u>AB</u>		<u>1GM</u>	<u>A079148 003</u>	Nov 25, 2008
<u>AB</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A205330 001</u>	Oct 31, 2017
<u>AB</u>		<u>850MG</u>	<u>A205330 002</u>	Oct 31, 2017
<u>AB</u>		<u>1GM</u>	<u>A205330 003</u>	Oct 31, 2017
<u>AB</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090888 001</u>	Mar 12, 2012
<u>AB</u>		<u>850MG</u>	<u>A090888 002</u>	Mar 12, 2012
<u>AB</u>		<u>1GM</u>	<u>A090888 003</u>	Mar 12, 2012
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A075973 001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075976 001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>A075973 002</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075976 002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>A075973 003</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075976 003</u>	Jan 24, 2002
<u>AB</u>	PROVIDENT PHARM	<u>500MG</u>	<u>A077853 001</u>	Jul 28, 2006
<u>AB</u>		<u>850MG</u>	<u>A077853 002</u>	Jul 28, 2006
<u>AB</u>		<u>1GM</u>	<u>A077853 003</u>	Jul 28, 2006
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A075965 001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075985 001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075965 002</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075985 002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075965 003</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075985 003</u>	Jan 25, 2002
<u>AB</u>	SCIEGEN PHARMS INC	<u>500MG</u>	<u>A203769 001</u>	Sep 11, 2013
<u>AB</u>		<u>850MG</u>	<u>A203769 002</u>	Sep 11, 2013
<u>AB</u>		<u>1GM</u>	<u>A203769 003</u>	Sep 11, 2013
<u>AB</u>	SUN PHARM INDNS INC	<u>500MG</u>	<u>A075967 001</u>	Jan 29, 2002
<u>AB</u>		<u>850MG</u>	<u>A075967 002</u>	Jan 29, 2002
<u>AB</u>		<u>1GM</u>	<u>A075967 003</u>	Jan 29, 2002
<u>AB</u>	SUN PHARM INDUSTRIES	<u>500MG</u>	<u>A076038 001</u>	Feb 21, 2002
<u>AB</u>		<u>850MG</u>	<u>A076038 002</u>	Feb 21, 2002
<u>AB</u>		<u>1GM</u>	<u>A076038 003</u>	Feb 21, 2002
<u>AB</u>	TEVA	<u>500MG</u>	<u>A075978 001</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>A075978 002</u>	Jan 25, 2002
<u>AB</u>		<u>1GM</u>	<u>A075978 003</u>	Nov 05, 2002
<u>AB</u>	TORRENT PHARMS	<u>500MG</u>	<u>A077711 001</u>	Jan 24, 2007
<u>AB</u>		<u>850MG</u>	<u>A077711 002</u>	Jan 24, 2007
<u>AB</u>		<u>1GM</u>	<u>A077711 003</u>	Jan 24, 2007
<u>AB</u>	ZYDUS HLTHCARE	<u>500MG</u>	<u>A203686 001</u>	Aug 28, 2014
<u>AB</u>		<u>850MG</u>	<u>A203686 002</u>	Aug 28, 2014
<u>AB</u>		<u>1GM</u>	<u>A203686 003</u>	Aug 28, 2014
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077064 001</u>	Apr 18, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-276 (of 436)

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>		<u>850MG</u>	<u>A077064 002</u>	Apr 18, 2005
<u>AB</u>		<u>1GM</u>	<u>A077064 003</u>	Apr 18, 2005
	CHARTWELL LIFE SCI	625MG	A075972 005	Jan 24, 2002
		750MG	A075972 004	Jan 24, 2002

TABLET, EXTENDED RELEASE; ORAL

GLUCOPHAGE XR

<u>AB</u>	+!	BRISTOL MYERS SQUIBB	<u>750MG</u>	<u>N021202 004</u>	Apr 11, 2003
-----------	----	----------------------	--------------	--------------------	--------------

METFORMIN HYDROCHLORIDE

<u>AB</u>	ACTAVIS LABS FL INC	<u>750MG</u>	<u>A076869 001</u>	Apr 12, 2005
<u>AB</u>	AMNEAL PHARMS NY	<u>750MG</u>	<u>A078596 002</u>	Jan 03, 2008
<u>AB</u>	APOTEX	<u>750MG</u>	<u>A076706 002</u>	Dec 29, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>750MG</u>	<u>A079118 002</u>	Jul 20, 2012
<u>AB</u>	BARR	<u>750MG</u>	<u>A076863 001</u>	Oct 14, 2004
<u>AB</u>	BEXIMCO PHARMS USA	<u>750MG</u>	<u>A207427 002</u>	Dec 13, 2016
<u>AB</u>	CSPC OUYI PHARM CO	<u>750MG</u>	<u>A078321 002</u>	Apr 17, 2008
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>750MG</u>	<u>A202306 002</u>	Feb 23, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>750MG</u>	<u>A206955 002</u>	Dec 07, 2016
<u>AB</u>	MARKSANS PHARMA	<u>750MG</u>	<u>A090295 002</u>	Apr 29, 2016
<u>AB</u>	NOSTRUM PHARMS LLC	<u>750MG</u>	<u>A076756 002</u>	Dec 12, 2011
<u>AB</u>	SUN PHARM IND (IN)	<u>750MG</u>	<u>A077336 002</u>	Feb 09, 2006
<u>AB</u>	TEVA	<u>750MG</u>	<u>A076864 001</u>	Apr 12, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>750MG</u>	<u>A077078 001</u>	Apr 21, 2005

GLUCOPHAGE XR

<u>AB1</u>	+	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N021202 001</u>	Oct 13, 2000
------------	---	----------------------	--------------	--------------------	--------------

METFORMIN HYDROCHLORIDE

<u>AB1</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A076172 001</u>	Jun 16, 2004
<u>AB1</u>	AMNEAL PHARMS NY	<u>500MG</u>	<u>A078596 001</u>	Jan 03, 2008
<u>AB1</u>	APOTEX	<u>500MG</u>	<u>A076706 001</u>	Dec 14, 2004
<u>AB1</u>	AUROBINDO PHARMA LTD	<u>500MG</u>	<u>A079118 001</u>	Jul 20, 2012
<u>AB1</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A207427 001</u>	Dec 13, 2016
<u>AB1</u>	CSPC OUYI PHARM CO	<u>500MG</u>	<u>A078321 001</u>	Apr 17, 2008
<u>AB1</u>	INTELLIPHARMACEUTIC S	<u>500MG</u>	<u>A202306 001</u>	Feb 23, 2017
<u>AB1</u>	INVENTIA HLTHCARE	<u>500MG</u>	<u>A201991 001</u>	Jan 18, 2012
<u>AB1</u>	MACLEODS PHARMS LTD	<u>500MG</u>	<u>A206955 001</u>	Dec 07, 2016
<u>AB1</u>	MARKSANS PHARMA	<u>500MG</u>	<u>A090295 001</u>	Apr 29, 2016
<u>AB1</u>	NOSTRUM PHARMS LLC	<u>500MG</u>	<u>A076756 001</u>	Jul 26, 2006
<u>AB1</u>	SANDOZ	<u>500MG</u>	<u>A076873 001</u>	Dec 14, 2004
<u>AB1</u>	SUN PHARM IND (IN)	<u>500MG</u>	<u>A077336 001</u>	Feb 09, 2006
<u>AB1</u>	TEVA	<u>500MG</u>	<u>A076269 001</u>	Jun 18, 2004
<u>AB1</u>	TORRENT PHARMS LTD	<u>500MG</u>	<u>A090014 001</u>	Dec 30, 2009
<u>AB1</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>A077060 001</u>	Apr 20, 2005

FORTAMET

<u>AB2</u>	+	ANDRX LABS LLC	<u>500MG</u>	<u>N021574 001</u>	Apr 27, 2004
<u>AB2</u>	+!		<u>1GM</u>	<u>N021574 002</u>	Apr 27, 2004

METFORMIN HYDROCHLORIDE

<u>AB2</u>	LUPIN LTD	<u>500MG</u>	<u>A090692 001</u>	Jun 29, 2011
<u>AB2</u>		<u>1GM</u>	<u>A090692 002</u>	Jun 29, 2011
<u>AB2</u>	MYLAN PHARMS INC	<u>500MG</u>	<u>A200690 001</u>	Aug 01, 2012
<u>AB2</u>		<u>1GM</u>	<u>A200690 002</u>	Aug 01, 2012
<u>AB2</u>	NOSTRUM LABS INC	<u>500MG</u>	<u>A203832 001</u>	Dec 26, 2017
<u>AB2</u>		<u>1GM</u>	<u>A203832 002</u>	Dec 26, 2017

GLUMETZA

<u>AB3</u>	+	SANTARUS INC	<u>500MG</u>	<u>N021748 001</u>	Jun 03, 2005
<u>AB3</u>	+!		<u>1GM</u>	<u>N021748 002</u>	Jun 03, 2005

METFORMIN HYDROCHLORIDE

<u>AB3</u>	ACTAVIS LABS FL INC	<u>500MG</u>	<u>A203755 001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A203755 002</u>	Aug 01, 2016
<u>AB3</u>	LUPIN LTD	<u>500MG</u>	<u>A091664 001</u>	Jul 19, 2013
<u>AB3</u>		<u>1GM</u>	<u>A091664 002</u>	Jul 19, 2013
<u>AB3</u>	SUN PHARMA GLOBAL	<u>500MG</u>	<u>A202917 001</u>	Aug 01, 2016
<u>AB3</u>		<u>1GM</u>	<u>A202917 002</u>	Aug 01, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-277 (of 436)

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

**ACTOPLUS MET**

<u>AB</u>	+	TAKEDA PHARMS USA	<u>500MG;EQ 15MG BASE</u>	<u>N021842 001</u>	Aug 29, 2005
<u>AB</u>	+		<u>850MG;EQ 15MG BASE</u>	<u>N021842 002</u>	Aug 29, 2005

**PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE**

<u>AB</u>	AUROBINDO PHARMA LTD	<u>500MG;EQ 15MG BASE</u>	<u>A200823 001</u>	Feb 13, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A200823 002</u>	Feb 13, 2013
<u>AB</u>	MACLEODS PHARMS LTD	<u>500MG;EQ 15MG BASE</u>	<u>A204802 001</u>	Nov 05, 2015
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A204802 002</u>	Nov 05, 2015
<u>AB</u>	MYLAN	<u>500MG;EQ 15MG BASE</u>	<u>A090406 001</u>	Feb 25, 2011
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A090406 002</u>	Feb 25, 2011
<u>AB</u>	SANDOZ	<u>500MG;EQ 15MG BASE</u>	<u>A091273 001</u>	Apr 16, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A091273 002</u>	Apr 16, 2013
<u>AB</u>	TEVA PHARMS USA	<u>500MG;EQ 15MG BASE</u>	<u>A091155 001</u>	Mar 10, 2014
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A091155 002</u>	Mar 10, 2014
<u>AB</u>	TORRENT PHARMS LTD	<u>500MG;EQ 15MG BASE</u>	<u>A202001 001</u>	Feb 13, 2013
<u>AB</u>		<u>850MG;EQ 15MG BASE</u>	<u>A202001 002</u>	Feb 13, 2013

TABLET, EXTENDED RELEASE; ORAL  
**ACTOPLUS MET XR**

+	TAKEDA PHARMS USA	1GM;EQ 15MG BASE	N022024 001	May 12, 2009
+		1GM;EQ 30MG BASE	N022024 002	May 12, 2009

METFORMIN HYDROCHLORIDE; REPAGLINIDE

TABLET; ORAL

REPAGLINIDE AND METFORMIN HYDROCHLORIDE

LUPIN LTD	500MG;1MG	A200624 001	Jul 15, 2015
!	500MG;2MG	A200624 002	Jul 15, 2015

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

ROSIGLITAZONE MALEATE AND METFORMIN HYDROCHLORIDE

TEVA	500MG;EQ 2MG BASE	A077337 001	May 07, 2014
	500MG;EQ 1MG BASE	A077337 005	May 19, 2017
	500MG;EQ 4MG BASE	A077337 002	May 07, 2014
!	1GM;EQ 4MG BASE	A077337 004	May 07, 2014
	1GM;EQ 2MG BASE	A077337 003	May 07, 2014

METFORMIN HYDROCHLORIDE; SAXagliptin HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

KOMBIGLYZE XR

+	ASTRAZENECA AB	500MG;EQ 5MG BASE	N200678 001	Nov 05, 2010
+		1GM;EQ 2.5MG BASE	N200678 003	Nov 05, 2010
+		1GM;EQ 5MG BASE	N200678 002	Nov 05, 2010

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUMET

+	MERCK SHARP DOHME	500MG;EQ 50MG BASE	N022044 001	Mar 30, 2007
+		1GM;EQ 50MG BASE	N022044 002	Mar 30, 2007

TABLET, EXTENDED RELEASE; ORAL

JANUMET XR

+	MERCK SHARP DOHME	500MG;EQ 50MG BASE	N202270 001	Feb 02, 2012
+		1GM;EQ 50MG BASE	N202270 002	Feb 02, 2012
+		1GM;EQ 100MG BASE	N202270 003	Feb 02, 2012

METHACHOLINE CHLORIDE

FOR SOLUTION; INHALATION

PROVOCHOLINE

+	METHAPHARM	100MG/VIAL	N019193 001	Oct 31, 1986
---	------------	------------	-------------	--------------

METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

**METHADONE HYDROCHLORIDE**

<u>AA</u>	VISTAPHARM	<u>10MG/ML</u>	<u>A040088 001</u>	Nov 30, 1994
<u>AA</u>	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A040180 001</u>	Apr 30, 1998

**METHADONE HYDROCHLORIDE INTENSOL**

<u>AA</u>	WEST-WARD PHARMS INT	<u>10MG/ML</u>	<u>A089897 001</u>	Sep 06, 1988
-----------	----------------------	----------------	--------------------	--------------

**METHADOSE**

<u>AA</u>	+! SPECGX LLC	<u>10MG/ML</u>	<u>N017116 002</u>
-----------	---------------	----------------	--------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-278 (of 436)

METHADONE HYDROCHLORIDE

INJECTABLE; INJECTION

METHADONE HYDROCHLORIDE

<u>AP</u>	AKORN	<u>10MG/ML</u>	<u>A208306 001</u> Oct 27, 2017
<u>AP</u> +!	MYLAN INSTITUTIONAL	<u>10MG/ML</u>	<u>N021624 001</u>

SOLUTION; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	VISTAPHARM	<u>5MG/5ML</u>	<u>A090707 001</u> Jun 30, 2010
<u>AA</u>		<u>10MG/5ML</u>	<u>A090707 002</u> Jun 30, 2010
<u>AA</u> !	WEST-WARD PHARMS INT	<u>5MG/5ML</u>	<u>A087393 001</u>
<u>AA</u> !		<u>10MG/5ML</u>	<u>A087997 001</u> Aug 30, 1982

TABLET; ORAL

DOLOPHINE HYDROCHLORIDE

<u>AA</u> +!	WEST-WARD PHARMS INT	<u>5MG</u>	<u>N006134 002</u>
<u>AA</u> +!		<u>10MG</u>	<u>N006134 010</u>

METHADONE HYDROCHLORIDE

<u>AA</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A203502 001</u> Aug 31, 2015
<u>AA</u>		<u>10MG</u>	<u>A203502 002</u> Aug 31, 2015
<u>AA</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A090065 001</u> Aug 18, 2015
<u>AA</u>		<u>10MG</u>	<u>A090065 002</u> Aug 18, 2015
<u>AA</u>	SANDOZ	<u>10MG</u>	<u>A04241 002</u> May 29, 1998
<u>AA</u>	SPECGX LLC	<u>5MG</u>	<u>A040517 001</u> Apr 27, 2004
<u>AA</u>		<u>10MG</u>	<u>A040517 002</u> Apr 27, 2004
<u>AA</u>	THE PHARMANETWORK	<u>10MG</u>	<u>A090635 001</u> Nov 25, 2009

METHADOSE

<u>AA</u>	SPECGX LLC	<u>5MG</u>	<u>A040050 001</u> Apr 15, 1993
<u>AA</u>		<u>10MG</u>	<u>A040050 002</u> Apr 15, 1993

TABLET, FOR SUSPENSION; ORAL

METHADONE HYDROCHLORIDE

<u>AA</u>	SANDOZ	<u>40MG</u>	<u>A075082 001</u> Mar 25, 1998
<u>AA</u>	SPECGX LLC	<u>40MG</u>	<u>A077142 001</u> Jul 12, 2005
<u>AA</u> +!	WEST-WARD PHARMS INT	<u>40MG</u>	<u>N017058 001</u>

METHADOSE

<u>AA</u>	SPECGX LLC	<u>40MG</u>	<u>A074184 001</u> Apr 29, 1993
-----------	------------	-------------	---------------------------------

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DESOXYN

<u>AA</u> +!	RECORDATI RARE	<u>5MG</u>	<u>N005378 002</u>
--------------	----------------	------------	--------------------

METHAMPHETAMINE HYDROCHLORIDE

<u>AA</u>	MAYNE PHARMA INC	<u>5MG</u>	<u>A091189 001</u> Apr 21, 2010
<u>AA</u>	WEST-WARD PHARMS INT	<u>5MG</u>	<u>A203846 001</u> Nov 17, 2015

METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

<u>AB</u>	ANI PHARMS INC	<u>25MG</u>	<u>A040001 001</u> Jun 30, 1993
<u>AB</u>		<u>50MG</u>	<u>A040001 002</u> Jun 30, 1993
<u>AB</u>	MIKART	<u>25MG</u>	<u>A040062 001</u> Jan 27, 1994
<u>AB</u> !		<u>50MG</u>	<u>A040062 002</u> Jan 27, 1994
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A040036 001</u> Jun 30, 1993
<u>AB</u>		<u>50MG</u>	<u>A040036 002</u> Jun 30, 1993

METHENAMINE HIPPURATE

TABLET; ORAL

HIPREX

<u>AB</u> +!	US PHARM HOLDINGS	<u>1GM</u>	<u>N017681 001</u>
<u>AB</u>	AUROBINDO PHARMA LTD	<u>1GM</u>	<u>A205661 001</u> Jul 05, 2016
<u>AB</u>	IMPAX LABS INC	<u>1GM</u>	<u>A076411 001</u> Jun 20, 2003
<u>AB</u>	UREX		
<u>AB</u>	CNTY LINE PHARMS	<u>1GM</u>	<u>N016151 001</u>

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

<u>AB</u>	ECI PHARMS LLC	<u>5MG</u>	<u>A040547 001</u> Feb 18, 2005
<u>AB</u>		<u>10MG</u>	<u>A040547 002</u> Feb 18, 2005
<u>AB</u>	HERITAGE PHARMA	<u>5MG</u>	<u>A040734 001</u> Dec 14, 2007
<u>AB</u>		<u>10MG</u>	<u>A040734 002</u> Dec 14, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-279 (of 436)

METHIMAZOLE

TABLET;ORAL

METHIMAZOLE

<u>AB</u>	MYLAN	<u>5MG</u>	<u>A040350</u> <u>001</u>	Mar 29, 2000
<u>AB</u>	!	<u>10MG</u>	<u>A040350</u> <u>002</u>	Mar 29, 2000
<u>AB</u>	RISING PHARMS INC	<u>5MG</u>	<u>A202068</u> <u>001</u>	Mar 07, 2012
<u>AB</u>		<u>10MG</u>	<u>A202068</u> <u>002</u>	Mar 07, 2012
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>A040411</u> <u>001</u>	Mar 27, 2001
<u>AB</u>		<u>10MG</u>	<u>A040411</u> <u>002</u>	Mar 27, 2001
<u>AB</u>	SUN PHARM INDNS INC	<u>5MG</u>	<u>A040870</u> <u>001</u>	Sep 25, 2007
<u>AB</u>		<u>10MG</u>	<u>A040870</u> <u>002</u>	Sep 25, 2007
<u>TAPAZOLE</u>				
<u>AB</u>	KING PHARMS LLC	<u>5MG</u>	<u>A040320</u> <u>001</u>	Mar 31, 2000
<u>AB</u>		<u>10MG</u>	<u>A040320</u> <u>002</u>	Mar 31, 2000

METHOCARBAMOL

SOLUTION; IM-IV

METHOCARBAMOL

<u>AP</u>	AUROBINDO PHARMA LTD	<u>1GM/10ML (100MG/ML)</u>	<u>A206128</u> <u>001</u>	May 27, 2016
<u>AP</u>	LUITPOLD PHARMS INC	<u>1GM/10ML (100MG/ML)</u>	<u>A207496</u> <u>001</u>	Jun 22, 2017
<u>AP</u>	MONTEREY PHARMS LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A205354</u> <u>001</u>	Oct 27, 2016
<u>AP</u>	MYLAN INSTITUTIONAL	<u>1GM/10ML (100MG/ML)</u>	<u>A204404</u> <u>001</u>	Dec 05, 2014
<u>AP</u>	NAVINTA LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A206071</u> <u>001</u>	Nov 24, 2017
<u>AP</u>	RENAISSANCE SSA LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A208116</u> <u>001</u>	Jan 19, 2017
<u>AP</u>	SAGENT PHARMS	<u>1GM/10ML (100MG/ML)</u>	<u>A205404</u> <u>001</u>	Jul 18, 2017
<u>AP</u>	SOMERSET THERAPS LLC	<u>1GM/10ML (100MG/ML)</u>	<u>A207522</u> <u>001</u>	Jul 31, 2017

ROBAXIN

<u>AP</u>	+! WEST-WARD PHARMS INT	<u>1GM/10ML (100MG/ML)</u>	<u>N011790</u> <u>001</u>
-----------	-------------------------	----------------------------	---------------------------

TABLET;ORAL

METHOCARBAMOL

<u>AA</u>	ATLAS PHARMS LLC	<u>500MG</u>	<u>A203550</u> <u>001</u>	Feb 08, 2017
<u>AA</u>		<u>750MG</u>	<u>A203550</u> <u>002</u>	Feb 08, 2017
<u>AA</u>	AUSTARPHARMA LLC	<u>500MG</u>	<u>A200958</u> <u>001</u>	Oct 21, 2011
<u>AA</u>		<u>750MG</u>	<u>A200958</u> <u>002</u>	Oct 21, 2011
<u>AA</u>	BEXIMCO PHARMS USA	<u>500MG</u>	<u>A208507</u> <u>001</u>	Jul 21, 2017
<u>AA</u>		<u>750MG</u>	<u>A208507</u> <u>002</u>	Jul 21, 2017
<u>AA</u>	HETERO LABS LTD III	<u>500MG</u>	<u>A090200</u> <u>001</u>	Nov 06, 2009
<u>AA</u>		<u>750MG</u>	<u>A090200</u> <u>002</u>	Nov 06, 2009
<u>AA</u>	HIKMA INTL PHARMS	<u>500MG</u>	<u>A085159</u> <u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A085123</u> <u>001</u>	
<u>AA</u>	PRINSTON INC	<u>500MG</u>	<u>A086989</u> <u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A086988</u> <u>001</u>	
<u>AA</u>	VINTAGE PHARMS	<u>500MG</u>	<u>A040489</u> <u>001</u>	Jan 29, 2003
<u>AA</u>		<u>750MG</u>	<u>A040489</u> <u>002</u>	Jan 29, 2003
<u>AA</u>	WATSON LABS	<u>500MG</u>	<u>A084277</u> <u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>A084276</u> <u>002</u>	

ROBAXIN

<u>AA</u>	+! AUXILIUM PHARMS LLC	<u>500MG</u>	<u>N011011</u> <u>004</u>
-----------	------------------------	--------------	---------------------------

ROBAXIN-750

<u>AA</u>	+! AUXILIUM PHARMS LLC	<u>750MG</u>	<u>N011011</u> <u>006</u>
-----------	------------------------	--------------	---------------------------

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

+!	PAR STERILE PRODUCTS	500MG/VIAL	N011559	001
+!		2.5GM/VIAL	N011559	002

METHOTREXATE

SOLUTION; SUBCUTANEOUS

OTREXUP

+!	ANTARES PHARMA INC	10MG/0.4ML (10MG/0.4ML)	N204824	001	Oct 11, 2013
+!		12.5MG/0.4ML (12.5MG/0.4ML)	N204824	006	Mar 24, 2016
+!		15MG/0.4ML (15MG/0.4ML)	N204824	002	Oct 11, 2013
+!		17.5MG/0.4ML (17.5MG/0.4ML)	N204824	007	Mar 24, 2016
+!		20MG/0.4ML (20MG/0.4ML)	N204824	003	Oct 11, 2013
+!		22.5MG/0.4ML (22.5MG/0.4ML)	N204824	008	Mar 24, 2016
+!		25MG/0.4ML (25MG/0.4ML)	N204824	004	Oct 11, 2013

RASUVO

+!	MEDAC PHARMA INC	7.5MG/0.15ML (7.5MG/0.15ML)	N205776	001	Jul 10, 2014
+!		10MG/0.20ML (10MG/0.20ML)	N205776	002	Jul 10, 2014
+!		12.5MG/0.25ML (12.5MG/0.25ML)	N205776	003	Jul 10, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-280 (of 436)

METHOTREXATE

SOLUTION; SUBCUTANEOUS

RASUVO

+	15MG/0.30ML (15MG/0.30ML)	N205776 004 Jul 10, 2014
+	17.5MG/0.35ML (17.5MG/0.35ML)	N205776 005 Jul 10, 2014
+	20MG/0.4ML (20MG/0.4ML)	N205776 006 Jul 10, 2014
+	22.5MG/0.45ML (22.5MG/0.45ML)	N205776 007 Jul 10, 2014
+	25MG/0.5ML (25MG/0.5ML)	N205776 008 Jul 10, 2014
+	30MG/0.6ML (30MG/0.6ML)	N205776 010 Jul 10, 2014

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE PRESERVATIVE FREE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 25MG BASE/ML</u>	A040265 001 Feb 26, 1999
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	A040266 001 Feb 26, 1999

METHOTREXATE SODIUM

<u>AP</u>	!	FRESENIUS KABI USA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A040263 001 Feb 26, 1999
<u>AP</u>	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A040263 002 Feb 26, 1999
<u>AP</u>	+!	HOSPIRA	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	N011719 010 Dec 15, 2004
<u>AP</u>	!	WEST-WARD PHARMS INT	<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	A089341 001 Sep 16, 1986

METHOTREXATE SODIUM PRESERVATIVE FREE

<u>AP</u>	!	ACCORD HLTHCARE	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A040767 001 Apr 30, 2007
<u>AP</u>	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A040768 001 Apr 30, 2007
<u>AP</u>	!		<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A040716 001 Apr 30, 2007
<u>AP</u>	+!	HOSPIRA	<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	N011719 012 Apr 13, 2005
<u>AP</u>		MYLAN LABS LTD	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A201529 001 Mar 29, 2012
<u>AP</u>			<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	A201529 002 Mar 29, 2012
<u>AP</u>			<u>EQ 200MG BASE/8ML (EQ 25MG BASE/ML)</u>	A201529 003 Mar 29, 2012
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A201529 004 Mar 29, 2012
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A201530 001 Mar 29, 2012
<u>AP</u>		PHARMACHEMIE BV	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A040843 002 Jan 11, 2010
<u>AP</u>			<u>EQ 100MG BASE/4ML (EQ 25MG BASE/ML)</u>	A040843 003 Feb 27, 2012
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A040843 004 Jan 11, 2010
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A040843 001 Jan 11, 2010
<u>AP</u>		SANDOZ INC	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A090039 001 Mar 31, 2009
<u>AP</u>			<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A090039 002 Mar 31, 2009
<u>AP</u>			<u>EQ 1GM BASE/40ML (EQ 25MG BASE/ML)</u>	A090029 001 Mar 31, 2009
<u>AP</u>	!	WEST-WARD PHARMS INT	<u>EQ 50MG BASE/2ML (EQ 25MG BASE/ML)</u>	A089340 001 Sep 16, 1986
<u>AP</u>	!		<u>EQ 250MG BASE/10ML (EQ 25MG BASE/ML)</u>	A089343 001 Sep 16, 1986

METHOTREXATE SODIUM

!	WEST-WARD PHARMS INT	EQ 200MG BASE/8ML (EQ 25MG BASE/ML)
---	----------------------	-------------------------------------

METHOTREXATE SODIUM PRESERVATIVE FREE

!	WEST-WARD PHARMS INT	EQ 1GM BASE/VIAL
---	----------------------	------------------

A040632 001 Aug 12, 2005

SOLUTION; ORAL

XATMEP

+!	SILVERGATE PHARMS	EQ 2.5MG BASE/ML
----	-------------------	------------------

N208400 001 Apr 25, 2017

TABLET; ORAL

METHOTREXATE SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>EQ 2.5MG BASE</u>	
<u>AB</u>	BARR	<u>EQ 2.5MG BASE</u>	
<u>AB</u>	+!	DAVA PHARMS INC	<u>EQ 2.5MG BASE</u>
<u>AB</u>		MYLAN	<u>EQ 2.5MG BASE</u>
<u>AB</u>		SUN PHARMA GLOBAL	<u>EQ 2.5MG BASE</u>
<u>AB</u>		WEST-WARD PHARMS INT	<u>EQ 2.5MG BASE</u>
<u>AB</u>		ZYDUS PHARMS USA INC	<u>EQ 2.5MG BASE</u>

A210040 001 Dec 22, 2017

A081099 001 Oct 15, 1990

N008085 002

A081235 001 May 15, 1992

A201749 001 May 21, 2015

A040054 001 Aug 01, 1994

A207812 001 Jan 13, 2017

TREXALL

BARR	EQ 5MG BASE
	EQ 7.5MG BASE
	EQ 10MG BASE
!	EQ 15MG BASE

A040385 001 Mar 21, 2001

A040385 002 Mar 21, 2001

A040385 003 Mar 21, 2001

A040385 004 Mar 21, 2001

METHOXSALEN

CAPSULE; ORAL

METHOXSALEN

<u>AB</u>	ACTAVIS INC	<u>10MG</u>
<u>AB</u>	STRIDES PHARMA	<u>10MG</u>
<u>AB</u>	OXSORALEN-ULTRA	
<u>AB</u>	+! DOW PHARM	<u>10MG</u>

A202603 001 Jun 09, 2015

A202687 001 Jun 05, 2014

N019600 001 Oct 30, 1986

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-281 (of 436)

METHOXSALEN

INJECTABLE; INJECTION

UVADEX

+! MALLINCKRODT HOSP 0.02MG/ML

N020969 001 Feb 25, 1999

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

<u>AA</u>	BAYSHORE PHARMS LLC	<u>2.5MG</u>	<u>A200602 001</u>	Sep 24, 2012
<u>AA</u>		<u>5MG</u>	<u>A200602 002</u>	Sep 24, 2012
<u>AA</u>	BRECKENRIDGE PHARM	<u>2.5MG</u>	<u>A040642 001</u>	Dec 06, 2011
<u>AA</u>		<u>5MG</u>	<u>A040642 002</u>	Dec 06, 2011
<u>AA</u> !	VINTAGE PHARMS	<u>2.5MG</u>	<u>A040624 001</u>	Dec 28, 2006
<u>AA</u> !		<u>5MG</u>	<u>A040624 002</u>	Dec 28, 2006

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

+ PARKE DAVIS 150MG  
+! 300MG

N010596 007  
N010596 008

METHYCLOTHIAZIDE

TABLET; ORAL

METHYCLOTHIAZIDE

! MYLAN PHARMS INC 5MG

A087672 001 Aug 17, 1982

METHYLDOPA

TABLET; ORAL

METHYLDOPA

<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A070084 001</u>	Oct 15, 1985
<u>AB</u>		<u>500MG</u>	<u>A070085 001</u>	Oct 15, 1985
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>250MG</u>	<u>A070098 001</u>	Feb 20, 1986
<u>AB</u>		<u>500MG</u>	<u>A070343 001</u>	Feb 20, 1986
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A070076 002</u>	Apr 18, 1985
<u>AB</u> !		<u>500MG</u>	<u>A070076 001</u>	Apr 18, 1985
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A070625 001</u>	Jun 06, 1986

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

METHYLDOPATE HYDROCHLORIDE

! LUITPOLD 50MG/ML

A071279 001 Oct 02, 1987

METHYLENE BLUE

SOLUTION; INTRAVENOUS

PROVAYBLUE

+! PROVEPHARM SAS 50MG/10ML (5MG/ML)

N204630 001 Apr 08, 2016

METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHSERGINE

<u>AP</u> +!	EDISON THERAPS LLC	<u>0.2MG/ML</u>	<u>N006035 004</u>	
<u>AP</u>	METHYLERGONOVINE MALEATE	<u>0.2MG/ML</u>	<u>A040889 001</u>	Sep 13, 2010
<u>AP</u>	BRECKENRIDGE PHARM	<u>0.2MG/ML</u>	<u>A090193 001</u>	Nov 24, 2008
<u>AP</u>	LUITPOLD	<u>0.2MG/ML</u>		
TABLET; ORAL				
METHYLERGONOVINE MALEATE				
! NOVEL LABS INC	0.2MG			
				A091577 001 May 02, 2011

METHYLNALTREXONE BROMIDE

SOLUTION; SUBCUTANEOUS

RELISTOR

+! SALIX PHARMS 8MG/0.4ML (8MG/0.4ML)  
+! 12MG/0.6ML (12MG/0.6ML)  
+! 12MG/0.6ML (12MG/0.6ML)

N021964 002 Sep 27, 2010  
N021964 001 Apr 24, 2008  
N021964 003 Sep 27, 2010

TABLET; ORAL

RELISTOR

+! SALIX PHARMS INC 150MG

N208271 001 Jul 19, 2016

METHYLPHENIDATE

FILM, EXTENDED RELEASE; TRANSDERMAL

DAYTRANA

+ NOVEN PHARMS INC 10MG/9HR (1.1MG/HR)  
+ 15MG/9HR (1.6MG/HR)  
+ 20MG/9HR (2.2MG/HR)  
+! 30MG/9HR (3.3MG/HR)

N021514 001 Apr 06, 2006  
N021514 002 Apr 06, 2006  
N021514 003 Apr 06, 2006  
N021514 004 Apr 06, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-282 (of 436)

METHYLPHENIDATE

TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE;ORAL  
 COTEMPLA XR-ODT  
 + NEOS THERAPS INC 8.6MG  
 + 17.3MG  
 + 25.9MG

N205489 001 Jun 19, 2017  
 N205489 002 Jun 19, 2017  
 N205489 003 Jun 19, 2017

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB1</u>	BARR LABS INC	<u>10MG</u>
<u>AB1</u>		<u>20MG</u>
<u>AB1</u>		<u>30MG</u>
<u>AB1</u>		<u>40MG</u>
<u>AB1</u>	MAYNE PHARMA	<u>20MG</u>
<u>AB1</u>		<u>30MG</u>
<u>AB1</u>		<u>40MG</u>

<u>A079031</u> <u>004</u>	Oct 15, 2014
<u>A079031</u> <u>001</u>	Jul 13, 2012
<u>A079031</u> <u>002</u>	Jul 13, 2012
<u>A079031</u> <u>003</u>	Jul 13, 2012
<u>A078458</u> <u>001</u>	Dec 01, 2011
<u>A078458</u> <u>002</u>	Dec 01, 2011
<u>A078458</u> <u>003</u>	Dec 01, 2011

RITALIN LA

<u>AB1</u> + NOVARTIS	<u>10MG</u>
<u>AB1</u> +	<u>20MG</u>
<u>AB1</u> +	<u>30MG</u>
<u>AB1</u> +	<u>40MG</u>

<u>N021284</u> <u>004</u>	Apr 10, 2004
<u>N021284</u> <u>001</u>	Jun 05, 2002
<u>N021284</u> <u>002</u>	Jun 05, 2002
<u>N021284</u> <u>003</u>	Jun 05, 2002

METADATE CD

<u>AB2</u> + UCB INC	<u>10MG</u>
<u>AB2</u> +	<u>20MG</u>
<u>AB2</u> +	<u>30MG</u>
<u>AB2</u> +	<u>40MG</u>
<u>AB2</u> +	<u>50MG</u>
<u>AB2</u> +!	<u>60MG</u>

<u>N021259</u> <u>003</u>	May 27, 2003
<u>N021259</u> <u>001</u>	Apr 03, 2001
<u>N021259</u> <u>002</u>	Jun 19, 2003
<u>N021259</u> <u>004</u>	Feb 19, 2006
<u>N021259</u> <u>005</u>	Feb 19, 2006
<u>N021259</u> <u>006</u>	Feb 19, 2006

METHYLPHENIDATE HYDROCHLORIDE

<u>AB2</u> IMPAX LABS INC	<u>10MG</u>
<u>AB2</u>	<u>20MG</u>
<u>AB2</u>	<u>30MG</u>
<u>AB2</u>	<u>40MG</u>
<u>AB2</u>	<u>50MG</u>
<u>AB2</u>	<u>60MG</u>

<u>A205105</u> <u>001</u>	Jul 28, 2016
<u>A205105</u> <u>002</u>	Jul 28, 2016
<u>A205105</u> <u>003</u>	Jul 28, 2016
<u>A205105</u> <u>004</u>	Jul 28, 2016
<u>A205105</u> <u>005</u>	Jul 28, 2016
<u>A205105</u> <u>006</u>	Jul 28, 2016

SPECGX LLC

<u>AB2</u>	<u>10MG</u>
<u>AB2</u>	<u>20MG</u>
<u>AB2</u>	<u>30MG</u>
<u>AB2</u>	<u>40MG</u>
<u>AB2</u>	<u>50MG</u>
<u>AB2</u>	<u>60MG</u>

<u>A203583</u> <u>001</u>	Sep 29, 2015
<u>A203583</u> <u>002</u>	Sep 29, 2015
<u>A203583</u> <u>003</u>	Sep 29, 2015
<u>A203583</u> <u>004</u>	Sep 29, 2015
<u>A203583</u> <u>005</u>	Sep 29, 2015
<u>A203583</u> <u>006</u>	Sep 29, 2015

TEVA PHARMS

<u>AB2</u>	<u>10MG</u>
<u>AB2</u>	<u>20MG</u>
<u>AB2</u>	<u>30MG</u>
<u>AB2</u>	<u>40MG</u>
<u>AB2</u>	<u>50MG</u>
<u>AB2</u>	<u>60MG</u>

<u>A077707</u> <u>001</u>	Jul 19, 2012
<u>A077707</u> <u>002</u>	Jul 19, 2012
<u>A077707</u> <u>003</u>	Jul 19, 2012
<u>A078873</u> <u>001</u>	Jul 19, 2012
<u>A078873</u> <u>002</u>	Jul 19, 2012
<u>A078873</u> <u>003</u>	Jul 19, 2012

APTENSIO XR

+ RHODES PHARMS	10MG
+	15MG
+	20MG
+	30MG
+	40MG
+	50MG
+!	60MG

N205831 001	Apr 17, 2015
N205831 002	Apr 17, 2015
N205831 003	Apr 17, 2015
N205831 004	Apr 17, 2015
N205831 005	Apr 17, 2015
N205831 006	Apr 17, 2015
N205831 007	Apr 17, 2015

METHYLPHENIDATE HYDROCHLORIDE

! MAYNE PHARMA 60MG

A078458 004 Jun 23, 2016

FOR SUSPENSION, EXTENDED RELEASE;ORAL

QUILLIVANT XR

+! NEXTWAVE PHARMS 5MG/ML

N202100 001 Sep 27, 2012

SOLUTION;ORAL

METHYLINE

<u>AA</u> +! SPECGX LLC	<u>5MG/5ML</u>
<u>AA</u> +!	<u>10MG/5ML</u>

<u>N021419</u> <u>001</u>	Dec 19, 2002
<u>N021419</u> <u>002</u>	Dec 19, 2002

METHYLPHENIDATE HYDROCHLORIDE

<u>AA</u>	ABHAI LLC	<u>5MG/5ML</u>
<u>AA</u>		<u>10MG/5ML</u>
<u>AA</u>	BRECKENRIDGE PHARM	<u>5MG/5ML</u>
<u>AA</u>		<u>10MG/5ML</u>
<u>AA</u>	NOVEL LABS INC	<u>5MG/5ML</u>
<u>AA</u>		<u>10MG/5ML</u>
<u>AA</u>	TRIS PHARMA INC	<u>5MG/5ML</u>

<u>A207485</u> <u>001</u>	Nov 18, 2016
<u>A207485</u> <u>002</u>	Nov 18, 2016
<u>A201466</u> <u>001</u>	Nov 12, 2013
<u>A201466</u> <u>002</u>	Nov 12, 2013
<u>A204602</u> <u>001</u>	Aug 14, 2015
<u>A204602</u> <u>002</u>	Aug 14, 2015
<u>A091601</u> <u>001</u>	Jul 23, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-283 (of 436)

METHYLPHENIDATE HYDROCHLORIDE

SOLUTION;ORAL

METHYLPHENIDATE HYDROCHLORIDE

AA 10MG/5ML

A091601 002 Jul 23, 2010

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ABHAI INC	<u>5MG</u>	<u>A206932 001</u>	May 11, 2017
<u>AB</u>		<u>10MG</u>	<u>A206932 002</u>	May 11, 2017
<u>AB</u>		<u>20MG</u>	<u>A206932 003</u>	May 11, 2017
<u>AB</u>	ACTAVIS LABS FL INC	<u>5MG</u>	<u>A040220 001</u>	Aug 29, 1997
<u>AB</u>		<u>10MG</u>	<u>A040220 002</u>	Aug 29, 1997
<u>AB</u>		<u>20MG</u>	<u>A040220 003</u>	Aug 29, 1997
<u>AB</u>	ASCENT PHARMS INC	<u>5MG</u>	<u>A207416 001</u>	Sep 22, 2015
<u>AB</u>		<u>10MG</u>	<u>A207416 002</u>	Sep 22, 2015
<u>AB</u>		<u>20MG</u>	<u>A207416 003</u>	Sep 22, 2015
<u>AB</u>	BRECKENRIDGE PHARM	<u>5MG</u>	<u>A207587 001</u>	Mar 03, 2017
<u>AB</u>		<u>10MG</u>	<u>A207587 002</u>	Mar 03, 2017
<u>AB</u>		<u>20MG</u>	<u>A207587 003</u>	Mar 03, 2017
<u>AB</u>	CNTY LINE PHARMS	<u>5MG</u>	<u>A206840 001</u>	Sep 15, 2016
<u>AB</u>		<u>10MG</u>	<u>A206840 002</u>	Sep 15, 2016
<u>AB</u>		<u>20MG</u>	<u>A206840 003</u>	Sep 15, 2016
<u>AB</u>	COREPHARMA	<u>5MG</u>	<u>A091159 001</u>	Mar 12, 2014
<u>AB</u>		<u>10MG</u>	<u>A091159 002</u>	Mar 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A091159 003</u>	Mar 12, 2014
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A207884 001</u>	Nov 13, 2015
<u>AB</u>		<u>10MG</u>	<u>A207884 002</u>	Nov 13, 2015
<u>AB</u>		<u>20MG</u>	<u>A207884 003</u>	Nov 13, 2015
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A040300 001</u>	Nov 27, 1998
<u>AB</u>		<u>10MG</u>	<u>A040300 002</u>	Nov 27, 1998
<u>AB</u>		<u>20MG</u>	<u>A040300 003</u>	Nov 27, 1998
<u>AB</u>	SUN PHARM INDs INC	<u>5MG</u>	<u>A090710 001</u>	Mar 15, 2012
<u>AB</u>		<u>10MG</u>	<u>A090710 002</u>	Mar 15, 2012
<u>AB</u>		<u>20MG</u>	<u>A090710 003</u>	Mar 15, 2012
<u>AB</u>	UCB INC	<u>5MG</u>	<u>A086429 001</u>	
<u>AB</u>		<u>10MG</u>	<u>A085799 001</u>	
<u>AB</u>		<u>20MG</u>	<u>A086428 001</u>	
<u>AB</u>	VINTAGE PHARMS	<u>5MG</u>	<u>A202892 001</u>	Sep 23, 2014
<u>AB</u>		<u>10MG</u>	<u>A202892 002</u>	Sep 23, 2014
<u>AB</u>		<u>20MG</u>	<u>A202892 003</u>	Sep 23, 2014

RITALIN

<u>AB</u> + NOVARTIS	<u>5MG</u>	<u>N010187 003</u>
<u>AB</u> +	<u>10MG</u>	<u>N010187 006</u>
<u>AB</u> +!	<u>20MG</u>	<u>N010187 010</u>

TABLET, CHEWABLE;ORAL

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	ASCENT PHARMS INC	<u>2.5MG</u>	<u>A210354 001</u>	Dec 29, 2017
<u>AB</u>		<u>5MG</u>	<u>A210354 002</u>	Dec 29, 2017
<u>AB</u>		<u>10MG</u>	<u>A210354 003</u>	Dec 29, 2017
<u>AB</u>	BRECKENRIDGE PHARM	<u>2.5MG</u>	<u>A204954 001</u>	Jan 26, 2017
<u>AB</u>		<u>5MG</u>	<u>A204954 002</u>	Jan 26, 2017
<u>AB</u>		<u>10MG</u>	<u>A204954 003</u>	Jan 26, 2017
<u>AB</u>	NOVEL LABS INC	<u>2.5MG</u>	<u>A204115 001</u>	Feb 25, 2015
<u>AB</u>		<u>5MG</u>	<u>A204115 002</u>	Feb 25, 2015
<u>AB</u>	!	<u>10MG</u>	<u>A204115 003</u>	Feb 25, 2015
<u>AB</u>	TEDOR PHARMA INC	<u>2.5MG</u>	<u>A205756 001</u>	Nov 07, 2016
<u>AB</u>		<u>5MG</u>	<u>A205756 002</u>	Nov 07, 2016
<u>AB</u>		<u>10MG</u>	<u>A205756 003</u>	Nov 07, 2016

TABLET, EXTENDED RELEASE;ORAL

CONCERTA

<u>AB</u> + JANSSEN PHARMS	<u>18MG</u>	<u>N021121 001</u>	Aug 01, 2000
<u>AB</u> +	<u>27MG</u>	<u>N021121 004</u>	Apr 01, 2002
<u>AB</u> +	<u>36MG</u>	<u>N021121 002</u>	Aug 01, 2000
<u>AB</u> +!	<u>54MG</u>	<u>N021121 003</u>	Dec 08, 2000

METADATE ER

<u>AB</u> ! UCB INC	<u>20MG</u>	<u>A089601 001</u>	Jun 01, 1988
---------------------	-------------	--------------------	--------------

METHYLIN ER

<u>AB</u> SPECGX LLC	<u>10MG</u>	<u>A075629 001</u>	May 09, 2000
<u>AB</u>	<u>20MG</u>	<u>A075629 002</u>	May 09, 2000

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u> ABHAI LLC	<u>10MG</u>	<u>A207488 001</u>	Jun 09, 2015
<u>AB</u>	<u>20MG</u>	<u>A207488 002</u>	Jun 09, 2015
<u>AB</u> CNTY LINE PHARMS	<u>10MG</u>	<u>A204772 001</u>	Feb 29, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-284 (of 436)

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

**METHYLPHENIDATE HYDROCHLORIDE**

<b>AB</b>		<b>20MG</b>	<b>A204772 002</b>	Feb 29, 2016
<b>AB</b>	COREPHARMA	<b>18MG</b>	<b>A208607 001</b>	Jul 14, 2017
<b>AB</b>		<b>27MG</b>	<b>A208607 002</b>	Jul 14, 2017
<b>AB</b>		<b>36MG</b>	<b>A208607 003</b>	Jul 14, 2017
<b>AB</b>		<b>54MG</b>	<b>A208607 004</b>	Jul 14, 2017
<b>AB</b>	MYLAN PHARMS INC	<b>18MG</b>	<b>A206726 001</b>	Oct 21, 2016
<b>AB</b>		<b>27MG</b>	<b>A206726 002</b>	Oct 21, 2016
<b>AB</b>		<b>36MG</b>	<b>A206726 003</b>	Oct 21, 2016
<b>AB</b>		<b>54MG</b>	<b>A206726 004</b>	Oct 21, 2016
<b>AB</b>	OSMOTICA	<b>18MG</b>	<b>A205327 001</b>	Jul 28, 2017
<b>AB</b>		<b>27MG</b>	<b>A205327 002</b>	Jul 28, 2017
<b>AB</b>		<b>36MG</b>	<b>A205327 003</b>	Jul 28, 2017
<b>AB</b>		<b>54MG</b>	<b>A205327 004</b>	Jul 28, 2017

**RITALIN-SR**

<b>AB</b>	+ NOVARTIS	<b>20MG</b>	<b>N018029 001</b>	Mar 30, 1982
	METHYLPHENIDATE HYDROCHLORIDE			
BX	KREMERS URBAN PHARMS	18MG	A091695 001	Jul 09, 2013
BX		27MG	A091695 002	Jul 09, 2013
BX		36MG	A091695 003	Sep 23, 2013
BX		54MG	A091695 004	Sep 23, 2013
BX	SPECGX LLC	27MG	A202608 001	Dec 28, 2012
BX		36MG	A202608 002	Dec 28, 2012
BX		54MG	A202608 003	Dec 28, 2012
	OSMOTICA	72MG	A205327 005	Jul 28, 2017
	TABLET, EXTENDED RELEASE, CHEWABLE; ORAL			
	QUILLICHEW ER			
	+ PFIZER INC	20MG	N207960 001	Dec 04, 2015
	+	30MG	N207960 002	Dec 04, 2015
	+!	40MG	N207960 003	Dec 04, 2015

METHYLPREDNISOLONE

TABLET; ORAL

**MEDROL**

<b>AB</b>	+ PHARMACIA AND UPJOHN	<b>4MG</b>	<b>N011153 001</b>	
<b>AB</b>	+	<b>8MG</b>	<b>N011153 004</b>	
<b>AB</b>	+	<b>16MG</b>	<b>N011153 003</b>	
<b>AB</b>	+!	<b>32MG</b>	<b>N011153 006</b>	

**METHYLPREDNISOLONE**

<b>AB</b>	DURAMED PHARMS BARR	<b>4MG</b>	<b>A088497 001</b>	Feb 21, 1984
<b>AB</b>	JUBILANT CADISTA	<b>4MG</b>	<b>A040189 001</b>	Oct 31, 1997
<b>AB</b>		<b>8MG</b>	<b>A040189 002</b>	Oct 31, 1997
<b>AB</b>		<b>16MG</b>	<b>A040189 003</b>	Jul 20, 2007
<b>AB</b>		<b>32MG</b>	<b>A040189 004</b>	Jul 20, 2007
<b>AB</b>	SANDOZ	<b>4MG</b>	<b>A040194 001</b>	Oct 31, 1997
<b>AB</b>	VINTAGE PHARMS	<b>4MG</b>	<b>A040183 001</b>	Dec 22, 1998
<b>AB</b>	WATSON LABS	<b>4MG</b>	<b>A040232 001</b>	Oct 16, 1997
	MEDROL			
	+ PHARMACIA AND UPJOHN	2MG	N011153 002	

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

**DEPO-MEDROL**

<b>AB</b>	+! PHARMACIA AND UPJOHN	<b>40MG/ML</b>	<b>N011757 001</b>	
<b>AB</b>	+	<b>80MG/ML</b>	<b>N011757 004</b>	

**METHYLPREDNISOLONE ACETATE**

<b>AB</b>	SANDOZ INC	<b>40MG/ML</b>	<b>A040719 001</b>	Jan 29, 2009
<b>AB</b>		<b>40MG/ML</b>	<b>A040794 001</b>	Mar 05, 2009
<b>AB</b>		<b>80MG/ML</b>	<b>A040719 002</b>	Jan 29, 2009
<b>AB</b>		<b>80MG/ML</b>	<b>A040794 002</b>	Mar 05, 2009
<b>AB</b>	TEVA PHARMS USA	<b>40MG/ML</b>	<b>A040557 001</b>	Feb 23, 2005
<b>AB</b>		<b>40MG/ML</b>	<b>A040620 001</b>	Oct 27, 2006
<b>AB</b>		<b>80MG/ML</b>	<b>A040557 002</b>	Feb 23, 2005
<b>AB</b>		<b>80MG/ML</b>	<b>A040620 002</b>	Oct 27, 2006

DEPO-MEDROL

+! PHARMACIA AND UPJOHN	20MG/ML	N011757 002
-------------------------	---------	-------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-285 (of 436)

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

<u>AP</u>	HOSPIRA	<u>EQ 40MG BASE/VIAL</u>	<u>A040664 001</u>	Dec 20, 2005
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040665 001</u>	Dec 20, 2005
<u>METHYLPREDNISOLONE SODIUM SUCCINATE</u>				
<u>AP</u>	AMNEAL PHARMS CO	<u>EQ 40MG BASE/VIAL</u>	<u>A207549 001</u>	Nov 09, 2016
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A207549 002</u>	Nov 09, 2016
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A207667 001</u>	Dec 15, 2015
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A207667 002</u>	Dec 15, 2015
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A207667 003</u>	Dec 15, 2015
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A207667 004</u>	Dec 15, 2015
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 40MG BASE/VIAL</u>	<u>A040583 001</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040583 002</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040612 001</u>	Aug 12, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 500MG BASE/VIAL</u>	<u>A202691 001</u>	Feb 16, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A202691 002</u>	Feb 16, 2016
<u>AP</u>	SAGENT PHARMS	<u>EQ 40MG BASE/VIAL</u>	<u>A040888 001</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>A040888 002</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A040888 003</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A040888 004</u>	Jul 18, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A040888 005</u>	Jul 18, 2011
<u>SOLU-MEDROL</u>				
<u>AP</u>	+! PHARMACIA AND UPJOHN	<u>EQ 40MG BASE/VIAL</u>	<u>N011856 003</u>	
<u>AP</u>	+!	<u>EQ 125MG BASE/VIAL</u>	<u>N011856 004</u>	
<u>AP</u>	+!	<u>EQ 500MG BASE/VIAL</u>	<u>N011856 005</u>	
<u>AP</u>	+!	<u>EQ 1GM BASE/VIAL</u>	<u>N011856 006</u>	
<u>AP</u>	+!	<u>EQ 2GM BASE/VIAL</u>	<u>N011856 007</u>	Feb 27, 1985

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

<u>AB</u>	IMPAK LABS INC	<u>10MG</u>	<u>A204851 001</u>	Sep 21, 2015
<u>TESTRED</u>				
<u>AB</u>	! VALEANT PHARM INTL	<u>10MG</u>	<u>A083976 001</u>	
	TABLET; ORAL			
	ANDROID 25			
BP	VALEANT PHARM INTL	25MG		A087147 001
	METHYLTESTOSTERONE			
BP	IMPAK LABS	10MG		A080767 002

METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

METIPRANOLOL

<u>AT</u>	SANDOZ INC	<u>0.3%</u>	<u>A075720 001</u>	Aug 06, 2001
<u>OPTIPRANOLOL</u>				
<u>AT</u>	+! BAUSCH AND LOMB	<u>0.3%</u>	<u>N019907 001</u>	Dec 29, 1989

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE

<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 5MG BASE/ML</u>	<u>A204756 001</u>	Dec 20, 2013
<u>METOCLOPRAMIDE HYDROCHLORIDE</u>				
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 5MG BASE/ML</u>	<u>A091392 001</u>	Apr 19, 2013
<u>AP</u>	! HOSPIRA	<u>EQ 5MG BASE/ML</u>	<u>A073118 001</u>	Jan 17, 1991
<u>AP</u>	TEVA PHARMS USA	<u>EQ 5MG BASE/ML</u>	<u>A073135 001</u>	Nov 27, 1991

SOLUTION; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AA</u>	ANI PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>A071402 001</u>	Jun 25, 1993
<u>AA</u>	PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>A072744 001</u>	May 28, 1991
<u>AA</u>	VISTAPHARM	<u>EQ 5MG BASE/5ML</u>	<u>A075051 001</u>	Jan 26, 2001
<u>AA</u>	+! WOCKHARDT BIO AG	<u>EQ 5MG BASE/5ML</u>	<u>A074703 001</u>	Oct 31, 1997

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AB</u>	IMPAX LABS INC	<u>EQ 5MG BASE</u>	<u>A071250 002</u>	Dec 28, 1995
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A071250 001</u>	Feb 03, 1988
<u>AB</u>	IPCA LABS LTD	<u>EQ 5MG BASE</u>	<u>A078807 001</u>	Jun 12, 2008
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078807 002</u>	Jun 12, 2008
<u>AB</u>	PAR PHARM INC	<u>EQ 10MG BASE</u>	<u>A070581 001</u>	Oct 17, 1985
<u>AB</u>	TEVA	<u>EQ 5MG BASE</u>	<u>A072801 001</u>	Jun 15, 1993
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A070184 001</u>	Jul 29, 1985
<u>AB</u>	VINTAGE PHARMS	<u>EQ 5MG BASE</u>	<u>A077878 001</u>	Aug 28, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-286 (of 436)

METOCLOPRAMIDE HYDROCHLORIDE

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077878 002</u>	Aug 28, 2006
	<u>REGLAN</u>			
<u>AB</u>	+ ANI PHARMS	<u>EQ 5MG BASE</u>	<u>N017854 002</u>	May 05, 1987
<u>AB</u>	+!	<u>EQ 10MG BASE</u>	<u>N017854 001</u>	
	TABLET, ORALLY DISINTEGRATING; ORAL			
	<u>METOCLOPRAMIDE HYDROCHLORIDE</u>			
<u>AB</u>	NOVEL LABS INC	<u>EQ 5MG BASE</u>	<u>A202191 001</u>	Aug 15, 2014
	<u>METOZOLV ODT</u>			
<u>AB</u>	+! SALIX PHARMS	<u>EQ 5MG BASE</u>	<u>N022246 001</u>	Sep 04, 2009
	METOCLOPRAMIDE HYDROCHLORIDE			
	NOVEL LABS INC	EQ 10MG BASE		
			<u>A202191 002</u>	Aug 15, 2014

METOLAZONE

TABLET; ORAL

METOLAZONE

<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>A076698 001</u>	Dec 23, 2003
<u>AB</u>		<u>5MG</u>	<u>A076698 002</u>	Oct 19, 2004
<u>AB</u>		<u>10MG</u>	<u>A076698 003</u>	Oct 19, 2004
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>A076732 001</u>	Dec 19, 2003
<u>AB</u>		<u>5MG</u>	<u>A076466 001</u>	Dec 19, 2003
<u>AB</u>		<u>10MG</u>	<u>A076466 002</u>	Dec 19, 2003
	<u>ZAROXOLYN</u>			
<u>AB</u>	+ UCB INC	<u>2.5MG</u>	<u>N017386 001</u>	
<u>AB</u>	+!	<u>5MG</u>	<u>N017386 002</u>	
<u>AB</u>	+!	<u>10MG</u>	<u>N017386 003</u>	

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>EQ 25MG TARTRATE</u>	<u>A204161 001</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A204161 002</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A204161 003</u>	Nov 25, 2016
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A204161 004</u>	Nov 25, 2016
<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 25MG TARTRATE</u>	<u>A077118 001</u>	Aug 03, 2009
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A076862 001</u>	Aug 03, 2009
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A077298 001</u>	Apr 15, 2010
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A077298 002</u>	Apr 15, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG TARTRATE</u>	<u>A090617 001</u>	Aug 01, 2012
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A090617 002</u>	Aug 01, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 25MG TARTRATE</u>	<u>A202033 001</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A202033 002</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A202033 003</u>	Dec 15, 2011
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A202033 004</u>	Dec 15, 2011
<u>AB</u>	REDDYS	<u>EQ 100MG TARTRATE</u>	<u>A078889 001</u>	Aug 15, 2012
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A078889 002</u>	Aug 15, 2012
<u>AB</u>	WOCKHARDT	<u>EQ 25MG TARTRATE</u>	<u>A090615 001</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 50MG TARTRATE</u>	<u>A090615 002</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 100MG TARTRATE</u>	<u>A090615 003</u>	Jul 22, 2010
<u>AB</u>		<u>EQ 200MG TARTRATE</u>	<u>A090615 004</u>	Jul 22, 2010
	<u>TOPROL-XL</u>			
<u>AB</u>	+ ARALEZ PHARMS	<u>EQ 25MG TARTRATE</u>	<u>N019962 004</u>	Feb 05, 2001
<u>AB</u>	+!	<u>EQ 50MG TARTRATE</u>	<u>N019962 001</u>	Jan 10, 1992
<u>AB</u>	+	<u>EQ 100MG TARTRATE</u>	<u>N019962 002</u>	Jan 10, 1992
<u>AB</u>	+!	<u>EQ 200MG TARTRATE</u>	<u>N019962 003</u>	Jan 10, 1992

METOPROLOL TARTRATE

INJECTABLE; INJECTION

LOPRESSOR

<u>AP</u>	+! NOVARTIS	<u>1MG/ML</u>	<u>N018704 001</u>	Mar 30, 1984
	<u>METOPROLOL TARTRATE</u>			
<u>AP</u>	BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>A078950 001</u>	Apr 29, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>1MG/ML</u>	<u>A091045 001</u>	Oct 25, 2010
<u>AP</u>	GLAND PHARMA LTD	<u>1MG/ML</u>	<u>A204205 001</u>	Aug 27, 2014
<u>AP</u>	HIKMA FARMACEUTICA	<u>1MG/ML</u>	<u>A077761 001</u>	May 30, 2007
<u>AP</u>	HOSPIRA	<u>1MG/ML</u>	<u>A074133 001</u>	Dec 21, 1993
<u>AP</u>		<u>1MG/ML</u>	<u>A075160 001</u>	Jul 06, 1998
<u>AP</u>		<u>1MG/ML</u>	<u>A078085 001</u>	Apr 29, 2008
<u>AP</u>	LUITPOLD	<u>1MG/ML</u>	<u>A090386 001</u>	Sep 30, 2009
<u>AP</u>		<u>1MG/ML</u>	<u>A091307 001</u>	Dec 29, 2010
<u>AP</u>	SAGENT STRIDES	<u>1MG/ML</u>	<u>A090317 001</u>	Apr 19, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-287 (of 436)

METOPROLOL TARTRATE

INJECTABLE; INJECTION

**METOPROLOL TARTRATE**

<u>AP</u>	SANDOZ INC	<u>1MG/ML</u>	<u>A077360 001</u>	Oct 02, 2007
<u>AP</u>	WEST-WARD PHARMS INT	<u>1MG/ML</u>	<u>A076495 001</u>	Jul 07, 2003
TABLET;ORAL				
<b>LOPRESSOR</b>				
<u>AB</u>	+ US PHARMS HOLDINGS I	<u>50MG</u>	<u>N017963 001</u>	
<u>AB</u>	+ I	<u>100MG</u>	<u>N017963 002</u>	
<b>METOPROLOL TARTRATE</b>				
<u>AB</u>	ALEMBIC PHARMS LTD	<u>25MG</u>	<u>A202871 001</u>	May 28, 2013
<u>AB</u>		<u>50MG</u>	<u>A202871 002</u>	May 28, 2013
<u>AB</u>		<u>100MG</u>	<u>A202871 003</u>	May 28, 2013
<u>AB</u>	AUROBINDO PHARMA	<u>25MG</u>	<u>A077739 001</u>	Sep 11, 2007
<u>AB</u>		<u>50MG</u>	<u>A077739 002</u>	Sep 11, 2007
<u>AB</u>		<u>100MG</u>	<u>A077739 003</u>	Sep 11, 2007
<u>AB</u>	IPCA LABS LTD	<u>25MG</u>	<u>A078459 001</u>	Jun 17, 2008
<u>AB</u>		<u>50MG</u>	<u>A078459 002</u>	Jun 17, 2008
<u>AB</u>		<u>100MG</u>	<u>A078459 003</u>	Jun 17, 2008
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A076704 001</u>	Jan 16, 2004
<u>AB</u>		<u>50MG</u>	<u>A076704 002</u>	Jan 16, 2004
<u>AB</u>	!	<u>100MG</u>	<u>A076704 003</u>	Jan 16, 2004
<u>AB</u>	RUBICON RES PVT LTD	<u>25MG</u>	<u>A200981 001</u>	Oct 28, 2014
<u>AB</u>		<u>50MG</u>	<u>A200981 002</u>	Oct 28, 2014
<u>AB</u>		<u>100MG</u>	<u>A200981 003</u>	Oct 28, 2014
<u>AB</u>	SUN PHARM INDNS INC	<u>25MG</u>	<u>A076670 001</u>	Jan 15, 2004
<u>AB</u>		<u>50MG</u>	<u>A074644 001</u>	Dec 10, 1996
<u>AB</u>		<u>100MG</u>	<u>A074644 002</u>	Dec 10, 1996
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A073654 002</u>	Jul 15, 2009
<u>AB</u>		<u>50MG</u>	<u>A073654 003</u>	Dec 21, 1993
<u>AB</u>		<u>100MG</u>	<u>A073654 001</u>	Dec 21, 1993
<u>AB</u>	TEVA	<u>50MG</u>	<u>A074141 001</u>	Jan 31, 1995
<u>AB</u>		<u>100MG</u>	<u>A074141 002</u>	Jan 31, 1995
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>A074217 001</u>	May 27, 1994
<u>AB</u>		<u>100MG</u>	<u>A074217 002</u>	May 27, 1994
	MYLAN	37.5MG	A076704 004	Mar 18, 2015
		75MG	A076704 005	Mar 18, 2015

METRONIDAZOLE

CAPSULE; ORAL

**FLAGYL**

<u>AB</u>	+! GD SEARLE LLC	<u>375MG</u>	<u>N020334 001</u>	May 03, 1995
<b>METRONIDAZOLE</b>				
<u>AB</u>	ALEMBIC PHARMS LTD	<u>375MG</u>	<u>A079065 001</u>	Jun 23, 2009
<u>AB</u>	PAR PHARM	<u>375MG</u>	<u>A076522 001</u>	Jan 29, 2004

CREAM;TOPICAL

**METROCREAM**

<u>AB</u>	+! GALDERMA LABS LP	<u>0.75%</u>	<u>N020531 001</u>	Sep 20, 1995
<b>METRONIDAZOLE</b>				

<u>AB</u>	FOUGERA PHARMS	<u>0.75%</u>	<u>A076408 001</u>	May 28, 2004
<u>AB</u>	G AND W LABS	<u>0.75%</u>	<u>A077549 001</u>	Dec 19, 2007

NORITATE

<u>AB</u>	+! VALEANT PHARMS NORTH	1%	N020743 001	Sep 26, 1997
<b>GEL;TOPICAL</b>				

**METROGEL**

<u>AB</u>	+! GALDERMA LABS LP	<u>0.75%</u>	<u>N019737 001</u>	Nov 22, 1988
<u>AB</u>	+!	<u>1%</u>	<u>N021789 001</u>	Jun 30, 2005

**METRONIDAZOLE**

<u>AB</u>	FOUGERA PHARMS	<u>0.75%</u>	<u>A077018 001</u>	Jun 06, 2006
<u>AB</u>	G AND W LABS INC	<u>0.75%</u>	<u>A078178 001</u>	Jan 19, 2011
<u>AB</u>	TARO	<u>0.75%</u>	<u>A077819 001</u>	Jul 18, 2006
<u>AB</u>	TARO PHARM	<u>1%</u>	<u>A204651 001</u>	Mar 14, 2017
<u>AB</u>	TOLMAR	<u>0.75%</u>	<u>A077547 001</u>	Jul 13, 2006
<u>AB</u>		<u>1%</u>	<u>A090903 001</u>	Jul 22, 2011

GEL;VAGINAL

**METROGEL-VAGINAL**

<u>AB</u>	+! MEDICIS	<u>0.75%</u>	<u>N020208 001</u>	Aug 17, 1992
<b>METRONIDAZOLE</b>				
<u>AB</u>	TOLMAR	<u>0.75%</u>	<u>A077264 001</u>	Oct 31, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-288 (of 436)

METRONIDAZOLE

GEL;VAGINAL			
VANDAZOLE			
BX TEVA PHARMS	0.75%		N021806 001 May 20, 2005
NUVESSA			
+! CHEMO RESEARCH SL	1.3%		N205223 001 Mar 24, 2014
INJECTABLE;INJECTION			
<b><u>FLAGYL I.V. RTU IN PLASTIC CONTAINER</u></b>			
<b>AP +!</b>	<b>BAXTER HLTHCARE</b>	<b>500MG/100ML</b>	<b>N018657 001</b>
<b><u>METRO I.V. IN PLASTIC CONTAINER</u></b>			
<b>AP +!</b>	<b>B BRAUN</b>	<b>500MG/100ML</b>	<b>N018900 001</b> Sep 29, 1983
<b><u>METRONIDAZOLE IN PLASTIC CONTAINER</u></b>			
<b>AP</b>	<b>BAXTER HLTHCARE</b>	<b>500MG/100ML</b>	<b>A078084 001</b> Mar 31, 2008
CORP			
<b>AP +!</b>	<b>HOSPIRA</b>	<b>500MG/100ML</b>	<b>N018890 002</b> Nov 18, 1983
<b>AP</b>	<b>MYLAN LABS LTD</b>	<b>500MG/100ML</b>	<b>A205531 001</b> May 09, 2017
LOTION;TOPICAL			
<b><u>METROLOTION</u></b>			
<b>AB +!</b>	<b>GALDERMA LABS LP</b>	<b>0.75%</b>	<b>N020901 001</b> Nov 24, 1998
<b><u>METRONIDAZOLE</u></b>			
<b>AB</b>	<b>FOUGERA PHARMS</b>	<b>0.75%</b>	<b>A077197 001</b> May 24, 2006
TABLET;ORAL			
<b><u>FLAGYL</u></b>			
<b>AB +</b>	<b>GD SEARLE LLC</b>	<b>250MG</b>	<b>N012623 001</b>
<b>AB +!</b>		<b>500MG</b>	<b>N012623 003</b>
<b><u>METRONIDAZOLE</u></b>			
<b>AB</b>	<b>ALEMBIC PHARMS LTD</b>	<b>250MG</b>	<b>A079067 001</b> Mar 13, 2009
<b>AB</b>		<b>500MG</b>	<b>A079067 002</b> Mar 13, 2009
<b>AB</b>	<b>APPCO PHARMA LLC</b>	<b>250MG</b>	<b>A205245 001</b> Sep 23, 2015
<b>AB</b>		<b>500MG</b>	<b>A205245 002</b> Sep 23, 2015
<b>AB</b>	<b>AUROBINDO PHARMA LTD</b>	<b>250MG</b>	<b>A203974 001</b> May 29, 2015
<b>AB</b>		<b>500MG</b>	<b>A203974 002</b> May 29, 2015
<b>AB</b>	<b>CADILA PHARMS LTD</b>	<b>250MG</b>	<b>A209794 001</b> Dec 12, 2017
<b>AB</b>		<b>500MG</b>	<b>A209794 002</b> Dec 12, 2017
<b>AB</b>	<b>FLAMINGO PHARMS</b>	<b>250MG</b>	<b>A207309 001</b> May 16, 2016
<b>AB</b>		<b>500MG</b>	<b>A207309 002</b> May 16, 2016
<b>AB</b>	<b>INNOGENIX</b>	<b>250MG</b>	<b>A070772 001</b> Jul 16, 1986
<b>AB</b>		<b>500MG</b>	<b>A070772 002</b> Jul 16, 1986
<b>AB</b>	<b>LUPIN LTD</b>	<b>250MG</b>	<b>A209096 001</b> Sep 12, 2017
<b>AB</b>		<b>500MG</b>	<b>A209096 002</b> Sep 12, 2017
<b>AB</b>	<b>ORIT LABS LLC</b>	<b>250MG</b>	<b>A208681 001</b> Jun 20, 2017
<b>AB</b>		<b>500MG</b>	<b>A208681 002</b> Jun 20, 2017
<b>AB</b>	<b>PLIVA</b>	<b>500MG</b>	<b>A070033 001</b> Dec 06, 1984
<b>AB</b>	<b>STRIDES PHARMA</b>	<b>250MG</b>	<b>A208162 001</b> May 25, 2016
<b>AB</b>		<b>500MG</b>	<b>A208162 002</b> May 25, 2016
<b>AB</b>	<b>TEVA PHARMS USA</b>	<b>250MG</b>	<b>A070027 001</b> Nov 06, 1984
<b>AB</b>	<b>UNICHEM LABS LTD</b>	<b>250MG</b>	<b>A203458 001</b> Jan 22, 2014
<b>AB</b>		<b>500MG</b>	<b>A203458 002</b> Jan 22, 2014
<b>AB</b>	<b>VIVIMED GLOBAL</b>	<b>250MG</b>	<b>A070040 001</b> Jan 29, 1985
<b>AB</b>		<b>500MG</b>	<b>A070039 001</b> Jan 29, 1985
<b>AB</b>	<b>WATSON LABS</b>	<b>250MG</b>	<b>A070035 001</b> Dec 20, 1984
<b>AB</b>	<b>WATSON LABS INC</b>	<b>500MG</b>	<b>A070044 001</b> Feb 08, 1985
<b>AB</b>	<b>ZYDUS PHARMS USA INC</b>	<b>250MG</b>	<b>A206560 001</b> Nov 16, 2016
<b>AB</b>		<b>500MG</b>	<b>A206560 002</b> Nov 16, 2016
TABLET, EXTENDED RELEASE;ORAL			
<b><u>FLAGYL ER</u></b>			
<b>AB +!</b>	<b>GD SEARLE LLC</b>	<b>750MG</b>	<b>N020868 001</b> Nov 26, 1997
<b><u>METRONIDAZOLE</u></b>			
<b>AB</b>	<b>ALEMBIC PHARMS LTD</b>	<b>750MG</b>	<b>A090222 001</b> May 05, 2010
<b><u>METYRAPONE</u></b>			
CAPSULE;ORAL			
METOPIRONE			
+! HRA PHARMA	250MG		N012911 002 Aug 09, 1996
<b><u>METYROSINE</u></b>			
CAPSULE;ORAL			
DEMSEER			
+! ATON PHARMA VPNA	250MG		N017871 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-289 (of 436)

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL		
MEXILETINE HYDROCHLORIDE		
TEVA	150MG	A074377 001 May 16, 1995
	200MG	A074377 002 May 16, 1995
!	250MG	A074377 003 May 16, 1995

MICAFUNGIN SODIUM

INJECTABLE; IV (INFUSION)		
MYCAMEINE		
+! ASTELLAS	EQ 50MG BASE/VIAL	N021506 002 Mar 16, 2005
+!	EQ 100MG BASE/VIAL	N021506 003 Jun 27, 2006

MICONAZOLE

TABLET; Buccal		
ORAVIG		
+! MIDATECH PHARMA US	50MG	N022404 001 Apr 16, 2010

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL		
<b>MICONAZOLE NITRATE</b>		
<b>AB</b> ACTAVIS PHARMA	<b>200MG</b>	<b>A073508 001</b> Nov 19, 1993
<b>MONISTAT 3</b>		
<b>AB</b> +! MEDTECH PRODUCTS	<b>200MG</b>	<b>N018888 001</b> Aug 15, 1984

MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL		
VUSION		
+! MYLAN PHARMS INC	0.25%, 81.35%; 15%	N021026 001 Feb 16, 2006

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION		
<b>MIDAZOLAM HYDROCHLORIDE</b>		
<b>AP</b> AKORN INC	<b>EQ 1MG BASE/ML</b>	<b>A075494 001</b> Jun 30, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075481 001</b> Jun 30, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075494 002</b> Jun 30, 2000
<b>AP</b> FRESENIUS KABI USA	<b>EQ 1MG BASE/ML</b>	<b>A075154 002</b> Jun 20, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075154 001</b> Jun 20, 2000
<b>AP</b> GLAND PHARMA LTD	<b>EQ 1MG BASE/ML</b>	<b>A090696 001</b> Feb 29, 2012
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A090850 001</b> Jan 25, 2012
<b>AP</b> ! HOSPIRA	<b>EQ 1MG BASE/ML</b>	<b>A075293 001</b> Jun 20, 2000
<b>AP</b>	<b>EQ 1MG BASE/ML</b>	<b>A075856 001</b> Jun 13, 2002
<b>AP</b> !	<b>EQ 5MG BASE/ML</b>	<b>A075293 002</b> Jun 20, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075856 002</b> Jun 13, 2002
<b>AP</b> WEST-WARD PHARMS INT	<b>EQ 1MG BASE/ML</b>	<b>A075243 001</b> Jun 20, 2000
<b>AP</b>	<b>EQ 1MG BASE/ML</b>	<b>A075247 002</b> Jun 23, 2000
<b>AP</b>	<b>EQ 1MG BASE/ML</b>	<b>A075324 001</b> Jun 20, 2000
<b>AP</b>	<b>EQ 1MG BASE/ML</b>	<b>A075421 002</b> Jun 20, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075243 002</b> Jun 20, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075247 001</b> Jun 23, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075324 002</b> Jun 20, 2000
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A075421 001</b> Jun 20, 2000

MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE

<b>AP</b> FRESENIUS KABI USA	<b>EQ 1MG BASE/ML</b>	<b>A203460 001</b> Aug 22, 2014
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A203460 002</b> Aug 22, 2014
<b>AP</b> ! HOSPIRA	<b>EQ 1MG BASE/ML</b>	<b>A075857 001</b> Jul 22, 2002
<b>AP</b> !	<b>EQ 5MG BASE/ML</b>	<b>A075857 002</b> Jul 22, 2002
<b>AP</b> SAGENT AGILA LLC	<b>EQ 1MG BASE/ML</b>	<b>A090315 001</b> Nov 29, 2010
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A090315 002</b> Nov 29, 2010

MIDOZALAM HYDROCHLORIDE

<b>AP</b> SAGENT STRIDES	<b>EQ 1MG BASE/ML</b>	<b>A090316 001</b> May 04, 2011
<b>AP</b>	<b>EQ 5MG BASE/ML</b>	<b>A090316 002</b> May 04, 2011
MIDAZOLAM HYDROCHLORIDE		
FRESENIUS KABI USA	EQ 5MG BASE/ML	
SYRUP; ORAL		
<b>MIDAZOLAM HYDROCHLORIDE</b>		
<b>AA</b> HI TECH PHARMA	<b>EQ 2MG BASE/ML</b>	<b>A075958 001</b> Sep 04, 2003
<b>AA</b> PADDICK LLC	<b>EQ 2MG BASE/ML</b>	<b>A076379 001</b> May 02, 2005
<b>AA</b> ! WEST-WARD PHARMS INT	<b>EQ 2MG BASE/ML</b>	<b>A075873 001</b> Apr 30, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-290 (of 436)

MIDODRINE HYDROCHLORIDE

TABLET;ORAL

MIDODRINE HYDROCHLORIDE

<u>AB</u>	IMPAK PHARMS	<u>2.5MG</u>	<u>A076449 001</u>	May 27, 2004
<u>AB</u>		<u>5MG</u>	<u>A076449 002</u>	May 27, 2004
<u>AB</u>		<u>10MG</u>	<u>A076449 003</u>	Dec 16, 2005
<u>AB</u>	MYLAN PHARMS INC	<u>2.5MG</u>	<u>A076577 001</u>	Sep 10, 2003
<u>AB</u>		<u>5MG</u>	<u>A076577 002</u>	Sep 10, 2003
<u>AB</u>		<u>10MG</u>	<u>A076577 003</u>	Sep 10, 2003
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>A076514 001</u>	Sep 11, 2003
<u>AB</u>		<u>5MG</u>	<u>A076514 002</u>	Sep 11, 2003
<u>AB</u>		<u>10MG</u>	<u>A076514 003</u>	Jul 02, 2004
<u>ORVATEN</u>				
<u>AB</u>	UPSHER-SMITH LABS	<u>2.5MG</u>	<u>A076725 001</u>	Nov 03, 2004
<u>AB</u>		<u>5MG</u>	<u>A076725 002</u>	Nov 03, 2004
<u>AB</u>		<u>10MG</u>	<u>A076725 003</u>	Nov 03, 2004
MIDODRINE HYDROCHLORIDE				
BX	APOTEX INC	2.5MG	A077746 001	Sep 12, 2006
BX		5MG	A077746 002	Sep 12, 2006
BX		10MG	A077746 003	Sep 12, 2006

MIDOSTAURIN

CAPSULE;ORAL

RYDAPT

+!	NOVARTIS PHARMS CORP	25MG	N207997 001	Apr 28, 2017
----	----------------------	------	-------------	--------------

MIFEPRISTONE

TABLET;ORAL

KORLYM

+!	CORCEPT THERAP	300MG	N202107 001	Feb 17, 2012
MIFEPRISTONE	DANCO LABS LLC	200MG	N020687 001	Sep 28, 2000

MIGLITOL

TABLET;ORAL

GLYSET

<u>AB</u>	+!	PHARMACIA AND UPJOHN	<u>25MG</u>	<u>N020682 001</u>	Dec 18, 1996
<u>AB</u>	+		<u>50MG</u>	<u>N020682 002</u>	Dec 18, 1996
<u>AB</u>	+		<u>100MG</u>	<u>N020682 003</u>	Dec 18, 1996
<u>MIGLITOL</u>					
<u>AB</u>	ORIENT PHARMA CO LTD	<u>25MG</u>	<u>A203965 001</u>	Feb 24, 2015	
<u>AB</u>		<u>50MG</u>	<u>A203965 002</u>	Feb 24, 2015	
<u>AB</u>		<u>100MG</u>	<u>A203965 003</u>	Feb 24, 2015	

MIGLUSTAT

CAPSULE;ORAL

ZAVESCA

+!	ACTELION PHARMS LTD	100MG	N021348 001	Jul 31, 2003
----	---------------------	-------	-------------	--------------

MILNACIPRAN HYDROCHLORIDE

TABLET;ORAL

MILNACIPRAN HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>12.5MG</u>	<u>A205081 001</u>	Apr 22, 2016
<u>AB</u>		<u>25MG</u>	<u>A205081 002</u>	Apr 22, 2016
<u>AB</u>		<u>50MG</u>	<u>A205081 003</u>	Apr 22, 2016
<u>AB</u>		<u>100MG</u>	<u>A205081 004</u>	Apr 22, 2016
<u>SAVELLA</u>				
<u>AB</u>	+ ALLERGAN SALES LLC	<u>12.5MG</u>	<u>N022256 001</u>	Jan 14, 2009
<u>AB</u>	+	<u>25MG</u>	<u>N022256 002</u>	Jan 14, 2009
<u>AB</u>	+!	<u>50MG</u>	<u>N022256 003</u>	Jan 14, 2009
<u>AB</u>	+	<u>100MG</u>	<u>N022256 004</u>	Jan 14, 2009

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 1MG BASE/ML</u>	<u>A075936 001</u>	May 28, 2002
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 1MG BASE/ML</u>	<u>A077190 001</u>	Oct 31, 2006
<u>AP</u>	! HIKMA FARMACEUTICA	<u>EQ 1MG BASE/ML</u>	<u>A077966 001</u>	Dec 03, 2010
<u>AP</u>	HOSPIRA INC	<u>EQ 1MG BASE/ML</u>	<u>A203280 001</u>	Sep 03, 2014
<u>AP</u>	INTL MEDICATED	<u>EQ 1MG BASE/ML</u>	<u>A076013 001</u>	Aug 02, 2002
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 1MG BASE/ML</u>	<u>A075530 001</u>	May 28, 2002
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>A075660 001</u>	May 28, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-291 (of 436)

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE IN DEXTROSE 5%

<u>AP</u>	RENAISSANCE SSA LLC	<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A077151 002</u>	Jul 20, 2005
<u>AP</u>		<u>MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
<u>AP</u>	! BAXTER HLTHCARE	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A075834 001</u>	May 28, 2002
<u>AP</u>	!	<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A075834 002</u>	May 28, 2002
<u>AP</u>	HOSPIRA	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A075885 001</u>	May 28, 2002
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A075885 002</u>	May 28, 2002
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A078113 001</u>	May 21, 2008
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A078113 002</u>	May 21, 2008
<u>AP</u>		<u>MILRINONE LACTATE IN PLASTIC CONTAINER</u>		
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)</u>	<u>A090038 001</u>	Jan 21, 2010
<u>AP</u>		<u>EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)</u>	<u>A090038 002</u>	Jan 21, 2010

MILTEFOSINE

CAPSULE; ORAL

IMPAVIDO

+! KNIGHT THERAPS

50MG

N204684 001 Mar 19, 2014

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DYNACIN

<u>AB</u>	CNTY LINE PHARMS	<u>EQ 50MG BASE</u>	<u>A063067 003</u>	Aug 14, 1990
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A063067 002</u>	Sep 15, 1999
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A063067 001</u>	Jul 31, 1990
<u>AB</u>		<u>MINOCIN</u>		
<u>AB</u>	+	PRECISION DERMAT	<u>EQ 50MG BASE</u>	<u>N050649 001</u> May 31, 1990
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N050649 002</u> May 31, 1990

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 50MG BASE</u>	<u>A065470 001</u>	Mar 11, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065470 002</u>	Mar 11, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065470 003</u>	Mar 11, 2008
<u>AB</u>	IMPAX LABS	<u>EQ 50MG BASE</u>	<u>A065005 001</u>	Mar 23, 1999
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065005 003</u>	Apr 18, 2001
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065005 002</u>	Mar 23, 1999
<u>AB</u>	SUN PHARM IND'S INC	<u>EQ 50MG BASE</u>	<u>A090867 001</u>	May 13, 2013
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090867 002</u>	May 13, 2013
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090867 003</u>	May 13, 2013
<u>AB</u>	TORRENT PHARMA INC	<u>EQ 50MG BASE</u>	<u>A065062 001</u>	Nov 30, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065062 002</u>	Nov 30, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065062 003</u>	Nov 30, 2000
<u>AB</u>	WATSON LABS	<u>EQ 75MG BASE</u>	<u>A063065 002</u>	Jun 10, 1999
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A063065 001</u>	Dec 30, 1991
<u>AB</u>	WATSON LABS TEVA	<u>EQ 50MG BASE</u>	<u>A063181 001</u>	Dec 30, 1991
<u>AB</u>	ZYDUS WORLDWIDE	<u>EQ 50MG BASE</u>	<u>A063011 001</u>	Mar 02, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A063009 002</u>	Aug 12, 2003
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A063009 001</u>	Mar 02, 1992

CAPSULE, EXTENDED RELEASE; ORAL

XIMINO

SUN PHARM IND'S LTD	<u>EQ 45MG BASE</u>	
	<u>EQ 90MG BASE</u>	
	<u>EQ 135MG BASE</u>	

N201922 001 Jul 11, 2012  
N201922 003 Jul 11, 2012  
N201922 005 Jul 11, 2012

INJECTABLE; INJECTION

MINOCIN

+! REMPEX PHARMS INC

EQ 100MG BASE/VIAL

N050444 001

POWDER, EXTENDED RELEASE; DENTAL

ARESTIN

+! ORAPHARMA

EQ 1MG BASE

N050781 001 Feb 16, 2001

TABLET; ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 50MG BASE</u>	<u>A065436 001</u>	Dec 26, 2007
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065436 002</u>	Dec 26, 2007
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065436 003</u>	Dec 26, 2007
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A065131 001</u>	Apr 16, 2003
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065131 002</u>	Apr 16, 2003
<u>AB</u>	!	<u>EQ 100MG BASE</u>	<u>A065131 003</u>	Apr 16, 2003
<u>AB</u>	SUN PHARM INDUSTRIES	<u>EQ 50MG BASE</u>	<u>A090217 001</u>	Jan 29, 2016
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090217 002</u>	Jan 29, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090217 003</u>	Jan 29, 2016
<u>AB</u>	TORRENT PHARMA INC	<u>EQ 50MG BASE</u>	<u>A065156 001</u>	Jan 06, 2004
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A065156 002</u>	Jan 06, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-292 (of 436)

MINOCYCLINE HYDROCHLORIDE

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A065156 003</u>	Jan 06, 2004	
	TABLET, EXTENDED RELEASE;ORAL				
<u>MINOCYCLINE HYDROCHLORIDE</u>					
<u>AB</u>	ALKEM LABS LTD	<u>EQ 45MG BASE</u>	<u>A204453 001</u>	Sep 28, 2016	
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204453 002</u>	Sep 28, 2016	
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A204453 003</u>	Sep 28, 2016	
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A204453 004</u>	Sep 28, 2016	
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A204453 005</u>	Sep 28, 2016	
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 45MG BASE</u>	<u>A202261 001</u>	Nov 19, 2012	
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A202261 006</u>	Jun 13, 2016	
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A202261 003</u>	Nov 19, 2012	
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A202261 007</u>	Jun 13, 2016	
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A202261 005</u>	Nov 19, 2012	
<u>AB</u>	LUPIN LTD	<u>EQ 45MG BASE</u>	<u>A091424 001</u>	Nov 30, 2011	
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A091424 003</u>	Nov 30, 2011	
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091424 004</u>	Nov 30, 2011	
<u>AB</u>	SANDOZ	<u>EQ 45MG BASE</u>	<u>A090422 001</u>	Aug 13, 2009	
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A090422 002</u>	Aug 13, 2009	
<u>AB</u>	!	<u>EQ 135MG BASE</u>	<u>A090422 003</u>	Aug 13, 2009	
<u>AB</u>	SIDMAK LABS INDIA	<u>EQ 45MG BASE</u>	<u>A204394 001</u>	Dec 30, 2015	
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A204394 004</u>	Dec 30, 2015	
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A204394 005</u>	Dec 30, 2015	
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A204394 007</u>	Dec 30, 2015	
<u>AB</u>	SUN PHARM INDS LTD	<u>EQ 45MG BASE</u>	<u>A091118 001</u>	Sep 25, 2014	
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A091118 004</u>	Sep 25, 2014	
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A091118 005</u>	Sep 25, 2014	
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A091118 006</u>	Sep 25, 2014	
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A091118 008</u>	Sep 25, 2014	
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 45MG BASE</u>	<u>A203553 001</u>	Nov 16, 2017	
<u>AB</u>		<u>EQ 80MG BASE</u>	<u>A203553 004</u>	Nov 16, 2017	
<u>AB</u>		<u>EQ 90MG BASE</u>	<u>A203553 005</u>	Nov 16, 2017	
<u>AB</u>		<u>EQ 105MG BASE</u>	<u>A203553 006</u>	Nov 16, 2017	
<u>AB</u>		<u>EQ 135MG BASE</u>	<u>A203553 008</u>	Nov 16, 2017	
<u>SOLODYN</u>					
<u>AB</u>	+	MEDICIS	<u>EQ 80MG BASE</u>	<u>N050808 007</u>	Aug 27, 2010
<u>AB</u>	+		<u>EQ 105MG BASE</u>	<u>N050808 006</u>	Aug 27, 2010
<u>MINOLIRA</u>					
	DR REDDYS LABS LTD	EQ 105MG BASE	N209269 001	May 08, 2017	
		EQ 135MG BASE	N209269 002	May 08, 2017	
<u>SOLODYN</u>					
	+	MEDICIS	EQ 55MG BASE	N050808 008	Aug 27, 2010
	+		EQ 65MG BASE	N050808 004	Jul 23, 2009
	+	!	EQ 115MG BASE	N050808 005	Jul 23, 2009
<u>MINOXIDIL</u>					
TABLET;ORAL					
<u>MINOXIDIL</u>					
<u>AB</u>	MUTUAL PHARM	<u>2.5MG</u>	<u>A072708 001</u>	Dec 14, 1995	
<u>AB</u>	PAR PHARM	<u>2.5MG</u>	<u>A071826 001</u>	Nov 14, 1988	
<u>AB</u>		<u>10MG</u>	<u>A071839 001</u>	Nov 14, 1988	
<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>	<u>A072709 001</u>	Dec 14, 1995	
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>A071344 001</u>	Mar 03, 1987	
<u>AB</u>	!	<u>10MG</u>	<u>A071345 001</u>	Mar 03, 1987	
<u>MIPOMERSEN SODIUM</u>					
SOLUTION;SUBCUTANEOUS					
KYNAMRO					
+!	KASTLE THERAPS LLC	200MG/ML (200MG/ML)	N203568 001	Jan 29, 2013	
<u>MIRABEGRON</u>					
TABLET, EXTENDED RELEASE;ORAL					
MYRBETRIQ					
+!	APGDI	25MG	N202611 001	Jun 28, 2012	
+!		50MG	N202611 002	Jun 28, 2012	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-293 (of 436)

MIRTAZAPINE

TABLET;ORAL

MIRTAZAPINE

<u>AB</u>	APOTEX INC	<u>15MG</u>	<u>A077666 001</u>	Aug 22, 2007
<u>AB</u>		<u>30MG</u>	<u>A077666 002</u>	Aug 22, 2007
<u>AB</u>		<u>45MG</u>	<u>A077666 003</u>	Aug 22, 2007
<u>AB</u>	AUROBINDO	<u>7.5MG</u>	<u>A076921 001</u>	Oct 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076921 002</u>	Oct 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076921 003</u>	Oct 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076921 004</u>	Oct 22, 2004
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A076122 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076122 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076122 003</u>	Jun 19, 2003
<u>AB</u>	SUN PHARM IND'S INC	<u>7.5MG</u>	<u>A076541 004</u>	Apr 22, 2004
<u>AB</u>		<u>15MG</u>	<u>A076541 001</u>	Apr 22, 2004
<u>AB</u>		<u>30MG</u>	<u>A076541 002</u>	Apr 22, 2004
<u>AB</u>		<u>45MG</u>	<u>A076541 003</u>	Apr 22, 2004
<u>AB</u>	TEVA	<u>15MG</u>	<u>A076119 001</u>	Jan 24, 2003
<u>AB</u>		<u>30MG</u>	<u>A076119 002</u>	Jan 24, 2003
<u>AB</u>		<u>45MG</u>	<u>A076119 003</u>	Jun 19, 2003
<u>AB</u>	UPSHER-SMITH LABS	<u>15MG</u>	<u>A076219 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076219 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076219 003</u>	Jun 19, 2003
<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>A076312 001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>A076312 002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>A076312 003</u>	Jun 19, 2003

REMERON

<u>AB</u>	+!	ORGANON USA INC	<u>15MG</u>	<u>N020415 001</u>	Jun 14, 1996
<u>AB</u>	+		<u>30MG</u>	<u>N020415 002</u>	Jun 14, 1996
<u>AB</u>	+		<u>45MG</u>	<u>N020415 003</u>	Mar 17, 1997

TABLET, ORALLY DISINTEGRATING;ORAL

MIRTAZAPINE

<u>AB</u>	ACTAVIS LABS FL INC	<u>15MG</u>	<u>A076307 001</u>	Dec 17, 2003
<u>AB</u>		<u>30MG</u>	<u>A076307 002</u>	Dec 17, 2003
<u>AB</u>		<u>45MG</u>	<u>A076307 003</u>	Feb 28, 2006
<u>AB</u>	AUROBINDO PHARMA LTD	<u>15MG</u>	<u>A077376 002</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>A077376 003</u>	Dec 08, 2005
<u>AB</u>		<u>45MG</u>	<u>A077376 004</u>	Feb 28, 2006
<u>AB</u>	IMPAX LABS INC	<u>15MG</u>	<u>A076901 001</u>	Jun 28, 2005
<u>AB</u>		<u>30MG</u>	<u>A076901 002</u>	Jun 28, 2005
<u>AB</u>		<u>45MG</u>	<u>A076901 003</u>	Jun 28, 2005
<u>AB</u>	ZYDUS PHARMS USA INC	<u>15MG</u>	<u>A205798 001</u>	Jun 01, 2017
<u>AB</u>		<u>30MG</u>	<u>A205798 002</u>	Jun 01, 2017
<u>AB</u>		<u>45MG</u>	<u>A205798 003</u>	Jun 01, 2017

REMERON SOLTAB

<u>AB</u>	+!	ORGANON USA INC	<u>15MG</u>	<u>N021208 001</u>	Jan 12, 2001
<u>AB</u>	+		<u>30MG</u>	<u>N021208 002</u>	Jan 12, 2001
<u>AB</u>	+		<u>45MG</u>	<u>N021208 003</u>	Jan 12, 2001

MISOPROSTOL

TABLET;ORAL

CYTOTEC

<u>AB</u>	+	GD SEARLE LLC	<u>0.1MG</u>	<u>N019268 003</u>	Sep 21, 1990
<u>AB</u>	+!		<u>0.2MG</u>	<u>N019268 001</u>	Dec 27, 1988

MISOPROSTOL

<u>AB</u>	IVAX SUB TEVA PHARMS	<u>0.1MG</u>	<u>A076095 001</u>	Jul 10, 2002
<u>AB</u>		<u>0.2MG</u>	<u>A076095 002</u>	Jul 10, 2002
<u>AB</u>	NOVEL LABS INC	<u>0.1MG</u>	<u>A091667 001</u>	Jul 25, 2012
<u>AB</u>		<u>0.2MG</u>	<u>A091667 002</u>	Jul 25, 2012

MITOMYCIN

FOR SOLUTION;TOPICAL

MITOSOL

<u>AB</u>	+!	MOBIUS THERAP	<u>0.2MG/VIAL</u>	<u>N022572 001</u>	Feb 07, 2012
-----------	----	---------------	-------------------	--------------------	--------------

INJECTABLE;INJECTION

MITOMYCIN

<u>AP</u>	!	ACCORD HLTHCARE	<u>5MG/VIAL</u>	<u>A064144 001</u>	Apr 30, 1998
<u>AP</u>	!		<u>20MG/VIAL</u>	<u>A064144 002</u>	Apr 30, 1998
<u>AP</u>	!		<u>40MG/VIAL</u>	<u>A064144 003</u>	Aug 11, 2009
<u>AP</u>	MYLAN LABS LTD		<u>5MG/VIAL</u>	<u>A202670 001</u>	Oct 13, 2017
<u>AP</u>			<u>20MG/VIAL</u>	<u>A202670 002</u>	Oct 13, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-294 (of 436)

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>		<u>40MG/VIAL</u>	<u>A203386 001</u>	Oct 13, 2017
<u>AP</u>	WEST-WARD PHARMS INT	<u>5MG/VIAL</u>	<u>A064117 001</u>	Apr 19, 1995
<u>AP</u>		<u>5MG/VIAL</u>	<u>A064180 001</u>	Dec 23, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A064117 002</u>	Apr 19, 1995
<u>AP</u>		<u>20MG/VIAL</u>	<u>A064180 002</u>	Dec 23, 1999

MITOTANE

TABLET; ORAL

LYSODREN

+!	BRISTOL MYERS SQUIBB	500MG	N016885 001
----	----------------------	-------	-------------

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077496 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077496 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077496 003</u>	Apr 11, 2006
<u>AP</u>	! HOSPIRA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076871 001</u>	Apr 11, 2006
<u>AP</u>	!	<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076871 002</u>	Apr 11, 2006
<u>AP</u>	!	<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076871 003</u>	Apr 11, 2006
<u>AP</u>	MYLAN INSTITUTIONAL	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A078980 001</u>	Apr 13, 2009
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A078980 002</u>	Apr 13, 2009
<u>AP</u>	MYLAN LABS LTD	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A201014 001</u>	Dec 11, 2012
<u>AP</u>	TEVA PHARMS USA	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A077356 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A077356 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A077356 003</u>	Apr 11, 2006
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 20MG BASE/10ML (EQ 2MG BASE/ML)</u>	<u>A076611 001</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)</u>	<u>A076611 002</u>	Apr 11, 2006
<u>AP</u>		<u>EQ 30MG BASE/15ML (EQ 2MG BASE/ML)</u>	<u>A076611 003</u>	Apr 11, 2006

MIVACURIUM CHLORIDE

SOLUTION; INTRAVENOUS

MIVACRON

+ ABBVIE	EQ 10MG BASE/5ML (EQ 2MG BASE/ML)	N020098 004
+!	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	N020098 005

Jan 22, 1992  
Jan 22, 1992

MODAFINIL

TABLET; ORAL

MODAFINIL

<u>AB</u>	ALEMBIC PHARMS LTD	<u>100MG</u>	<u>A202700 001</u>	Oct 18, 2012
<u>AB</u>		<u>200MG</u>	<u>A202700 002</u>	Oct 18, 2012
<u>AB</u>	APOTEX INC	<u>100MG</u>	<u>A077667 001</u>	Feb 03, 2014
<u>AB</u>		<u>200MG</u>	<u>A077667 002</u>	Feb 03, 2014
<u>AB</u>	APPCO PHARMA LLC	<u>100MG</u>	<u>A207196 001</u>	Aug 16, 2017
<u>AB</u>		<u>200MG</u>	<u>A207196 002</u>	Aug 16, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A202566 001</u>	Sep 27, 2012
<u>AB</u>		<u>200MG</u>	<u>A202566 002</u>	Sep 27, 2012
<u>AB</u>	HIKMA PHARMS	<u>100MG</u>	<u>A090543 001</u>	Sep 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A090543 002</u>	Sep 26, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>100MG</u>	<u>A076594 001</u>	Jul 16, 2012
<u>AB</u>		<u>200MG</u>	<u>A076594 002</u>	Jul 16, 2012
<u>AB</u>	ORCHID HLTHCARE	<u>100MG</u>	<u>A078963 001</u>	Sep 26, 2012
<u>AB</u>		<u>200MG</u>	<u>A078963 002</u>	Sep 26, 2012
<u>AB</u>	WATSON LABS INC	<u>100MG</u>	<u>A076715 001</u>	Nov 01, 2012
<u>AB</u>		<u>200MG</u>	<u>A076715 002</u>	Nov 01, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>100MG</u>	<u>A209966 001</u>	Sep 14, 2017
<u>AB</u>		<u>200MG</u>	<u>A209966 002</u>	Sep 14, 2017
	<u>PROVIGIL</u>			
<u>AB</u>	+ CEPHALON	<u>100MG</u>	<u>N020717 001</u>	Dec 24, 1998
<u>AB</u>	+!	<u>200MG</u>	<u>N020717 002</u>	Dec 24, 1998

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>7.5MG</u>	<u>A078454 001</u>	Jun 02, 2008
<u>AB</u>		<u>15MG</u>	<u>A078454 002</u>	Jun 02, 2008
<u>AB</u>	CHARTWELL RX	<u>7.5MG</u>	<u>A077536 001</u>	Nov 30, 2006
<u>AB</u>		<u>15MG</u>	<u>A077536 002</u>	Nov 30, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-295 (of 436)

MOEXIPRIL HYDROCHLORIDE

TABLET;ORAL

MOEXIPRIL HYDROCHLORIDE

<u>AB</u>	GLENMARK GENERICS	<u>7.5MG</u>	<u>A090416 001</u>	Mar 30, 2010
<u>AB</u>		<u>15MG</u>	<u>A090416 002</u>	Mar 30, 2010
<u>AB</u>	TEVA	<u>7.5MG</u>	<u>A076204 001</u>	May 08, 2003
<u>AB</u>	!	<u>15MG</u>	<u>A076204 002</u>	May 08, 2003

MOLINDONE HYDROCHLORIDE

TABLET;ORAL

MOLINDONE HYDROCHLORIDE

EPIC PHARMA LLC	5MG	A090453 001	Mar 20, 2015
	10MG	A090453 002	Mar 20, 2015
!	25MG	A090453 003	Mar 20, 2015

MOMETASONE FUROATE

AEROSOL, METERED;INHALATION

ASMANEX HFA

+	MERCK SHARP DOHME	0.10MG/INH	N205641 001	Apr 25, 2014
+!		0.20MG/INH	N205641 002	Apr 25, 2014

CREAM;TOPICAL

ELOC CON

<u>AB</u>	+!	MERCK SHARP DOHME	<u>0.1%</u>	<u>N019625 002</u>	Apr 19, 2013
-----------	----	-------------------	-------------	--------------------	--------------

MOMETASONE FUROATE

<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A076171 001</u>	Apr 08, 2005
<u>AB</u>	G AND W LABS	<u>0.1%</u>	<u>A077447 001</u>	May 22, 2006
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A078541 001</u>	May 28, 2008
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076679 001</u>	Dec 21, 2004
<u>AB</u>	TOLMAR	<u>0.1%</u>	<u>A076591 001</u>	Apr 18, 2007

IMPLANT;IMPLANTATION

SINUVA

+	INTERSECT ENT INC	1.35MG	N209310 001	Dec 08, 2017
---	-------------------	--------	-------------	--------------

LOTION;TOPICAL

ELOC CON

<u>AB</u>	+!	MERCK SHARP DOHME	<u>0.1%</u>	<u>N019796 001</u>	Mar 30, 1989
-----------	----	-------------------	-------------	--------------------	--------------

MOMETASONE FUROATE

<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A075919 001</u>	Nov 29, 2007
<u>AB</u>	G AND W LABS	<u>0.1%</u>	<u>A077678 001</u>	Nov 21, 2007
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A090506 001</u>	Aug 09, 2010
<u>AB</u>	PERRIGO ISRAEL	<u>0.1%</u>	<u>A077180 001</u>	Apr 06, 2005
<u>AB</u>	TARO	<u>0.1%</u>	<u>A076788 001</u>	Mar 15, 2006
<u>AB</u>	TOLMAR	<u>0.1%</u>	<u>A076499 001</u>	Nov 21, 2007

OINTMENT;TOPICAL

ELOC CON

<u>AB</u>	+!	MERCK SHARP DOHME	<u>0.1%</u>	<u>N019543 001</u>	Apr 30, 1987
-----------	----	-------------------	-------------	--------------------	--------------

MOMETASONE FUROATE

<u>AB</u>	FOUGERA PHARMS	<u>0.1%</u>	<u>A077061 001</u>	Mar 28, 2005
<u>AB</u>	G AND W LABS	<u>0.1%</u>	<u>A077401 001</u>	Jun 20, 2006
<u>AB</u>	GLENMARK GENERICS	<u>0.1%</u>	<u>A078571 001</u>	May 28, 2008
<u>AB</u>	PERRIGO NEW YORK	<u>0.1%</u>	<u>A076067 001</u>	Mar 18, 2002
<u>AB</u>	TOLMAR	<u>0.1%</u>	<u>A076481 001</u>	Nov 14, 2003

POWDER;INHALATION

ASMANEX TWISTHALER

+	MERCK SHARP DOHME	0.11MG/INH	N021067 002	Feb 01, 2008
+		0.22MG/INH	N021067 001	Mar 30, 2005

SPRAY, METERED;NASAL

MOMETASONE FUROATE

<u>AB</u>	AMNEAL PHARMS	<u>EQ 0.05MG BASE/SPRAY</u>	<u>A207989 001</u>	Apr 03, 2017
<u>AB</u>	APOTEX INC	<u>EQ 0.05MG BASE/SPRAY</u>	<u>A091161 001</u>	Mar 22, 2016

NASONEX

<u>AB</u>	+!	MERCK SHARP DOHME	<u>EQ 0.05MG BASE/SPRAY</u>	<u>N020762 001</u>	Oct 01, 1997
-----------	----	-------------------	-----------------------------	--------------------	--------------

MONTELUKAST SODIUM

GRANULE;ORAL

MONTELUKAST SODIUM

<u>AB</u>	AJANTA PHARMA LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A203438 001</u>	Jul 31, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/PACKET</u>	<u>A202906 001</u>	Sep 17, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 4MG BASE/PACKET</u>	<u>A202776 001</u>	Dec 18, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 4MG BASE/PACKET</u>	<u>A090955 001</u>	Aug 03, 2012

SINGULAIR

<u>AB</u>	+!	MERCK	<u>EQ 4MG BASE/PACKET</u>	<u>N021409 001</u>	Jul 26, 2002
-----------	----	-------	---------------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-296 (of 436)

**MONTELUKAST SODIUM**

TABLET;ORAL

**MONTELUKAST SODIUM**

<b><u>AB</u></b>	ACCORD HLTHCARE	<b><u>EQ 10MG BASE</u></b>	<b><u>A202717 001</u></b>	Sep 21, 2012
<b><u>AB</u></b>	AJANTA PHARMA LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A203432 001</u></b>	Jul 31, 2015
<b><u>AB</u></b>	AMNEAL PHARMS	<b><u>EQ 10MG BASE</u></b>	<b><u>A204604 001</u></b>	Sep 04, 2015
<b><u>AB</u></b>	ANBISON LAB CO LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A205683 001</u></b>	Jan 12, 2016
<b><u>AB</u></b>	APOTEX CORP	<b><u>EQ 10MG BASE</u></b>	<b><u>A201294 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A202468 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	CIPILA LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A207463 001</u></b>	Oct 28, 2016
<b><u>AB</u></b>	CSPC OUYI PHARM CO	<b><u>EQ 10MG BASE</u></b>	<b><u>A209012 001</u></b>	Apr 24, 2017
<b><u>AB</u></b>	DR REDDYS LABS LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A201582 001</u></b>	Aug 06, 2012
<b><u>AB</u></b>	GLENMARK GENERICS	<b><u>EQ 10MG BASE</u></b>	<b><u>A090926 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	HETERO LABS LTD V	<b><u>EQ 10MG BASE</u></b>	<b><u>A202843 001</u></b>	Sep 10, 2014
<b><u>AB</u></b>	KREMERS URBAN PHARMS	<b><u>EQ 10MG BASE</u></b>	<b><u>A201522 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	MACLEODS PHARMS LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A203366 001</u></b>	Sep 11, 2014
<b><u>AB</u></b>	MYLAN PHARMS INC	<b><u>EQ 10MG BASE</u></b>	<b><u>A079103 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	PERRIGO R AND D	<b><u>EQ 10MG BASE</u></b>	<b><u>A206112 001</u></b>	Apr 26, 2017
<b><u>AB</u></b>	SANDOZ INC	<b><u>EQ 10MG BASE</u></b>	<b><u>A200889 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	TEVA PHARMS	<b><u>EQ 10MG BASE</u></b>	<b><u>A078605 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	TORRENT PHARMS LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A201515 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	UNICHEM LABS LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A204290 001</u></b>	Oct 08, 2015
<b><u>AB</u></b>	UNIMARK REMEDIES LTD	<b><u>EQ 10MG BASE</u></b>	<b><u>A202859 001</u></b>	Oct 30, 2014
<b><u>AB</u></b>	VINTAGE PHARMS LLC	<b><u>EQ 10MG BASE</u></b>	<b><u>A091576 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>	WEST-WARD PHARMS INT	<b><u>EQ 10MG BASE</u></b>	<b><u>A090655 001</u></b>	Aug 03, 2012

**SINGULAIR**

<b><u>AB</u></b>	+! MSD MERCK CO	<b><u>EQ 10MG BASE</u></b>	<b><u>N020829 002</u></b>	Feb 20, 1998
------------------	-----------------	----------------------------	---------------------------	--------------

**MONTELUKAST SODIUM**

<b><u>AB</u></b>	AJANTA PHARMA LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A203328 001</u></b>	Jul 31, 2015
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A203328 002</u></b>	Jul 31, 2015
<b><u>AB</u></b>	ANBISON LAB CO LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A205695 001</u></b>	Nov 05, 2015
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A205695 002</u></b>	Nov 05, 2015
<b><u>AB</u></b>	APOTEX INC	<b><u>EQ 4MG BASE</u></b>	<b><u>A201508 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A201508 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A202096 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A202096 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	CSPC OUYI PHARM CO	<b><u>EQ 4MG BASE</u></b>	<b><u>A209011 001</u></b>	Apr 18, 2017
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A209011 002</u></b>	Apr 18, 2017
<b><u>AB</u></b>	DR REDDYS LABS LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A201581 001</u></b>	Aug 06, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A201581 002</u></b>	Aug 06, 2012
<b><u>AB</u></b>	HETERO LABS LTD V	<b><u>EQ 4MG BASE</u></b>	<b><u>A204093 001</u></b>	May 22, 2015
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A204093 002</u></b>	May 22, 2015
<b><u>AB</u></b>	JUBILANT GENERICS	<b><u>EQ 4MG BASE</u></b>	<b><u>A203795 001</u></b>	Feb 27, 2015
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A203795 002</u></b>	Feb 27, 2015
<b><u>AB</u></b>	KREMERS URBAN PHARMS	<b><u>EQ 4MG BASE</u></b>	<b><u>A200405 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A200405 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	MACLEODS PHARMS LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A203582 001</u></b>	Mar 12, 2015
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A203582 002</u></b>	Mar 12, 2015
<b><u>AB</u></b>	MYLAN PHARMS INC	<b><u>EQ 4MG BASE</u></b>	<b><u>A079142 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A079142 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	SANDOZ INC	<b><u>EQ 4MG BASE</u></b>	<b><u>A091414 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A091414 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	TEVA PHARMS	<b><u>EQ 4MG BASE</u></b>	<b><u>A078723 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A078723 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	TORRENT PHARMS LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A090984 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A090984 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	UNIMARK REMEDIES LTD	<b><u>EQ 4MG BASE</u></b>	<b><u>A203037 001</u></b>	Oct 30, 2014
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A203037 002</u></b>	Oct 30, 2014
<b><u>AB</u></b>	VINTAGE PHARMS LLC	<b><u>EQ 4MG BASE</u></b>	<b><u>A091588 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A091588 002</u></b>	Aug 03, 2012
<b><u>AB</u></b>	WEST-WARD PHARMS INT	<b><u>EQ 4MG BASE</u></b>	<b><u>A091128 001</u></b>	Aug 03, 2012
<b><u>AB</u></b>		<b><u>EQ 5MG BASE</u></b>	<b><u>A091128 002</u></b>	Aug 03, 2012

**SINGULAIR**

<b><u>AB</u></b>	+! MSD MERCK CO	<b><u>EQ 4MG BASE</u></b>	<b><u>N020830 002</u></b>	Mar 03, 2000
<b><u>AB</u></b>	++!	<b><u>EQ 5MG BASE</u></b>	<b><u>N020830 001</u></b>	Feb 20, 1998

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-297 (of 436)

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

KADIAN

<u>AB</u>	+!	ALLERGAN SALES LLC	<u>10MG</u>	<u>N020616 008</u>	Apr 20, 2007
<u>AB</u>	+		<u>20MG</u>	<u>N020616 001</u>	Jul 03, 1996
<u>AB</u>	+		<u>30MG</u>	<u>N020616 004</u>	Mar 09, 2001
<u>AB</u>	+		<u>40MG</u>	<u>N020616 009</u>	Jul 09, 2012
<u>AB</u>	+		<u>50MG</u>	<u>N020616 002</u>	Jul 03, 1996
<u>AB</u>	+		<u>60MG</u>	<u>N020616 005</u>	Mar 09, 2001
<u>AB</u>	+		<u>70MG</u>	<u>N020616 010</u>	Jul 09, 2012
<u>AB</u>	+		<u>80MG</u>	<u>N020616 006</u>	Oct 27, 2006
<u>AB</u>	+!		<u>100MG</u>	<u>N020616 003</u>	Jul 03, 1996

MORPHINE SULFATE

<u>AB</u>		IMPAK LABS INC	<u>20MG</u>	<u>A200411 001</u>	Apr 12, 2016
<u>AB</u>			<u>30MG</u>	<u>A200411 002</u>	Apr 12, 2016
<u>AB</u>			<u>50MG</u>	<u>A200411 003</u>	Apr 12, 2016
<u>AB</u>			<u>60MG</u>	<u>A200411 004</u>	Apr 12, 2016
<u>AB</u>			<u>80MG</u>	<u>A200411 005</u>	Apr 12, 2016
<u>AB</u>			<u>100MG</u>	<u>A200411 006</u>	Apr 12, 2016
<u>AB</u>		PAR PHARM INC	<u>20MG</u>	<u>A200812 001</u>	Nov 10, 2011
<u>AB</u>			<u>30MG</u>	<u>A200812 002</u>	Nov 10, 2011
<u>AB</u>			<u>50MG</u>	<u>A200812 003</u>	Nov 10, 2011
<u>AB</u>			<u>60MG</u>	<u>A200812 004</u>	Nov 10, 2011
<u>AB</u>			<u>80MG</u>	<u>A200812 005</u>	Nov 10, 2011
<u>AB</u>			<u>100MG</u>	<u>A200812 006</u>	Nov 10, 2011
<u>AB</u>		TEVA PHARMS USA	<u>20MG</u>	<u>A202718 001</u>	Dec 29, 2014
<u>AB</u>			<u>30MG</u>	<u>A202718 002</u>	Dec 29, 2014
<u>AB</u>			<u>40MG</u>	<u>A202718 007</u>	Jun 03, 2015
<u>AB</u>			<u>50MG</u>	<u>A202718 003</u>	Dec 29, 2014
<u>AB</u>			<u>60MG</u>	<u>A202718 004</u>	Dec 29, 2014
<u>AB</u>			<u>70MG</u>	<u>A202718 008</u>	Jun 03, 2015
<u>AB</u>			<u>80MG</u>	<u>A202718 005</u>	Dec 29, 2014
<u>AB</u>			<u>100MG</u>	<u>A202718 006</u>	Dec 29, 2014
<u>AB</u>		UPSHER-SMITH LABS	<u>10MG</u>	<u>A202104 001</u>	Jun 03, 2013
<u>AB</u>			<u>20MG</u>	<u>A202104 002</u>	Jun 03, 2013
<u>AB</u>			<u>30MG</u>	<u>A202104 003</u>	Jun 03, 2013
<u>AB</u>			<u>50MG</u>	<u>A202104 004</u>	Jun 03, 2013
<u>AB</u>			<u>60MG</u>	<u>A202104 005</u>	Jun 03, 2013
<u>AB</u>			<u>80MG</u>	<u>A202104 006</u>	Jun 03, 2013
<u>AB</u>			<u>100MG</u>	<u>A202104 007</u>	Jun 03, 2013

KADIAN

<u>AB</u>	+	ALLERGAN SALES LLC	<u>130MG</u>	<u>N020616 011</u>	Jul 09, 2012
<u>AB</u>	+		<u>150MG</u>	<u>N020616 012</u>	Jul 09, 2012
<u>AB</u>	+!		<u>200MG</u>	<u>N020616 007</u>	Feb 27, 2007

MORPHINE SULFATE

	ACTAVIS ELIZABETH	<u>30MG</u>	<u>A079040 001</u>	Jan 16, 2013
		<u>45MG</u>	<u>A079040 002</u>	Jan 16, 2013
		<u>60MG</u>	<u>A079040 003</u>	Jan 16, 2013
		<u>75MG</u>	<u>A079040 004</u>	Jan 16, 2013
		<u>90MG</u>	<u>A079040 005</u>	Jan 16, 2013
	!	<u>120MG</u>	<u>A079040 006</u>	Jan 16, 2013

INJECTABLE; INJECTION

ASTRAMORPH PF

<u>AP</u>		FRESENIUS KABI USA	<u>0.5MG/ML</u>	<u>A071050 001</u>	Oct 07, 1986
<u>AP</u>			<u>1MG/ML</u>	<u>A071051 001</u>	Oct 07, 1986
<u>AP</u>			<u>1MG/ML</u>	<u>A071052 001</u>	Oct 07, 1986
<u>AP</u>			<u>1MG/ML</u>	<u>A071053 001</u>	Oct 07, 1986

DURAMORPH PF

<u>AP</u>	+!	WEST-WARD PHARMS INT	<u>0.5MG/ML</u>	<u>N018565 001</u>	Sep 18, 1984
<u>AP</u>	+!		<u>1MG/ML</u>	<u>N018565 002</u>	Sep 18, 1984

MORPHINE SULFATE

<u>AP</u>		EUROHLTH INTL SARL	<u>4MG/ML</u>	<u>A205758 001</u>	May 21, 2015
<u>AP</u>			<u>8MG/ML</u>	<u>A205758 002</u>	May 21, 2015
<u>AP</u>			<u>10MG/ML</u>	<u>A205758 003</u>	May 21, 2015
<u>AP</u>		HOSPIRA	<u>0.5MG/ML</u>	<u>A071849 001</u>	May 11, 1988
<u>AP</u>			<u>0.5MG/ML</u>	<u>A073509 001</u>	Sep 30, 1992
<u>AP</u>			<u>1MG/ML</u>	<u>A071850 001</u>	May 11, 1988
<u>AP</u>			<u>1MG/ML</u>	<u>A073510 001</u>	Sep 30, 1992
<u>AP</u>	+!	HOSPIRA INC	<u>4MG/ML</u>	<u>N202515 002</u>	Nov 14, 2011
<u>AP</u>	+!		<u>8MG/ML</u>	<u>N202515 003</u>	Nov 14, 2011
<u>AP</u>	+!		<u>10MG/ML</u>	<u>N202515 004</u>	Nov 14, 2011
<u>AP</u>	+!	ICU MEDICAL INC	<u>1MG/ML</u>	<u>N019916 001</u>	Oct 30, 1992

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-298 (of 436)

MORPHINE SULFATE

INJECTABLE; INJECTION

INFUMORPH

+!	WEST-WARD PHARMS INT	10MG/ML	N018565 003	Jul 19, 1991
+!		25MG/ML	N018565 004	Jul 19, 1991
MORPHINE SULFATE				
+!	HOSPIRA INC	2MG/ML	N202515 001	Nov 14, 2011
+!	ICU MEDICAL INC	5MG/ML	N019916 002	Oct 27, 2006
+!	MERIDIAN MEDCL TECHN	15MG/ML	N019999 001	Jul 12, 1990

SOLUTION; INTRAMUSCULAR, INTRAVENOUS

MORPHINE SULFATE

+!	FRESENIUS KABI USA	2MG/ML (2MG/ML)	N204223 001	Oct 30, 2013
+!		4MG/ML (4MG/ML)	N204223 002	Oct 30, 2013
+!		5MG/ML (5MG/ML)	N204223 003	Oct 30, 2013
+!		8MG/ML (8MG/ML)	N204223 004	Oct 30, 2013
+!		10MG/ML (10MG/ML)	N204223 005	Oct 30, 2013

SOLUTION; ORAL

MORPHINE SULFATE

<u>AA</u>	HI-TECH PHARMACAL	<u>100MG/5ML</u>	<u>A208809 001</u>	Jul 06, 2017
<u>AA</u>	NOSTRUM LABS INC	<u>10MG/5ML</u>	<u>A201011 001</u>	Feb 05, 2014
<u>AA</u>		<u>20MG/5ML</u>	<u>A201011 002</u>	Feb 05, 2014
<u>AA</u>		<u>100MG/5ML</u>	<u>A204053 001</u>	Oct 06, 2016
<u>AA</u>	PADDOCK LLC	<u>100MG/5ML</u>	<u>A201574 001</u>	Aug 06, 2012
<u>AA</u>	PHARM ASSOC	<u>100MG/5ML</u>	<u>A206573 001</u>	Nov 14, 2016
<u>AA</u>	RHODES PHARMS	<u>10MG/5ML</u>	<u>A206308 001</u>	Jun 22, 2017
<u>AA</u>		<u>20MG/5ML</u>	<u>A206420 001</u>	Jul 12, 2016
<u>AA</u>		<u>100MG/5ML</u>	<u>A206308 002</u>	Jun 22, 2017
<u>AA</u>	SPECGX LLC	<u>100MG/5ML</u>	<u>A202348 001</u>	Jul 15, 2011
<u>AA</u>	TRIS PHARMA INC	<u>10MG/5ML</u>	<u>A203518 001</u>	May 12, 2015
<u>AA</u>		<u>20MG/5ML</u>	<u>A203519 001</u>	May 18, 2016
<u>AA</u>		<u>100MG/5ML</u>	<u>A203518 002</u>	May 12, 2015
<u>AA</u>	VINTAGE PHARMS LLC	<u>10MG/5ML</u>	<u>A202309 001</u>	Nov 25, 2015
<u>AA</u>		<u>20MG/5ML</u>	<u>A202310 001</u>	Oct 30, 2015
<u>AA</u>	VISTAPHARM	<u>10MG/5ML</u>	<u>A201947 001</u>	Jan 05, 2012
<u>AA</u>		<u>20MG/5ML</u>	<u>A201947 002</u>	Jan 05, 2012
<u>AA</u>	+ WEST-WARD PHARMS INT	<u>10MG/5ML</u>	<u>N022195 001</u>	Mar 17, 2008
<u>AA</u>	+ +	<u>20MG/5ML</u>	<u>N022195 002</u>	Mar 17, 2008
<u>AA</u>	+ !	<u>100MG/5ML</u>	<u>N022195 003</u>	Jan 25, 2010
	LANNETT HOLDINGS INC	100MG/5ML	N201517 001	Jun 23, 2011

TABLET; ORAL

MORPHINE SULFATE

+ WEST-WARD PHARMS INT	15MG	N022207 001	Mar 17, 2008
+ !	30MG	N022207 002	Mar 17, 2008

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

<u>AB</u>	ACTAVIS ELIZABETH	<u>15MG</u>	<u>A203849 001</u>	Apr 06, 2015
<u>AB</u>		<u>30MG</u>	<u>A203849 002</u>	Apr 06, 2015
<u>AB</u>		<u>60MG</u>	<u>A203849 003</u>	Apr 06, 2015
<u>AB</u>		<u>100MG</u>	<u>A203849 004</u>	Apr 06, 2015
<u>AB</u>		<u>200MG</u>	<u>A203849 005</u>	Apr 06, 2015
<u>AB</u>	DAVA PHARMS INC	<u>15MG</u>	<u>A075407 001</u>	Jan 28, 2000
<u>AB</u>	EPIC PHARMA LLC	<u>15MG</u>	<u>A091357 001</u>	Jun 23, 2016
<u>AB</u>		<u>30MG</u>	<u>A091357 002</u>	Jun 23, 2016
<u>AB</u>		<u>60MG</u>	<u>A091357 003</u>	Jun 23, 2016
<u>AB</u>		<u>100MG</u>	<u>A091357 004</u>	Jun 23, 2016
<u>AB</u>		<u>200MG</u>	<u>A091357 005</u>	Jun 23, 2016
<u>AB</u>	MAYNE PHARMA INC	<u>15MG</u>	<u>A205386 001</u>	Oct 28, 2016
<u>AB</u>		<u>30MG</u>	<u>A205386 002</u>	Oct 28, 2016
<u>AB</u>		<u>60MG</u>	<u>A205386 003</u>	Oct 28, 2016
<u>AB</u>		<u>100MG</u>	<u>A205386 004</u>	Oct 28, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>15MG</u>	<u>A200824 001</u>	Oct 18, 2011
<u>AB</u>		<u>30MG</u>	<u>A200824 002</u>	Oct 18, 2011
<u>AB</u>		<u>60MG</u>	<u>A200824 003</u>	Oct 18, 2011
<u>AB</u>		<u>100MG</u>	<u>A200824 004</u>	Oct 18, 2011
<u>AB</u>		<u>200MG</u>	<u>A200824 005</u>	Oct 18, 2011
<u>AB</u>	NESHER PHARMS	<u>15MG</u>	<u>A076733 001</u>	May 19, 2004
<u>AB</u>		<u>30MG</u>	<u>A076720 002</u>	Dec 23, 2005
<u>AB</u>		<u>60MG</u>	<u>A076720 001</u>	May 19, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-299 (of 436)

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

**MORPHINE SULFATE**

<u>AB</u>		<u>100MG</u>	<u>A077855</u>	<u>001</u>	Sep 27, 2007
<u>AB</u>		<u>200MG</u>	<u>A077855</u>	<u>002</u>	Sep 27, 2007
<u>AB</u>	NOVEL LABS INC	<u>15MG</u>	<u>A203602</u>	<u>001</u>	Dec 16, 2015
<u>AB</u>		<u>30MG</u>	<u>A203602</u>	<u>002</u>	Dec 16, 2015
<u>AB</u>		<u>60MG</u>	<u>A203602</u>	<u>003</u>	Dec 16, 2015
<u>AB</u>		<u>100MG</u>	<u>A203602</u>	<u>004</u>	Dec 16, 2015
<u>AB</u>		<u>200MG</u>	<u>A203602</u>	<u>005</u>	Dec 16, 2015
<u>AB</u>	RHODES PHARMS	<u>15MG</u>	<u>A074862</u>	<u>001</u>	Jul 07, 1998
<u>AB</u>		<u>30MG</u>	<u>A074862</u>	<u>002</u>	Jul 07, 1998
<u>AB</u>		<u>60MG</u>	<u>A074862</u>	<u>003</u>	Jul 07, 1998
<u>AB</u>		<u>100MG</u>	<u>A074769</u>	<u>001</u>	Jul 02, 1998
<u>AB</u>		<u>200MG</u>	<u>A074769</u>	<u>002</u>	Jul 02, 1998
<u>AB</u>	SPECGX LLC	<u>15MG</u>	<u>A076412</u>	<u>001</u>	Jul 31, 2003
<u>AB</u>		<u>30MG</u>	<u>A076412</u>	<u>002</u>	Jul 31, 2003
<u>AB</u>		<u>60MG</u>	<u>A076412</u>	<u>003</u>	Jul 31, 2003
<u>AB</u>		<u>100MG</u>	<u>A076438</u>	<u>001</u>	Jul 03, 2003
<u>AB</u>		<u>200MG</u>	<u>A076438</u>	<u>002</u>	Jul 03, 2003
<u>AB</u>	SUN PHARM INDS LTD	<u>15MG</u>	<u>A078761</u>	<u>001</u>	May 11, 2012
<u>AB</u>		<u>30MG</u>	<u>A078761</u>	<u>002</u>	May 11, 2012
<u>AB</u>		<u>60MG</u>	<u>A078761</u>	<u>003</u>	May 11, 2012
<u>AB</u>		<u>100MG</u>	<u>A078761</u>	<u>004</u>	May 11, 2012
<u>AB</u>		<u>200MG</u>	<u>A078761</u>	<u>005</u>	May 11, 2012
<u>AB</u>	SUN PHARM INDUSTRIES	<u>15MG</u>	<u>A205634</u>	<u>001</u>	Aug 25, 2016
<u>AB</u>		<u>30MG</u>	<u>A205634</u>	<u>002</u>	Aug 25, 2016
<u>AB</u>		<u>60MG</u>	<u>A205634</u>	<u>003</u>	Aug 25, 2016
<u>AB</u>		<u>100MG</u>	<u>A205634</u>	<u>004</u>	Aug 25, 2016
<u>AB</u>		<u>200MG</u>	<u>A205634</u>	<u>005</u>	Aug 25, 2016
<u>AB</u>	VINTAGE PHARMS LLC	<u>15MG</u>	<u>A075295</u>	<u>001</u>	Oct 28, 1998
<u>AB</u>		<u>30MG</u>	<u>A075295</u>	<u>002</u>	Oct 28, 1998
<u>AB</u>		<u>60MG</u>	<u>A075295</u>	<u>003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>A075295</u>	<u>004</u>	Sep 15, 2000
<u>AB</u>		<u>200MG</u>	<u>A075295</u>	<u>005</u>	Sep 15, 2000
<b>MS CONTIN</b>					
<u>AB</u>	+ PURDUE PHARMA LP	<u>15MG</u>	<u>N019516</u>	<u>003</u>	Sep 12, 1989
<u>AB</u>	+	<u>30MG</u>	<u>N019516</u>	<u>001</u>	May 29, 1987
<u>AB</u>	+	<u>60MG</u>	<u>N019516</u>	<u>002</u>	Apr 08, 1988
<u>AB</u>	+!	<u>100MG</u>	<u>N019516</u>	<u>004</u>	Jan 16, 1990
<u>AB</u>	+	<u>200MG</u>	<u>N019516</u>	<u>005</u>	Nov 08, 1993
ARYMO ER					
	+ EGALET	15MG	N208603	001	Jan 09, 2017
	+	30MG	N208603	002	Jan 09, 2017
	+	60MG	N208603	003	Jan 09, 2017
MORPHABOND ER					
	+ DAIICHI SANKYO INC	15MG	N206544	001	Oct 02, 2015
	+	30MG	N206544	002	Oct 02, 2015
	+	60MG	N206544	003	Oct 02, 2015
	+!	100MG	N206544	004	Oct 02, 2015
<b>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE</b>					
CAPSULE, EXTENDED RELEASE; ORAL					
EMBEDA					
	+ ALPHARMA PHARMS	20MG; 0.8MG	N022321	001	Aug 13, 2009
	+	30MG; 1.2MG	N022321	002	Aug 13, 2009
	+	50MG; 2MG	N022321	003	Aug 13, 2009
	+!	60MG; 2.4MG	N022321	004	Aug 13, 2009
	+	80MG; 3.2MG	N022321	005	Aug 13, 2009
	+	100MG; 4MG	N022321	006	Aug 13, 2009
<b>MOXIFLOXACIN HYDROCHLORIDE</b>					
SOLUTION; IV (INFUSION)					
MOXIFLOXACIN HYDROCHLORIDE					
+!	FRESENIUS KABI USA	EQ 400MG BASE/250ML (EQ 1.6MG BASE/ML)	N205572	001	Apr 03, 2015
MOXIFLOXACIN HYDROCHLORIDE		IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER			
!	MYLAN LABS LTD	400MG/250ML (1.6MG/ML)	A205833	001	May 05, 2017
SOLUTION/DROPS; OPHTHALMIC					
<b>MOXIFLOXACIN HYDROCHLORIDE</b>					
<b>AT1</b>	AKORN	<u>EQ 0.5% BASE</u>	<u>A202916</u>	<u>001</u>	Nov 09, 2017
<b>AT1</b>	APOTEX INC	<u>EQ 0.5% BASE</u>	<u>A090080</u>	<u>001</u>	Jun 30, 2017
<b>AT1</b>	AUROBINDO PHARMA LTD	<u>EQ 0.5% BASE</u>	<u>A206242</u>	<u>001</u>	Oct 04, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-300 (of 436)

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

MOXIFLOXACIN HYDROCHLORIDE

<u>AT1</u>	LUPIN LTD	<u>EQ 0.5% BASE</u>	<u>A202867 001</u>	Sep 04, 2014
<u>AT1</u>	WATSON LABS INC	<u>EQ 0.5% BASE</u>	<u>A202525 001</u>	Mar 06, 2015
	VIGAMOX			
<u>AT1</u>	+! NOVARTIS PHARMS CORP	<u>EQ 0.5% BASE</u>	<u>N021598 001</u>	Apr 15, 2003
	MOXEZA			
<u>AT2</u>	+! NOVARTIS PHARMS CORP	<u>EQ 0.5% BASE</u>	<u>N022428 001</u>	Nov 19, 2010
	<u>MOXIFLOXACIN HYDROCHLORIDE</u>			
<u>AT2</u>	LUPIN LTD	<u>EQ 0.5% BASE</u>	<u>A204079 001</u>	May 28, 2015
	TABLET;ORAL			
	AVELOX			
<u>AB</u>	+! BAYER HLTHCARE	<u>EQ 400MG BASE</u>	<u>N021085 001</u>	Dec 10, 1999
	<u>MOXIFLOXACIN HYDROCHLORIDE</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 400MG BASE</u>	<u>A202632 001</u>	Mar 04, 2014
<u>AB</u>	CROSSMEDIKA SA	<u>EQ 400MG BASE</u>	<u>A205348 001</u>	Jan 14, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 400MG BASE</u>	<u>A076938 001</u>	Mar 04, 2014
<u>AB</u>	MSN LABS PVT LTD	<u>EQ 400MG BASE</u>	<u>A208682 001</u>	Sep 22, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 400MG BASE</u>	<u>A204635 001</u>	Aug 31, 2015
<u>AB</u>	NOVEL LABS INC	<u>EQ 400MG BASE</u>	<u>A207285 001</u>	Feb 13, 2017
<u>AB</u>	TEVA PHARMS USA	<u>EQ 400MG BASE</u>	<u>A077437 001</u>	Feb 18, 2014
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 400MG BASE</u>	<u>A200160 001</u>	Apr 03, 2014

MUPIROCIN

OINTMENT;TOPICAL

MUPIROCIN

<u>AB</u>	FOUGERA PHARMS	<u>2%</u>	<u>A065192 001</u>	Nov 30, 2005
<u>AB</u>	GLENMARK PHARMS	<u>2%</u>	<u>A090480 001</u>	Jun 08, 2011
<u>AB</u>	! PERRIGO NEW YORK	<u>2%</u>	<u>A065123 001</u>	Nov 07, 2003
<u>AB</u>	TARO	<u>2%</u>	<u>A065170 001</u>	Sep 23, 2005
<u>AB</u>	TEVA	<u>2%</u>	<u>A065085 001</u>	Nov 07, 2003
	CENTANY			
BX	PERRIGO NEW YORK	2%		

MUPIROCIN CALCIUM

CREAM;TOPICAL

BACTROBAN

<u>AB</u>	+! GLAXOSMITHKLINE	<u>EQ 2% BASE</u>	<u>N050746 001</u>	Dec 11, 1997
	<u>MUPIROCIN</u>			
<u>AB</u>	GLENMARK PHARMS INC	<u>EQ 2% BASE</u>	<u>A201587 001</u>	Jan 24, 2013
	OINTMENT;NASAL			
	BACTROBAN			
	+! GLAXOSMITHKLINE	EQ 2% BASE		

MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>250MG</u>	<u>N050722 001</u>	May 03, 1995
	<u>MYCOPHENOLATE MOFETIL</u>			
<u>AB</u>	ACCORD HLTHCARE	<u>250MG</u>	<u>A090253 001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>250MG</u>	<u>A200197 001</u>	Jun 13, 2013
<u>AB</u>	APOTEX CORP	<u>250MG</u>	<u>A090419 001</u>	Apr 22, 2009
<u>AB</u>	MYLAN	<u>250MG</u>	<u>A065520 001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>A065379 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>250MG</u>	<u>A090055 001</u>	Jun 10, 2010
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>A065491 001</u>	May 06, 2009
<u>AB</u>	VINTAGE PHARMS LLC	<u>250MG</u>	<u>A090111 001</u>	Dec 22, 2009
<u>AB</u>	WEST-WARD PHARMS INT	<u>250MG</u>	<u>A065410 001</u>	Jul 29, 2008
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>250MG</u>	<u>A204077 001</u>	Nov 13, 2017

SUSPENSION;ORAL

CELLCEPT

<u>AB</u>	+! ROCHE PALO	<u>200MG/ML</u>	<u>N050759 001</u>	Oct 01, 1998
	<u>MYCOPHENOLATE MOFETIL</u>			
<u>AB</u>	ALKEM LABS LTD	<u>200MG/ML</u>	<u>A203005 001</u>	Nov 14, 2014
	TABLET;ORAL			
	<u>CELLCEPT</u>			
<u>AB</u>	+! ROCHE PALO	<u>500MG</u>	<u>N050723 001</u>	Jun 19, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-301 (of 436)

MYCOPHENOLATE MOFETIL

TABLET; ORAL

MYCOPHENOLATE MOFETIL

<u>AB</u>	ACCORD HLTHCARE	<u>500MG</u>	<u>A065416 001</u>	May 04, 2009
<u>AB</u>	ALKEM LABS LTD	<u>500MG</u>	<u>A091249 001</u>	Nov 04, 2011
<u>AB</u>	APOTEX	<u>500MG</u>	<u>A090499 001</u>	Apr 22, 2009
<u>AB</u>	MYLAN	<u>500MG</u>	<u>A065521 001</u>	May 04, 2009
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A065451 001</u>	Oct 15, 2008
<u>AB</u>	STRIDES PHARMA	<u>500MG</u>	<u>A090456 001</u>	Jun 10, 2010
<u>AB</u>	TEVA PHARMS	<u>500MG</u>	<u>A065457 001</u>	May 04, 2009
<u>AB</u>	VINTAGE PHARMS LLC	<u>500MG</u>	<u>A090606 001</u>	Jul 16, 2010
<u>AB</u>	WEST-WARD PHARMS INT	<u>500MG</u>	<u>A065413 001</u>	Jul 29, 2008
<u>AB</u>	ZHEJIANG HISUN PHARM	<u>500MG</u>	<u>A204076 001</u>	Nov 16, 2017

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

CELLCEPT

<u>AP</u>	+!	ROCHE PALO	<u>500MG/VIAL</u>	<u>N050758 001</u>	Aug 12, 1998
<u>MYCOPHENOLATE MOFETIL HYDROCHLORIDE</u>					
<u>AP</u>		AKORN INC	<u>500MG/VIAL</u>	<u>A204043 001</u>	Feb 28, 2017
<u>AP</u>		MYLAN LABS LTD	<u>500MG/VIAL</u>	<u>A203859 001</u>	Mar 31, 2017
<u>AP</u>		PAR STERILE PRODUCTS	<u>500MG/VIAL</u>	<u>A203575 001</u>	Oct 28, 2016
<u>AP</u>		ZYDUS PHARMS USA INC	<u>500MG/VIAL</u>	<u>A204473 001</u>	Aug 31, 2017

MYCOPHENOLIC ACID

TABLET, DELAYED RELEASE; ORAL

MYCOPHENOLIC ACID

<u>AB</u>	ACCORD HLTHCARE	<u>180MG</u>	<u>A202555 001</u>	Aug 23, 2017	
<u>AB</u>		<u>360MG</u>	<u>A202555 002</u>	Aug 23, 2017	
<u>AB</u>	APOTEX INC	<u>180MG</u>	<u>A091558 001</u>	Aug 21, 2012	
<u>AB</u>		<u>360MG</u>	<u>A091558 002</u>	Aug 19, 2014	
<u>AB</u>	MYLAN PHARMS INC	<u>180MG</u>	<u>A091248 002</u>	Jan 08, 2014	
<u>AB</u>		<u>360MG</u>	<u>A091248 001</u>	Jan 08, 2014	
<u>AB</u>	TEVA PHARMS USA	<u>180MG</u>	<u>A202720 001</u>	Oct 30, 2014	
<u>AB</u>		<u>360MG</u>	<u>A202720 002</u>	Oct 30, 2014	
<u>MYFORTIC</u>					
<u>AB</u>	+	NOVARTIS	<u>180MG</u>	<u>N050791 001</u>	Feb 27, 2004
<u>AB</u>	+		<u>360MG</u>	<u>N050791 002</u>	Feb 27, 2004

NABILONE

CAPSULE; ORAL

CESAMET

+! MYLAN SPECIALITY LP 1MG

N018677 001 Dec 26, 1985

NABUMETONE

TABLET; ORAL

NABUMETONE

<u>AB</u>	APOTEX INC	<u>500MG</u>	<u>A090427 001</u>	Dec 30, 2011
<u>AB</u>		<u>750MG</u>	<u>A090427 002</u>	Dec 30, 2011
<u>AB</u>	CHARTWELL MOLECULES	<u>500MG</u>	<u>A076009 001</u>	Jan 24, 2003
<u>AB</u>		<u>750MG</u>	<u>A076009 002</u>	Jan 24, 2003
<u>AB</u>	IMPAX LABS INC	<u>500MG</u>	<u>A075189 001</u>	May 26, 2000
<u>AB</u>	!	<u>750MG</u>	<u>A075189 002</u>	Sep 24, 2001
<u>AB</u>	INVAGEN PHARMS	<u>500MG</u>	<u>A078671 001</u>	Mar 07, 2008
<u>AB</u>		<u>750MG</u>	<u>A078671 002</u>	Mar 07, 2008
<u>AB</u>	LUPIN LTD	<u>500MG</u>	<u>A090445 001</u>	Jan 12, 2011
<u>AB</u>		<u>750MG</u>	<u>A090445 002</u>	Jan 12, 2011
<u>AB</u>	MYLAN PHARMS INC	<u>500MG</u>	<u>A090516 001</u>	Jul 12, 2010
<u>AB</u>		<u>750MG</u>	<u>A090516 002</u>	Jul 12, 2010
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>A075280 001</u>	Feb 25, 2002
<u>AB</u>		<u>750MG</u>	<u>A075280 002</u>	Feb 25, 2002
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>A091083 001</u>	Jun 13, 2011
<u>AB</u>		<u>750MG</u>	<u>A091083 002</u>	Jun 13, 2011

NADOLOL

TABLET; ORAL

CORGARD

<u>AB</u>	+	US WORLDMEDS LLC	<u>20MG</u>	<u>N018063 005</u>	Oct 28, 1986
<u>AB</u>	+		<u>40MG</u>	<u>N018063 001</u>	
<u>AB</u>	+		<u>80MG</u>	<u>N018063 002</u>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-302 (of 436)

NADOLOL

TABLET;ORAL

**NADOLOL**

<u>AB</u>	AMNEAL PHARMS CO	<u>20MG</u>	<u>A208832</u> <u>001</u>	Jun 02, 2017
<u>AB</u>		<u>40MG</u>	<u>A208832</u> <u>002</u>	Jun 02, 2017
<u>AB</u>		<u>80MG</u>	<u>A208832</u> <u>003</u>	Jun 02, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A201893</u> <u>001</u>	Sep 16, 2015
<u>AB</u>		<u>80MG</u>	<u>A201893</u> <u>002</u>	Sep 16, 2015
<u>AB</u>	INVAGEN PHARMS	<u>20MG</u>	<u>A203455</u> <u>001</u>	Dec 18, 2015
<u>AB</u>		<u>40MG</u>	<u>A203455</u> <u>002</u>	Dec 18, 2015
<u>AB</u>		<u>80MG</u>	<u>A203455</u> <u>003</u>	Dec 18, 2015
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>20MG</u>	<u>A074229</u> <u>001</u>	Aug 30, 1996
<u>AB</u>		<u>40MG</u>	<u>A074229</u> <u>002</u>	Aug 30, 1996
<u>AB</u>		<u>80MG</u>	<u>A074255</u> <u>001</u>	Jan 24, 1996
<u>AB</u>	LUPIN LTD	<u>20MG</u>	<u>A209309</u> <u>001</u>	Oct 05, 2017
<u>AB</u>		<u>40MG</u>	<u>A209309</u> <u>002</u>	Oct 05, 2017
<u>AB</u>		<u>80MG</u>	<u>A209309</u> <u>003</u>	Oct 05, 2017
<u>AB</u>	MYLAN	<u>20MG</u>	<u>A074172</u> <u>001</u>	Oct 31, 1993
<u>AB</u>		<u>40MG</u>	<u>A074172</u> <u>002</u>	Oct 31, 1993
<u>AB</u>		<u>80MG</u>	<u>A074172</u> <u>003</u>	Oct 31, 1993
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>A074501</u> <u>001</u>	Nov 09, 1995
<u>AB</u>		<u>40MG</u>	<u>A074501</u> <u>002</u>	Nov 09, 1995
<u>AB</u>		<u>80MG</u>	<u>A074501</u> <u>003</u>	Nov 09, 1995
<u>AB</u>	ZYDUS PHARMS USA INC	<u>20MG</u>	<u>A207761</u> <u>001</u>	Jul 28, 2017
<u>AB</u>		<u>40MG</u>	<u>A207761</u> <u>002</u>	Jul 28, 2017
<u>AB</u>		<u>80MG</u>	<u>A207761</u> <u>003</u>	Jul 28, 2017

NAFARELIN ACETATE

SPRAY, METERED;NASAL  
 SYNAREL

+! GD SEARLE LLC EQ 0.2MG BASE/SPRAY

N019886 001 Feb 13, 1990

NAFCILLIN SODIUM

INJECTABLE;INJECTION

**NAFCILLIN SODIUM**

<u>AP</u>	ANTIBOTICE	<u>EQ 1GM BASE/VIAL</u>	<u>A090560</u> <u>001</u>	Oct 03, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090560</u> <u>002</u>	Oct 03, 2011
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A091613</u> <u>001</u>	Dec 26, 2012
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A091613</u> <u>002</u>	Dec 26, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A091614</u> <u>001</u>	Dec 26, 2012
<u>AP</u>	ISTITUTO BIO ITA SPA	<u>EQ 1GM BASE/VIAL</u>	<u>A090002</u> <u>001</u>	Jun 30, 2011
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090002</u> <u>002</u>	Jun 30, 2011
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090005</u> <u>001</u>	Apr 20, 2011
<u>AP</u>	MYLAN LABS LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A200002</u> <u>001</u>	Apr 07, 2014
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A200002</u> <u>002</u>	Apr 07, 2014
<u>AP</u>	SAGENT PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A090582</u> <u>001</u>	Aug 24, 2012
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A090582</u> <u>002</u>	Aug 24, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A090580</u> <u>001</u>	Aug 24, 2012
<u>AP</u>	! SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A062527</u> <u>002</u>	Aug 02, 1984
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062732</u> <u>001</u>	Dec 23, 1986
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062527</u> <u>003</u>	Aug 02, 1984
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062732</u> <u>002</u>	Dec 23, 1986
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A062527</u> <u>004</u>	Aug 02, 1984

NALLPEN IN PLASTIC CONTAINER

+! BAXTER HLTHCARE EQ 20MG BASE/ML  
 +! EQ 2GM BASE/100ML

N050655 001 Oct 31, 1989  
 N050655 002 Oct 31, 1989

NAFTIFINE HYDROCHLORIDE

CREAM;TOPICAL

**NAFTIFINE HYDROCHLORIDE**

<u>AB</u>	TARO	<u>1%</u>	<u>A205975</u> <u>001</u>	Sep 08, 2016
<u>AB</u>		<u>2%</u>	<u>A206901</u> <u>001</u>	Jan 06, 2016
<u>AB</u>	TOLMAR	<u>2%</u>	<u>A206960</u> <u>001</u>	Apr 10, 2017

**NAFTIN**

<u>AB</u>	+! SEBELA IRELAND LTD	<u>1%</u>	<u>N019599</u> <u>001</u>	Feb 29, 1988
<u>AB</u>	+!	<u>2%</u>	<u>N019599</u> <u>002</u>	Jan 13, 2012

GEL;TOPICAL

NAFTIN

+!	SEBELA IRELAND LTD	1%	N019356	001 Jun 18, 1990
+!		2%	N204286	001 Jun 27, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-303 (of 436)

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

<u>AP</u>	!	HOSPIRA	<u>10MG/ML</u>	<u>A070914 001</u>	Feb 03, 1989
<u>AP</u>	!		<u>10MG/ML</u>	<u>A070915 001</u>	Feb 03, 1989
<u>AP</u>	!		<u>20MG/ML</u>	<u>A070916 001</u>	Feb 03, 1989
<u>AP</u>	!		<u>20MG/ML</u>	<u>A070918 001</u>	Feb 03, 1989

NALDEMEDINE TOSYLATE

TABLET; ORAL

SYMPROIC

+! SHIONOGI INC

EQ 0.2MG BASE

N208854 001 Mar 23, 2017

NALOXEGOL OXALATE

TABLET; ORAL

MOVANTIK

+ ASTRAZENECA PHARMS

EQ 12.5MG BASE

N204760 001 Sep 16, 2014

+!

EQ 25MG BASE

N204760 002 Sep 16, 2014

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

<u>AP</u>		WEST-WARD PHARMS	<u>0 . 4MG/ML</u>	<u>A070299 001</u>	Sep 24, 1986
	INT				

NALOXONE HYDROCHLORIDE

<u>AP</u>		AKORN	<u>0 . 4MG/ML</u>	<u>A208871 001</u>	Feb 28, 2017
<u>AP</u>			<u>0 . 4MG/ML</u>	<u>A208872 001</u>	Mar 14, 2017
<u>AP</u>	!	HOSPIRA	<u>0 . 4MG/ML</u>	<u>A070172 001</u>	Sep 24, 1986
<u>AP</u>	!		<u>0 . 4MG/ML</u>	<u>A070254 001</u>	Jan 07, 1987
<u>AP</u>	!		<u>0 . 4MG/ML</u>	<u>A070256 001</u>	Jan 07, 1987
<u>AP</u>	!		<u>0 . 4MG/ML</u>	<u>A070257 001</u>	Jan 07, 1987
<u>AP</u>		INTL MEDICATION	<u>0 . 4MG/ML</u>	<u>A070639 001</u>	Sep 24, 1986
<u>AP</u>		MYLAN INSTITUTIONAL	<u>0 . 4MG/ML</u>	<u>A204997 001</u>	Mar 06, 2014
<u>AP</u>			<u>0 . 4MG/ML</u>	<u>A205014 001</u>	Jun 29, 2016
<u>AP</u>		SOMERSET THERAPS LLC	<u>0 . 4MG/ML</u>	<u>A207633 001</u>	Aug 08, 2017
<u>AP</u>			<u>0 . 4MG/ML</u>	<u>A207634 001</u>	Jul 26, 2017
	!	INTL MEDICATION	1MG/ML	A072076 001	Mar 24, 1988
		SOLUTION; INTRAMUSCULAR, SUBCUTANEOUS			
		EVZIO			
		+! KALEO INC	2MG/0.4ML (2MG/0.4ML)	N209862 001	Oct 19, 2016
		SPRAY, METERED; NASAL			
		NARCAN			
		+ ADAPT	2MG/SPRAY	N208411 002	Jan 24, 2017
		+!	4MG/SPRAY	N208411 001	Nov 18, 2015

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE

<u>AB</u>		GAVIS PHARMS	<u>EQ 0 . 5MG BASE;EQ 50MG BASE</u>	<u>A075735 001</u>	Jul 11, 2001
<u>AB</u>		SUN PHARM INDs LTD	<u>EQ 0 . 5MG BASE;EQ 50MG BASE</u>	<u>A075523 001</u>	Mar 17, 2000
<u>AB</u>	!	WATSON LABS	<u>EQ 0 . 5MG BASE;EQ 50MG BASE</u>	<u>A074736 001</u>	Jan 21, 1997

NALTREXONE

FOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

VIVITROL

+! ALKERMES

380MG/VIAL

N021897 001 Apr 13, 2006

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

<u>AB</u>		ACCORD HLTHCARE	<u>50MG</u>	<u>A091205 001</u>	Aug 17, 2011
<u>AB</u>		APOTEX INC	<u>50MG</u>	<u>A207905 001</u>	Jul 21, 2017
<u>AB</u>		BARR	<u>50MG</u>	<u>A074918 001</u>	May 08, 1998
<u>AB</u>		ELITE LABS	<u>50MG</u>	<u>A075274 001</u>	May 26, 1999
<u>AB</u>	!	SPECGX LLC	<u>50MG</u>	<u>A076264 002</u>	Mar 22, 2002
<u>AB</u>		SUN PHARMA GLOBAL	<u>50MG</u>	<u>A090356 001</u>	Feb 24, 2012
		SPECGX LLC	25MG	A076264 001	Mar 22, 2002
			100MG	A076264 003	Mar 22, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-304 (of 436)

NANDROLONE DECANOATE

INJECTABLE; INJECTION  
 NANDROLONE DECANOATE  
 ! LUITPOLD PHARMS INC 200MG/ML

A091252 001 Aug 30, 2010

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC  
 NAPHAZOLINE HYDROCHLORIDE  
 ! AKORN INC 0.1%

A083590 001

NAPROXEN

SUSPENSION; ORAL

NAPROSYN

AB +! ATNAHS PHARMA US 25MG/ML

N018965 001 Mar 23, 1987

NAPROXEN

AB WEST-WARD PHARMS INT 25MG/ML

A074190 001 Mar 30, 1994

TABLET; ORAL

NAPROSYN

AB + ATNAHS PHARMA US 250MG

N017581 002

AB + 375MG

N017581 003

AB +! 500MG

N017581 004 Apr 15, 1982

NAPROXEN

AB AMNEAL PHARMS NY 250MG

A075927 001 Dec 18, 2001

AB 375MG

A075927 002 Dec 18, 2001

AB 500MG

A075927 003 Dec 18, 2001

AB AUROBINDO PHARMA LTD 250MG

A200429 001 Nov 08, 2011

AB 375MG

A200429 002 Nov 08, 2011

AB 500MG

A200429 003 Nov 08, 2011

AB GLENMARK GENERICS 250MG

A078250 001 Mar 28, 2007

AB 375MG

A078250 002 Mar 28, 2007

AB 500MG

A078250 003 Mar 28, 2007

AB INVAGEN PHARMS 250MG

A091305 001 Aug 24, 2011

AB 375MG

A091305 002 Aug 24, 2011

AB 500MG

A091305 003 Aug 24, 2011

AB MARKSANS PHARMA 250MG

A091416 001 Feb 14, 2011

AB 375MG

A091416 002 Feb 14, 2011

AB 500MG

A091416 003 Feb 14, 2011

AB MYLAN 250MG

A074121 001 Dec 21, 1993

AB 375MG

A074121 002 Dec 21, 1993

AB 500MG

A074121 003 Dec 21, 1993

AB PERRIGO R AND D 250MG

A077339 001 Apr 27, 2005

AB 375MG

A077339 002 Apr 27, 2005

AB 500MG

A077339 003 Apr 27, 2005

AB TEVA 250MG

A074201 001 Dec 21, 1993

AB 375MG

A074201 002 Dec 21, 1993

AB 500MG

A074201 003 Dec 21, 1993

AB ZYDUS PHARMS USA 250MG

A078620 001 Jun 07, 2007

AB 375MG

A078620 002 Jun 07, 2007

AB 500MG

A078620 003 Jun 07, 2007

TABLET, DELAYED RELEASE; ORAL

EC-NAPROSYN

AB +! ATNAHS PHARMA US 375MG

N020067 002 Oct 14, 1994

AB +! 500MG

N020067 003 Oct 14, 1994

NAPROXEN

AB INVAGEN PHARMS 375MG

A091432 001 Sep 19, 2011

AB 500MG

A091432 002 Sep 19, 2011

AB PLIVA 375MG

A075337 001 May 26, 1999

AB 500MG

A075337 002 May 26, 1999

AB TEVA 375MG

A075227 001 Jun 30, 1998

AB 500MG

A075227 002 Jun 30, 1998

NAPROXEN SODIUM

TABLET; ORAL

ANAPROX

AB + ATNAHS PHARMA US EQ 250MG BASE

N018164 001

ANAPROX DS

AB +! ATNAHS PHARMA US EQ 500MG BASE

N018164 003 Sep 30, 1987

NAPROXEN SODIUM

AB AMNEAL PHARMS NY EQ 250MG BASE

A078432 001 Apr 25, 2007

AB EQ 500MG BASE

A078432 002 Apr 25, 2007

AB AUROBINDO PHARMA LTD EQ 250MG BASE

A200629 001 Oct 31, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-305 (of 436)

NAPROXEN SODIUM

TABLET;ORAL

NAPROXEN SODIUM

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 500MG BASE</u>	<u>A200629 002</u>	Oct 31, 2011
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A078486 001</u>	Jul 26, 2007
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A078486 002</u>	Jul 26, 2007
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 250MG BASE</u>	<u>A078314 001</u>	Apr 27, 2007
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A078314 002</u>	Apr 27, 2007
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>A074198 001</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A074198 002</u>	Dec 21, 1993

TABLET, EXTENDED RELEASE;ORAL

NAPRELAN

<u>AB</u>	+	ALVOGEN MALTA	<u>EQ 375MG BASE</u>	<u>N020353 001</u>	Jan 05, 1996
<u>AB</u>	+		<u>EQ 500MG BASE</u>	<u>N020353 002</u>	Jan 05, 1996
<u>AB</u>	+!		<u>EQ 750MG BASE</u>	<u>N020353 003</u>	Jan 05, 1996

NAPROXEN SODIUM

<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 375MG BASE</u>	<u>A075416 002</u>	Apr 23, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>A075416 001</u>	Aug 27, 2002
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>A075416 003</u>	Aug 11, 2016

NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE

TABLET;ORAL

TREXIMET

+	PERNIX IRELAND LTD	60MG;EQ 10MG BASE	N021926 002	May 14, 2015
+!		500MG;EQ 85MG BASE	N021926 001	Apr 15, 2008

NARatriptan Hydrochloride

TABLET;ORAL

AMERGE

<u>AB</u>	+	GLAXOSMITHKLINE LLC	<u>EQ 1MG BASE</u>	<u>N020763 002</u>	Feb 10, 1998
<u>AB</u>	+!		<u>EQ 2.5MG BASE</u>	<u>N020763 001</u>	Feb 10, 1998

NARatriptan

<u>AB</u>	APOTEX CORP	<u>EQ 1MG BASE</u>	<u>A091373 001</u>	Apr 22, 2011
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A091373 002</u>	Apr 22, 2011
<u>AB</u>	HERITAGE PHARMS INC	<u>EQ 1MG BASE</u>	<u>A200502 001</u>	Feb 28, 2011
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A200502 002</u>	Feb 28, 2011
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 1MG BASE</u>	<u>A202431 001</u>	May 31, 2012
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A202431 002</u>	May 31, 2012
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 1MG BASE</u>	<u>A091441 001</u>	Apr 30, 2012
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A091441 002</u>	Apr 30, 2012
<u>AB</u>	PADDOCK LLC	<u>EQ 1MG BASE</u>	<u>A091326 001</u>	Jul 08, 2010
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A091326 002</u>	Jul 08, 2010
<u>AB</u>	SANDOZ	<u>EQ 1MG BASE</u>	<u>A090288 001</u>	Jul 07, 2010
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A090288 002</u>	Jul 07, 2010
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 2.5MG BASE</u>	<u>A091552 001</u>	Feb 14, 2011
<u>AB</u>	TEVA PHARMS	<u>EQ 1MG BASE</u>	<u>A078751 001</u>	Jul 07, 2010
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A078751 002</u>	Jul 07, 2010
<u>AB</u>	WEST-WARD PHARMS INT	<u>EQ 1MG BASE</u>	<u>A090381 001</u>	Jul 07, 2010
<u>AB</u>		<u>EQ 2.5MG BASE</u>	<u>A090381 002</u>	Jul 07, 2010

NATAMYCIN

SUSPENSION;OPHTHALMIC

NATACYN

+!	NOVARTIS PHARMS CORP	5%	N050514 001
----	----------------------	----	-------------

NATEGLINIDE

TABLET;ORAL

NATEGLINIDE

<u>AB</u>	ALVOGEN MALTA	<u>60MG</u>	<u>A205055 001</u>	Dec 11, 2015
<u>AB</u>		<u>120MG</u>	<u>A205055 002</u>	Dec 11, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>60MG</u>	<u>A077461 001</u>	Sep 09, 2009
<u>AB</u>		<u>120MG</u>	<u>A077461 002</u>	Sep 09, 2009
<u>AB</u>	PAR PHARM	<u>60MG</u>	<u>A077463 001</u>	Sep 09, 2009
<u>AB</u>		<u>120MG</u>	<u>A077463 002</u>	Sep 09, 2009
<u>AB</u>	WATSON LABS	<u>60MG</u>	<u>A077462 001</u>	Mar 30, 2011
<u>AB</u>		<u>120MG</u>	<u>A077462 002</u>	Mar 30, 2011
<u>AB</u>	ZYDUS PHARMS USA INC	<u>60MG</u>	<u>A205248 001</u>	Jul 06, 2016
<u>AB</u>		<u>120MG</u>	<u>A205248 002</u>	Jul 06, 2016

STARLIX

<u>AB</u>	+	NOVARTIS	<u>60MG</u>	<u>N021204 001</u>	Dec 22, 2000
<u>AB</u>	+!		<u>120MG</u>	<u>N021204 002</u>	Dec 22, 2000

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-306 (of 436)

NEBIVOLOL HYDROCHLORIDE

TABLET;ORAL

BYSTOLIC

+ FOREST LABS	EQ 2.5MG BASE	N021742 002 Dec 17, 2007
+	EQ 5MG BASE	N021742 003 Dec 17, 2007
+	EQ 10MG BASE	N021742 004 Dec 17, 2007
+!	EQ 20MG BASE	N021742 005 Oct 08, 2008

NEBIVOLOL HYDROCHLORIDE; VALSARTAN

TABLET;ORAL

BYVALSON

+! FOREST LABS LLC	EQ 5MG BASE;80MG	N206302 001 Jun 03, 2016
--------------------	------------------	--------------------------

NEDOCROMIL SODIUM

SOLUTION/DROPS;OPHTHALMIC

ALOCRIL

<u>AT +! ALLERGAN</u>	<u>2%</u>	<u>N021009 001</u> Dec 08, 1999
-----------------------	-----------	---------------------------------

NEDOCROMIL SODIUM

<u>AT AKORN</u>	<u>2%</u>	<u>A090638 001</u> Aug 22, 2012
-----------------	-----------	---------------------------------

NEFAZODONE HYDROCHLORIDE

TABLET;ORAL

NEFAZODONE HYDROCHLORIDE

TEVA	50MG	A076037 001 Sep 16, 2003
	100MG	A076037 002 Sep 16, 2003
	150MG	A076037 003 Sep 16, 2003
	200MG	A076037 004 Sep 16, 2003
!	250MG	A076037 005 Sep 16, 2003

NELARABINE

INJECTABLE;IV (INFUSION)

ARRANON

+! NOVARTIS PHARMS CORP	250MG/50ML (5MG/ML)	N021877 001 Oct 28, 2005
-------------------------	---------------------	--------------------------

NELFINAVIR MESYLATE

TABLET;ORAL

VIRACEPT

+! AGOURON PHARMS	EQ 250MG BASE	N020779 001 Mar 14, 1997
+	EQ 625MG BASE	N021503 001 Apr 30, 2003

NEOMYCIN SULFATE

TABLET;ORAL

NEOMYCIN SULFATE

<u>AA BRECKENRIDGE PHARM</u>	<u>500MG</u>	<u>A065468 001</u> Mar 29, 2010
<u>AA LANNETT HOLDINGS INC</u>	<u>500MG</u>	<u>A204435 001</u> Jun 10, 2016
<u>AA ! TEVA</u>	<u>500MG</u>	<u>A060304 001</u>
<u>AA X GEN PHARMS</u>	<u>500MG</u>	<u>A065220 001</u> Jul 28, 2006

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION;IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

<u>AT WATSON LABS</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A062664 001</u> Apr 08, 1986
<u>AT X GEN PHARMS</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A065106 001</u> Jan 31, 2006
<u>AT</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A065108 001</u> Jan 31, 2006
<u>NEOSPORIN G.U. IRRIGANT</u>		
<u>AT ! MONARCH PHARMS</u>	<u>EQ 40MG BASE/ML;200,000 UNITS/ML</u>	<u>A060707 001</u>

NEOSTIGMINE METHYLSULFATE

SOLUTION;INTRAVENOUS

BLOXIVERZ

<u>AP +! ECLAT PHARMS LLC</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>N204078 001</u> May 31, 2013
<u>AP +!</u>	<u>10MG/10ML (1MG/ML)</u>	<u>N204078 002</u> May 31, 2013

NEOSTIGMINE METHYLSULFATE

<u>AP AMPHASTAR PHARMS INC</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A209933 001</u> Sep 25, 2017
<u>AP EUROHLTH INTL SARL</u>	<u>10MG/10ML (1MG/ML)</u>	<u>A209933 002</u> Sep 25, 2017
<u>AP</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A207042 001</u> Dec 28, 2015
<u>AP</u>	<u>10MG/10ML (1MG/ML)</u>	<u>A207042 002</u> Dec 28, 2015
<u>AP PAR STERILE PRODUCTS</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>A208405 001</u> Apr 26, 2017
<u>AP</u>	<u>10MG/10ML (1MG/ML)</u>	<u>A208405 002</u> Apr 26, 2017
<u>FRESENIUS KABI USA</u>	<u>5MG/10ML (0.5MG/ML)</u>	<u>N203629 001</u> Jan 08, 2015
	<u>10MG/10ML (1MG/ML)</u>	<u>N203629 002</u> Jan 08, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-307 (of 436)

NEPAFENAC

SUSPENSION/DROPS;OPHTHALMIC

ILEVRO

+! NOVARTIS PHARMS CORP

NEVANAC

+! NOVARTIS PHARMS CORP

N203491 001 Oct 16, 2012

N021862 001 Aug 19, 2005

NERATINIB MALEATE

TABLET;ORAL

NERLYNX

+! PUMA BIOTECH EQ 40MG BASE

N208051 001 Jul 17, 2017

NESIRITIDE RECOMBINANT

FOR SOLUTION;INTRAVENOUS

NATRECOR

+! SCIOS LLC 1.5MG/VIAL

N020920 001 Aug 10, 2001

NETARSUDIL DIMESYLATE

SOLUTION/DROPS;OPHTHALMIC

RHOPRESSA

+! AERIE PHARMS INC EQ 0.02% BASE

N208254 001 Dec 18, 2017

NETUPITANT; PALONOSETRON HYDROCHLORIDE

CAPSULE;ORAL

AKYNZEO

+! HELSINN HLTHCARE 300MG;EQ 0.5MG BASE

N205718 001 Oct 10, 2014

NEVIRAPINE

SUSPENSION;ORAL

NEVIRAPINE

AA AUROBINDO

50MG/5ML

A077702 001 May 22, 2012

AA CIPLA LTD

50MG/5ML

A207684 001 Aug 03, 2017

VIRAMUNE

AA +! BOEHRINGER

INGELHEIM

50MG/5ML

N020933 001 Sep 11, 1998

TABLET;ORAL

NEVIRAPINE

AB APOTEX INC

200MG

A203021 001 May 22, 2012

AB AUROBINDO

200MG

A077521 001 May 22, 2012

AB CIPLA

200MG

A077956 001 May 22, 2012

AB HETERO LABS LTD III

200MG

A078584 001 May 22, 2012

AB MICRO LABS LTD

200MG

A203080 001 May 22, 2012

AB MYLAN LABS

200MG

A078864 001 May 22, 2012

AB MYLAN PHARMS INC

200MG

A202523 001 May 22, 2012

AB PRINSTON INC

200MG

A078644 001 May 22, 2012

AB STRIDES PHARMA

200MG

A078195 001 May 22, 2012

AB TECH ORGANIZED

200MG

A203176 001 May 22, 2012

VIRAMUNE

AB +! BOEHRINGER

INGELHEIM

200MG

N020636 001 Jun 21, 1996

TABLET, EXTENDED RELEASE;ORAL

NEVIRAPINE

AB ALVOGEN MALTA

100MG

A204621 002 Nov 09, 2015

AB APOTEX INC

400MG

A204621 001 Jul 10, 2015

AB AUROBINDO PHARMA LTD

100MG

A205258 001 Apr 03, 2014

AB

400MG

A208616 001 Nov 23, 2016

AB

400MG

A207698 001 Feb 28, 2017

AB CIPLA LTD

400MG

A206448 001 Oct 15, 2015

AB MACLEODS PHARMS LTD

400MG

A206879 001 Oct 06, 2017

AB MYLAN PHARMS INC

100MG

A206271 001 Nov 09, 2015

AB

400MG

A205651 001 Oct 27, 2014

AB SANDOZ INC

400MG

A203411 001 Apr 03, 2014

AB TECH ORGANIZED

100MG

A207467 001 Jul 31, 2017

AB

400MG

A207467 002 Jul 31, 2017

VIRAMUNE XR

AB +! BOEHRINGER

INGELHEIM

100MG

N201152 002 Nov 08, 2012

AB +!

400MG

N201152 001 Mar 25, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-308 (of 436)

NIACIN

TABLET;ORAL

<b>AA</b>	WOCKHARDT	<u>500MG</u>	<b>A081134 001</b>	Apr 28, 1992
	<b>NIACOR</b>			
<b>AA</b>	AVONDALE PHARMS	<u>500MG</u>	<b>A040378 001</b>	May 03, 2000
	TABLET, EXTENDED RELEASE;ORAL			
	<b>NIACIN</b>			
<b>AB</b>	AMNEAL PHARMS	<u>500MG</u>	<b>A203578 001</b>	Jul 24, 2015
<b>AB</b>		<u>750MG</u>	<b>A204178 001</b>	Dec 11, 2015
<b>AB</b>		<u>1GM</u>	<b>A203578 002</b>	Jul 24, 2015
<b>AB</b>	BARR	<u>500MG</u>	<b>A076378 001</b>	Apr 26, 2005
<b>AB</b>		<u>750MG</u>	<b>A076378 002</b>	Apr 26, 2005
<b>AB</b>		<u>1GM</u>	<b>A076250 001</b>	Apr 14, 2005
<b>AB</b>	LANNETT CO INC	<u>500MG</u>	<b>A203899 001</b>	Jun 16, 2017
<b>AB</b>		<u>1GM</u>	<b>A203899 002</b>	Jun 16, 2017
<b>AB</b>	LUPIN LTD	<u>500MG</u>	<b>A090860 001</b>	Mar 20, 2014
<b>AB</b>		<u>750MG</u>	<b>A090892 001</b>	Mar 20, 2014
<b>AB</b>		<u>1GM</u>	<b>A090446 001</b>	Mar 20, 2014
<b>AB</b>	SUN PHARMA GLOBAL	<u>500MG</u>	<b>A200484 001</b>	Apr 23, 2014
<b>AB</b>		<u>750MG</u>	<b>A201273 001</b>	Apr 23, 2014
<b>AB</b>		<u>1GM</u>	<b>A200484 002</b>	Apr 23, 2014
	<b>NIASPAN</b>			
<b>AB</b>	+ ABBVIE	<u>500MG</u>	<b>N020381 002</b>	Jul 28, 1997
<b>AB</b>	+!	<u>750MG</u>	<b>N020381 003</b>	Jul 28, 1997
<b>AB</b>	+!	<u>1GM</u>	<b>N020381 004</b>	Jul 28, 1997

NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

	<b>NICARDIPINE HYDROCHLORIDE</b>			
<b>AB</b>	ANI PHARMS INC	<u>20MG</u>	<b>A074439 001</b>	Dec 10, 1996
<b>AB</b>		<u>20MG</u>	<b>A074540 001</b>	Oct 28, 1996
<b>AB</b>		<u>30MG</u>	<b>A074439 002</b>	Dec 10, 1996
<b>AB</b>		<u>30MG</u>	<b>A074540 002</b>	Oct 28, 1996
<b>AB</b>	EPIC PHARMA	<u>20MG</u>	<b>A074928 001</b>	Mar 19, 1998
<b>AB</b>		<u>30MG</u>	<b>A074928 002</b>	Mar 19, 1998
<b>AB</b>	MYLAN	<u>20MG</u>	<b>A074642 001</b>	Jul 18, 1996
<b>AB</b>	!	<u>30MG</u>	<b>A074642 002</b>	Jul 18, 1996

INJECTABLE;INJECTION

	<b>CARDENE</b>			
<b>AP</b>	+! CHIESI USA INC	<u>25MG/10ML (2.5MG/ML)</u>	<b>N019734 001</b>	Jan 30, 1992
	<b>NICARDIPINE HYDROCHLORIDE</b>			
<b>AP</b>	EXELA PHARMA SCIENCE	<u>25MG/10ML (2.5MG/ML)</u>	<b>N022276 001</b>	Jul 24, 2008
<b>AP</b>	LUITPOLD PHARMS INC	<u>25MG/10ML (2.5MG/ML)</u>	<b>A090534 001</b>	Nov 17, 2009
<b>AP</b>	MYLAN INSTITUTIONAL	<u>25MG/10ML (2.5MG/ML)</u>	<b>A090664 001</b>	Nov 17, 2009
<b>AP</b>	NAVINTA LLC	<u>25MG/10ML (2.5MG/ML)</u>	<b>A090125 001</b>	Nov 17, 2009
<b>AP</b>	SUN PHARMA GLOBAL	<u>25MG/10ML (2.5MG/ML)</u>	<b>N078405 001</b>	Nov 17, 2009
<b>AP</b>	WEST-WARD PHARMS INT	<u>25MG/10ML (2.5MG/ML)</u>	<b>A078714 001</b>	Dec 28, 2009
<b>AP</b>	WOCKHARDT	<u>25MG/10ML (2.5MG/ML)</u>	<b>A090671 001</b>	Nov 17, 2009

INJECTABLE;INTRAVENOUS

	CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER			
+!	CHIESI USA INC	40MG/200ML (0.2MG/ML)	N019734 004	Nov 07, 2008
	CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER			
+!	CHIESI USA INC	20MG/200ML (0.1MG/ML)	N019734 003	Jul 31, 2008
	CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER			
+!	CHIESI USA INC	20MG/200ML (0.1MG/ML)	N019734 002	Jul 31, 2008
	CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER			
+!	CHIESI USA INC	40MG/200ML (0.2MG/ML)	N019734 005	Nov 07, 2008
	NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE			
	EXELA PHARMA SCIENCE	20MG/200ML (0.1MG/ML)	N022276 002	Apr 07, 2016
		40MG/200ML (0.2MG/ML)	N022276 003	Apr 07, 2016

NICOTINE

INHALANT;ORAL

	<b>NICOTROL</b>			
+!	PHARMACIA AND UPJOHN	4MG/CARTRIDGE	N020714 001	May 02, 1997
	SPRAY, METERED;NASAL			
NICOTROL				
+!	Pfizer Inc	0.5MG/SPRAY	N020385 001	Mar 22, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-309 (of 436)

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072579</u> <u>001</u>	Jan 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A072556</u> <u>001</u>	Sep 20, 1990
<u>AB</u>	HERITAGE PHARMA	<u>10MG</u>	<u>A202644</u> <u>001</u>	Apr 25, 2013
<u>AB</u>		<u>20MG</u>	<u>A202644</u> <u>002</u>	Apr 25, 2013
<u>AB</u>	INTERGEL PHARM	<u>10MG</u>	<u>A072781</u> <u>001</u>	Jul 30, 1993
<u>AB</u>	VALIDUS PHARMS	<u>10MG</u>	<u>A073250</u> <u>001</u>	Oct 08, 1991
<u>AB</u>		<u>20MG</u>	<u>A074045</u> <u>001</u>	Apr 30, 1992

PROCARDIA

<u>AB</u> +!	PFIZER	<u>10MG</u>	<u>N018482</u> <u>001</u>
--------------	--------	-------------	---------------------------

TABLET, EXTENDED RELEASE; ORAL

ADALAT CC

<u>AB1</u> +	ALVOGEN	<u>30MG</u>	<u>N020198</u> <u>001</u>	Apr 21, 1993
<u>AB1</u> +!		<u>60MG</u>	<u>N020198</u> <u>002</u>	Apr 21, 1993
<u>AB1</u> +!		<u>90MG</u>	<u>N020198</u> <u>003</u>	Apr 21, 1993

AFFECTAB CR

<u>AB1</u>	WATSON LABS	<u>60MG</u>	<u>A075659</u> <u>001</u>	Oct 26, 2001
<u>AB1</u>	WATSON LABS TEVA	<u>30MG</u>	<u>A075128</u> <u>001</u>	Mar 10, 2000

NIFEDIPINE

<u>AB1</u>	MYLAN	<u>30MG</u>	<u>A201071</u> <u>001</u>	Dec 03, 2010
<u>AB1</u>		<u>60MG</u>	<u>A201071</u> <u>002</u>	Dec 03, 2010
<u>AB1</u>		<u>90MG</u>	<u>A201071</u> <u>003</u>	Dec 03, 2010
<u>AB1</u>	NOVAST LABS LTD	<u>30MG</u>	<u>A202987</u> <u>001</u>	Aug 25, 2016
<u>AB1</u>		<u>60MG</u>	<u>A202987</u> <u>002</u>	Aug 25, 2016
<u>AB1</u>		<u>90MG</u>	<u>A202987</u> <u>003</u>	Aug 25, 2016
<u>AB1</u>	PAR PHARM	<u>30MG</u>	<u>A077899</u> <u>001</u>	Dec 13, 2006
<u>AB1</u>		<u>60MG</u>	<u>A077899</u> <u>002</u>	Dec 13, 2006
<u>AB1</u>		<u>90MG</u>	<u>A077899</u> <u>003</u>	May 25, 2012
<u>AB1</u>	VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075269</u> <u>001</u>	Dec 04, 2000
<u>AB1</u>		<u>60MG</u>	<u>A075269</u> <u>002</u>	Dec 04, 2000
<u>AB1</u>		<u>90MG</u>	<u>A076070</u> <u>001</u>	Aug 16, 2002
<u>AB2</u>	MYLAN	<u>30MG</u>	<u>A090649</u> <u>001</u>	Jun 21, 2010
<u>AB2</u>		<u>60MG</u>	<u>A090649</u> <u>002</u>	Jun 21, 2010
<u>AB2</u>		<u>90MG</u>	<u>A090649</u> <u>003</u>	Jun 21, 2010
<u>AB2</u>	OSMOTICA PHARM US	<u>30MG</u>	<u>A077127</u> <u>001</u>	Nov 21, 2005
<u>AB2</u>		<u>60MG</u>	<u>A077127</u> <u>002</u>	Nov 21, 2005
<u>AB2</u>		<u>90MG</u>	<u>A077410</u> <u>001</u>	Oct 03, 2007
<u>AB2</u>	TWI PHARMS INC	<u>30MG</u>	<u>A203126</u> <u>001</u>	Apr 03, 2014
<u>AB2</u>		<u>60MG</u>	<u>A203126</u> <u>002</u>	Apr 03, 2014
<u>AB2</u>		<u>90MG</u>	<u>A203126</u> <u>003</u>	Apr 03, 2014
<u>AB2</u>	VALEANT PHARMS NORTH	<u>30MG</u>	<u>A075289</u> <u>002</u>	Feb 06, 2001
<u>AB2</u>		<u>60MG</u>	<u>A075289</u> <u>001</u>	Sep 27, 2000
<u>AB2</u>	ZYDUS PHARMS USA INC	<u>30MG</u>	<u>A210012</u> <u>001</u>	Dec 19, 2017
<u>AB2</u>		<u>60MG</u>	<u>A210012</u> <u>002</u>	Dec 19, 2017
<u>AB2</u>		<u>90MG</u>	<u>A210012</u> <u>003</u>	Dec 19, 2017

PROCARDIA XL

<u>AB2</u> +	PFIZER	<u>30MG</u>	<u>N019684</u> <u>001</u>	Sep 06, 1989
<u>AB2</u> +		<u>60MG</u>	<u>N019684</u> <u>002</u>	Sep 06, 1989
<u>AB2</u> +!		<u>90MG</u>	<u>N019684</u> <u>003</u>	Sep 06, 1989

NILOTINIB HYDROCHLORIDE MONOHYDRATE

CAPSULE; ORAL

TASIGNA

+ NOVARTIS		EQ 150MG BASE	N022068	002	Jun 17, 2010
+!		EQ 200MG BASE	N022068	001	Oct 29, 2007

NILUTAMIDE

TABLET; ORAL

NILANDRON

<u>AB</u> +!	CONCORDIA PHARMS INC	<u>150MG</u>	<u>N020169</u> <u>002</u>	Apr 30, 1999
--------------	----------------------	--------------	---------------------------	--------------

NILUTAMIDE

<u>AB</u>	ANI PHARMS INC	<u>150MG</u>	<u>A207631</u> <u>001</u>	Jul 15, 2016
-----------	----------------	--------------	---------------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-310 (of 436)

NIMODIPINE

CAPSULE;ORAL

NIMODIPINE

<u>AB</u>	!	BIONPHARMA INC	<u>30MG</u>	<u>A076740 001</u>	Jan 17, 2008
<u>AB</u>		HERITAGE PHARMS INC	<u>30MG</u>	<u>A077811 001</u>	May 02, 2007
<u>AB</u>		SOFGEN PHARMS	<u>30MG</u>	<u>A201832 001</u>	Jul 24, 2015
<u>AB</u>		SUN PHARM INDs INC	<u>30MG</u>	<u>A077067 001</u>	Apr 17, 2007
<u>AB</u>		THEPHARMANETWORK LLC	<u>30MG</u>	<u>A090103 001</u>	Apr 07, 2014
		SOLUTION;ORAL			
		NYMALIZE			
+!		ARBOR PHARMS LLC	60MG/20ML	N203340 001	May 10, 2013

NINTEDANIB ESYLADE

CAPSULE;ORAL

OFEV

+	BOEHRINGER INGELHEIM	EQ 100MG BASE	N205832 001	Oct 15, 2014
+!		EQ 150MG BASE	N205832 002	Oct 15, 2014

NIRAPARIB TOSYLATE

CAPSULE;ORAL

ZEJULA

+!	TESARO INC	EQ 100MG BASE	N208447 001	Mar 27, 2017
----	------------	---------------	-------------	--------------

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

NISOLDIPINE

<u>AB</u>		MYLAN	<u>8 . 5MG</u>	<u>A091001 001</u>	Jan 26, 2011
<u>AB</u>			<u>17MG</u>	<u>A091001 002</u>	Jan 26, 2011
<u>AB</u>			<u>34MG</u>	<u>A091001 004</u>	Jan 26, 2011
		<u>SULAR</u>			
<u>AB</u>	+	COVIS PHARMA BV	<u>8 . 5MG</u>	<u>N020356 008</u>	Jan 02, 2008
<u>AB</u>	+		<u>17MG</u>	<u>N020356 007</u>	Jan 02, 2008
<u>AB</u>	+		<u>34MG</u>	<u>N020356 005</u>	Jan 02, 2008
		NISOLDIPINE			
		MYLAN	20MG	A079051 001	Jul 25, 2008
			25.5MG	A091001 003	Jan 26, 2011
	!		30MG	A079051 002	Jul 25, 2008
	!		40MG	A079051 003	Jul 25, 2008

NITAZOXANIDE

FOR SUSPENSION;ORAL

ALINIA

+!	ROMARK	100MG/5ML	N021498 001	Nov 22, 2002
----	--------	-----------	-------------	--------------

TABLET;ORAL

ALINIA

+!	ROMARK	500MG	N021497 001	Jul 21, 2004
----	--------	-------	-------------	--------------

NITISINONE

CAPSULE;ORAL

ORFADIN

+	SWEDISH ORPHAN	2MG	N021232 001	Jan 18, 2002
+		5MG	N021232 002	Jan 18, 2002
+		10MG	N021232 003	Jan 18, 2002
+		20MG	N021232 004	Jun 13, 2016

SUSPENSION;ORAL

ORFADIN

+!	SWEDISH ORPHAN	4MG/ML	N206356 001	Apr 22, 2016
----	----------------	--------	-------------	--------------

TABLET;ORAL

NITYR

+	CYCLE PHARMS LTD	2MG	N209449 001	Jul 26, 2017
+		5MG	N209449 002	Jul 26, 2017
+		10MG	N209449 003	Jul 26, 2017

NITRIC OXIDE

GAS;INHALATION

INOMAX

+!	MALLINCKRODT HOSP	800PPM	N020845 003	Dec 23, 1999
----	-------------------	--------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-311 (of 436)

NITROFURANTOIN

SUSPENSION; ORAL

FURADANTIN

<u>AB</u>	+!	CASPER PHARMA LLC	<u>25MG/5ML</u>	<u>N009175 001</u>
<u>AB</u>		<u>NITROFURANTOIN</u>		
<u>AB</u>		ACTAVIS MID ATLANTIC	<u>25MG/5ML</u>	<u>A205180 001</u> May 03, 2016
<u>AB</u>		AMNEAL PHARMS	<u>25MG/5ML</u>	<u>A201679 001</u> May 11, 2011
<u>AB</u>		NOSTRUM LABS INC	<u>25MG/5ML</u>	<u>A201355 001</u> Aug 14, 2013
<u>AB</u>		NOVEL LABS INC	<u>25MG/5ML</u>	<u>A201693 001</u> Sep 08, 2014

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACRODANTIN

<u>AB</u>	+	ALVOGEN MALTA	<u>25MG</u>	<u>N016620 003</u>
<u>AB</u>	+		<u>50MG</u>	<u>N016620 001</u>
<u>AB</u>	+		<u>100MG</u>	<u>N016620 002</u>
<u>AB</u>		<u>NITROFURANTOIN</u>		
<u>AB</u>		ACTAVIS LABS FL INC	<u>25MG</u>	<u>A091095 001</u> Jun 18, 2015
<u>AB</u>			<u>50MG</u>	<u>A091095 002</u> Jun 18, 2015
<u>AB</u>			<u>100MG</u>	<u>A091095 003</u> Jun 18, 2015
<u>AB</u>		IMPAX LABS INC	<u>50MG</u>	<u>A073671 001</u> Jan 28, 1993
<u>AB</u>			<u>100MG</u>	<u>A073652 001</u> Jan 28, 1993
<u>AB</u>		MYLAN	<u>50MG</u>	<u>A074967 001</u> Jul 09, 1997
<u>AB</u>			<u>100MG</u>	<u>A077025 001</u> Aug 18, 2004
<u>AB</u>		SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A201722 001</u> Feb 16, 2016
<u>AB</u>			<u>50MG</u>	<u>A201722 002</u> Feb 16, 2016
<u>AB</u>			<u>100MG</u>	<u>A201722 003</u> Feb 16, 2016
<u>AB</u>		ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A205005 001</u> Dec 12, 2017
<u>AB</u>			<u>100MG</u>	<u>A205005 002</u> Dec 12, 2017

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

MACROBID

<u>AB</u>	+!	ALVOGEN MALTA	<u>75MG;25MG</u>	<u>N020064 001</u> Dec 24, 1991
		<u>NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)</u>		
<u>AB</u>		AMNEAL PHARMS	<u>75MG;25MG</u>	<u>A207372 001</u> May 15, 2017
<u>AB</u>		MYLAN	<u>75MG;25MG</u>	<u>A076648 001</u> Mar 22, 2004
<u>AB</u>		SANDOZ	<u>75MG;25MG</u>	<u>A077066 001</u> Apr 05, 2005
<u>AB</u>		WATSON LABS INC	<u>75MG;25MG</u>	<u>A202250 001</u> Jul 08, 2015

NITROGLYCERIN

AEROSOL, METERED; SUBLINGUAL

NITROMIST

	+!	MIST PHARMS LLC	0.4MG/SPRAY	
		FILM, EXTENDED RELEASE; TRANSDERMAL		

MINITRAN

<u>AB1</u>		MEDICIS	<u>0 . 4MG/H R</u>	<u>A089773 001</u> Aug 30, 1996
<u>AB1</u>		VALEANT PHARMS	<u>0 . 1MG/H R</u>	<u>A089771 001</u> Aug 30, 1996
<u>AB1</u>			<u>0 . 6MG/H R</u>	<u>A089774 001</u> Aug 30, 1996
<u>AB1</u>		VALEANT PHARMS NORTH	<u>0 . 2MG/H R</u>	<u>A089772 001</u> Aug 30, 1996

NITRO-DUR

<u>AB1</u>	+!	USPHARMA	<u>0 . 1MG/H R</u>	<u>N020145 001</u> Apr 04, 1995
<u>AB1</u>	+!		<u>0 . 2MG/H R</u>	<u>N020145 002</u> Apr 04, 1995
<u>AB1</u>	+!		<u>0 . 4MG/H R</u>	<u>N020145 004</u> Apr 04, 1995
<u>AB1</u>	+!		<u>0 . 6MG/H R</u>	<u>N020145 005</u> Apr 04, 1995

NITROGLYCERIN

<u>AB2</u>		HERCON PHARM	<u>0 . 2MG/H R</u>	<u>A089884 001</u> Oct 30, 1998
<u>AB2</u>			<u>0 . 4MG/H R</u>	<u>A089885 001</u> Oct 30, 1998
<u>AB2</u>			<u>0 . 6MG/H R</u>	<u>A089886 001</u> Oct 30, 1998
<u>AB2</u>	!	MYLAN TECHNOLOGIES	<u>0 . 2MG/H R</u>	<u>A074559 003</u> Aug 30, 1996
<u>AB2</u>	!		<u>0 . 4MG/H R</u>	<u>A074559 002</u> Aug 30, 1996
<u>AB2</u>	!		<u>0 . 6MG/H R</u>	<u>A074559 001</u> Aug 30, 1996

NITRO-DUR

	+!	USPHARMA	0.3MG/HR	N020145 003 Apr 04, 1995
	+!		0.8MG/HR	N020145 006 Apr 04, 1995

NITROGLYCERIN

	!	MYLAN TECHNOLOGIES	0.1MG/HR	A074559 004 Feb 06, 1998
		INJECTABLE; INJECTION		

NITROGLYCERIN IN DEXTROSE 5%

<u>AP</u>	+!	BAXTER HLTHCARE	<u>10MG/100ML</u>	<u>N019970 001</u> Dec 29, 1989
<u>AP</u>	+!		<u>20MG/100ML</u>	<u>N019970 002</u> Dec 29, 1989

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-312 (of 436)

**NITROGLYCERIN**

INJECTABLE; INJECTION

**NITROGLYCERIN IN DEXTROSE 5%**

<b>AP</b>	+!	<b><u>40MG/100ML</u></b>	<b><u>N019970 003</u></b>	Dec 29, 1989
<b>AP</b>	HOSPIRA	<b><u>10MG/100ML</u></b>	<b><u>A071846 001</u></b>	Aug 31, 1990
<b>AP</b>		<b><u>20MG/100ML</u></b>	<b><u>A071847 001</u></b>	Aug 31, 1990
<b>AP</b>		<b><u>40MG/100ML</u></b>	<b><u>A071848 001</u></b>	Aug 31, 1990
NITROGLYCERIN				
! LUITPOLD		5MG/ML	A072034 001	May 24, 1988
OINTMENT; INTRA-ANAL				
RECTIV				
+! FOREST LABS INC		0.4%	N021359 001	Jun 21, 2011
OINTMENT; TRANSDERMAL				
NITROGLYCERIN				
! FOUGERA PHARMS INC		2%	A087355 001	Jul 08, 1988
POWDER; SUBLINGUAL				
GONITRO				
+! POHL BOSKAMP		0.4MG/PACKET	N208424 001	Jun 08, 2016
SPRAY, METERED; SUBLINGUAL				
<b><u>NITROGLYCERIN</u></b>				
<b>AB</b>	PERRIGO ISRAEL	<b><u>0.4MG/SPRAY</u></b>	<b><u>A091496 001</u></b>	Sep 20, 2013
<b><u>NITROLINGUAL PUMPS SPRAY</u></b>				
<b>AB</b>	+! POHL BOSKAMP	<b><u>0.4MG/SPRAY</u></b>	<b><u>N018705 002</u></b>	Jan 10, 1997
TABLET; SUBLINGUAL				
<b><u>NITROGLYCERIN</u></b>				
<b>AB</b>	ACTAVIS LABS FL INC	<b><u>0.3MG</u></b>	<b><u>A203693 001</u></b>	Oct 16, 2017
<b>AB</b>		<b><u>0.4MG</u></b>	<b><u>A203693 002</u></b>	Oct 16, 2017
<b>AB</b>		<b><u>0.6MG</u></b>	<b><u>A203693 003</u></b>	Oct 16, 2017
<b>AB</b>	DR REDDYS LABS INC	<b><u>0.3MG</u></b>	<b><u>A208191 001</u></b>	Aug 26, 2016
<b>AB</b>		<b><u>0.4MG</u></b>	<b><u>A208191 002</u></b>	Aug 26, 2016
<b>AB</b>		<b><u>0.6MG</u></b>	<b><u>A208191 003</u></b>	Aug 26, 2016
<b>AB</b>	GLENMARK PHARMS LTD	<b><u>0.3MG</u></b>	<b><u>A206391 001</u></b>	Sep 19, 2017
<b>AB</b>		<b><u>0.4MG</u></b>	<b><u>A206391 002</u></b>	Sep 19, 2017
<b>AB</b>		<b><u>0.6MG</u></b>	<b><u>A206391 003</u></b>	Sep 19, 2017
<b><u>NITROSTAT</u></b>				
<b>AB</b>	+ PFIZER PHARMS	<b><u>0.3MG</u></b>	<b><u>N021134 001</u></b>	May 01, 2000
<b>AB</b>	+	<b><u>0.4MG</u></b>	<b><u>N021134 002</u></b>	May 01, 2000
<b>AB</b>	+!	<b><u>0.6MG</u></b>	<b><u>N021134 003</u></b>	May 01, 2000

**NIZATIDINE**

CAPSULE; ORAL

**NIZATIDINE**

<b>AB</b>	ANI PHARMS INC	<b><u>150MG</u></b>	<b><u>A075668 001</u></b>	Sep 12, 2002
<b>AB</b>		<b><u>300MG</u></b>	<b><u>A075668 002</u></b>	Sep 12, 2002
<b>AB</b>	DR REDDYS LABS LTD	<b><u>150MG</u></b>	<b><u>A077314 001</u></b>	Sep 15, 2005
<b>AB</b>		<b><u>300MG</u></b>	<b><u>A077314 002</u></b>	Sep 15, 2005
<b>AB</b>	GLENMARK GENERICS	<b><u>150MG</u></b>	<b><u>A090618 001</u></b>	Jul 15, 2011
<b>AB</b>		<b><u>300MG</u></b>	<b><u>A090618 002</u></b>	Jul 15, 2011
<b>AB</b>	MYLAN PHARMS INC	<b><u>150MG</u></b>	<b><u>A075806 001</u></b>	Jul 05, 2002
<b>AB</b>	!	<b><u>300MG</u></b>	<b><u>A075806 002</u></b>	Jul 05, 2002
<b>AB</b>	SANDOZ	<b><u>150MG</u></b>	<b><u>A076178 001</u></b>	Jul 05, 2002
<b>AB</b>		<b><u>300MG</u></b>	<b><u>A076178 002</u></b>	Jul 05, 2002
<b>AB</b>	WATSON LABS	<b><u>150MG</u></b>	<b><u>A075616 001</u></b>	Jul 09, 2002
<b>AB</b>		<b><u>300MG</u></b>	<b><u>A075616 002</u></b>	Jul 09, 2002
SOLUTION; ORAL				
NIZATIDINE				
! AMNEAL PHARMS		15MG/ML	A090576 001	Nov 18, 2009

**NOREpinephrine Bitartrate**

INJECTABLE; INJECTION

**LEVOPHED**

<b>AP</b>	+! HOSPIRA	<b><u>EQ 1MG BASE/ML</u></b>	<b><u>N007513 001</u></b>	
<b>AP</b>	<b><u>NOREpinephrine Bitartrate</u></b>	<b><u>EQ 1MG BASE/ML</u></b>	<b><u>A040859 001</u></b>	Mar 27, 2012
<b>AP</b>	BAXTER HLTHCARE CORP	<b><u>EQ 1MG BASE/ML</u></b>	<b><u>A040455 001</u></b>	Mar 03, 2003
<b>AP</b>	TEVA PHARMS USA	<b><u>EQ 1MG BASE/ML</u></b>	<b><u>A040462 001</u></b>	Oct 31, 2003
<b>AP</b>	WEST-WARD PHARMS INT	<b><u>EQ 1MG BASE/ML</u></b>		

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-313 (of 436)

NORETHINDRONE

TABLET;ORAL-28

CAMILA

<u>AB1</u>	MAYNE PHARMA	<u>0 . 35MG</u>	<u>A076177 001</u> Oct 21, 2002
------------	--------------	-----------------	---------------------------------

HEATHER

<u>AB1</u>	GLENMARK GENERICS	<u>0 . 35MG</u>	<u>A090454 001</u> Apr 23, 2010
------------	-------------------	-----------------	---------------------------------

INCASSIA

<u>AB1</u>	AUROBINDO PHARMA LTD	<u>0 . 35MG</u>	<u>A207304 001</u> Sep 23, 2016
------------	----------------------	-----------------	---------------------------------

NOR-QD

<u>AB1</u>	+! APIL	<u>0 . 35MG</u>	<u>N017060 001</u>
------------	---------	-----------------	--------------------

NORETHINDRONE

<u>AB1</u>	ACCORD HLTHCARE	<u>0 . 35MG</u>	<u>A206807 001</u> Dec 13, 2016
------------	-----------------	-----------------	---------------------------------

<u>AB1</u>	AMNEAL PHARMS	<u>0 . 35MG</u>	<u>A202260 001</u> Aug 01, 2013
------------	---------------	-----------------	---------------------------------

<u>AB1</u>	LUPIN LTD	<u>0 . 35MG</u>	<u>A091325 001</u> Sep 19, 2011
------------	-----------	-----------------	---------------------------------

<u>AB1</u>	MYLAN LABS LTD	<u>0 . 35MG</u>	<u>A201483 001</u> Jun 24, 2013
------------	----------------	-----------------	---------------------------------

<u>AB1</u>	NOVAST LABS LTD	<u>0 . 35MG</u>	<u>A202014 001</u> Sep 13, 2013
------------	-----------------	-----------------	---------------------------------

ERRIN

<u>AB2</u>	MAYNE PHARMA	<u>0 . 35MG</u>	<u>A076225 001</u> Oct 21, 2002
------------	--------------	-----------------	---------------------------------

JENCYCLA

<u>AB2</u>	LUPIN LTD	<u>0 . 35MG</u>	<u>A091323 001</u> Mar 28, 2013
------------	-----------	-----------------	---------------------------------

MICRONOR

<u>AB2</u>	+! JANSSEN PHARMS	<u>0 . 35MG</u>	<u>N016954 001</u>
------------	-------------------	-----------------	--------------------

NORETHINDRONE

<u>AB2</u>	GLENMARK GENERICS	<u>0 . 35MG</u>	<u>A091209 001</u> Jul 22, 2010
------------	-------------------	-----------------	---------------------------------

<u>AB2</u>	MYLAN LABS LTD	<u>0 . 35MG</u>	<u>A200980 001</u> Jun 12, 2013
------------	----------------	-----------------	---------------------------------

<u>AB2</u>	NOVAST LABS	<u>0 . 35MG</u>	<u>A200961 001</u> Sep 13, 2013
------------	-------------	-----------------	---------------------------------

NORETHINDRONE ACETATE

TABLET;ORAL

AYGESTIN

<u>AB</u>	+! DURAMED RES	<u>5MG</u>	<u>N018405 001</u> Apr 21, 1982
-----------	----------------	------------	---------------------------------

NORETHINDRONE ACETATE

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A200275 001</u> Jul 30, 2012
-----------	---------------	------------	---------------------------------

<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A204236 001</u> Jan 08, 2016
-----------	----------------------	------------	---------------------------------

<u>AB</u>	BARR	<u>5MG</u>	<u>A075951 001</u> May 25, 2001
-----------	------	------------	---------------------------------

<u>AB</u>	GLENMARK GENERICS	<u>5MG</u>	<u>A091090 001</u> Jul 21, 2010
-----------	-------------------	------------	---------------------------------

<u>AB</u>	MYLAN LABS LTD	<u>5MG</u>	<u>A205278 001</u> Nov 10, 2016
-----------	----------------	------------	---------------------------------

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE;ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MAYNE PHARMA	<u>EQ 10MG BASE</u>	<u>A073553 001</u> Mar 30, 1992
-----------	--------------	---------------------	---------------------------------

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A073554 001</u> Mar 30, 1992
-----------	--	---------------------	---------------------------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A073555 001</u> Mar 30, 1992
-----------	--	---------------------	---------------------------------

<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A073556 001</u> Mar 30, 1992
-----------	--	---------------------	---------------------------------

<u>AB</u>	TARO PHARM	<u>EQ 10MG BASE</u>	<u>A075520 004</u> May 08, 2000
-----------	------------	---------------------	---------------------------------

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A075520 003</u> May 08, 2000
-----------	--	---------------------	---------------------------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A075520 001</u> May 08, 2000
-----------	--	---------------------	---------------------------------

<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A075520 002</u> May 08, 2000
-----------	--	---------------------	---------------------------------

<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A074132 001</u> Mar 27, 1995
-----------	------	---------------------	---------------------------------

<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A074132 002</u> Mar 27, 1995
-----------	--	---------------------	---------------------------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A074132 003</u> Mar 27, 1995
-----------	--	---------------------	---------------------------------

<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A074132 004</u> Mar 27, 1995
-----------	--	---------------------	---------------------------------

PAMELOR

<u>AB</u>	+ SPECGX LLC	<u>EQ 10MG BASE</u>	<u>N018013 001</u>
-----------	--------------	---------------------	--------------------

<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N018013 002</u>
-----------	---	---------------------	--------------------

<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N018013 004</u>
-----------	---	---------------------	--------------------

<u>AB</u>	+!	<u>EQ 75MG BASE</u>	<u>N018013 003</u>
-----------	----	---------------------	--------------------

SOLUTION;ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AA</u>	! PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>A075606 001</u> Aug 28, 2000
-----------	---------------	-------------------------	---------------------------------

<u>AA</u>	TARO	<u>EQ 10MG BASE/5ML</u>	<u>A077965 001</u> Jun 20, 2006
-----------	------	-------------------------	---------------------------------

NUSINERSEN SODIUM

SOLUTION;INTRATHECAL

SPINRAZA

+!	BIOGEN IDEC	12MG/5ML (2.4MG/ML)	N209531 001 Dec 23, 2016
----	-------------	---------------------	--------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-314 (of 436)

NYSTATIN

CREAM;TOPICAL

**NYSTATIN**

<b>AT</b>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<b>A062949 001</b>	Jun 13, 1988
<b>AT</b>	CROWN LABS INC	<u>100,000 UNITS/GM</u>	<b>A207733 001</b>	Sep 26, 2017
<b>AT</b>	FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<b>A062129 001</b>	
<b>AT</b>	G AND W LABS INC	<u>100,000 UNITS/GM</u>	<b>A061966 001</b>	
<b>AT</b>	PERRIGO NEW YORK	<u>100,000 UNITS/GM</u>	<b>A062225 001</b>	
<b>AT</b>	TARO	<u>100,000 UNITS/GM</u>	<b>A064022 001</b>	Jan 29, 1993
<b>AT</b>	VINTAGE	<u>100,000 UNITS/GM</u>	<b>A065315 001</b>	May 31, 2006

OINTMENT;TOPICAL

**NYSTATIN**

<b>AT</b>	ACTAVIS MID ATLANTIC	<u>100,000 UNITS/GM</u>	<b>A062840 001</b>	Nov 13, 1987
<b>AT</b>	FOUGERA PHARMS	<u>100,000 UNITS/GM</u>	<b>A062124 002</b>	Sep 23, 1982
<b>AT</b>	G AND W LABS INC	<u>100,000 UNITS/GM</u>	<b>A209114 001</b>	Oct 06, 2017
<b>AT</b>	PERRIGO NEW YORK	<u>100,000 UNITS/GM</u>	<b>A062472 001</b>	Feb 13, 1984

POWDER;TOPICAL

**NYSTATIN**

<b>AT</b>	GAVIS PHARMS	<u>100,000 UNITS/GM</u>	<b>A065138 001</b>	Jul 23, 2004
<b>AT</b>	LYNE	<u>100,000 UNITS/GM</u>	<b>A208838 001</b>	May 30, 2017
<b>AT</b>	MAYNE PHARMA INC	<u>100,000 UNITS/GM</u>	<b>A065203 001</b>	Jul 15, 2004
<b>AT</b>	NESHER PHARMS	<u>100,000 UNITS/GM</u>	<b>A208581 001</b>	Jun 08, 2017
<b>AT</b>	UPSHER-SMITH LABS	<u>100,000 UNITS/GM</u>	<b>A065183 001</b>	May 03, 2005
<b>AT</b>	X GEN PHARMS	<u>100,000 UNITS/GM</u>	<b>A065175 001</b>	Dec 17, 2004

**NYSTOP**

<b>AT</b>	PADDOCK LLC	<u>100,000 UNITS/GM</u>	<b>A064118 001</b>	Aug 16, 1996
	SUSPENSION;ORAL			

**NYSTATIN**

<b>AA</b>	FOUGERA PHARMS INC	<u>100,000 UNITS/ML</u>	<b>A062517 001</b>	Jun 07, 1984
<b>AA</b>	G AND W LABS INC	<u>100,000 UNITS/ML</u>	<b>A062349 001</b>	Jul 14, 1982
<b>AA</b>	HI TECH PHARMA	<u>100,000 UNITS/ML</u>	<b>A064042 001</b>	Feb 28, 1994
<b>AA</b>	PHARM ASSOC	<u>100,000 UNITS/ML</u>	<b>A203621 001</b>	Jan 07, 2016
<b>AA</b>	TARO PHARM	<u>100,000 UNITS/ML</u>	<b>A062876 001</b>	Feb 29, 1988
<b>AA</b>	VINTAGE PHARMS	<u>100,000 UNITS/ML</u>	<b>A065148 001</b>	Jun 28, 2005
<b>AA</b>	VISTAPHARM	<u>100,000 UNITS/ML</u>	<b>A064142 001</b>	Jun 25, 1998
<b>AA</b>		<u>100,000 UNITS/ML</u>	<b>A065422 001</b>	Mar 07, 2011
<b>AA</b>	WOCKHARDT BIO AG	<u>100,000 UNITS/ML</u>	<b>A062512 001</b>	Oct 29, 1984

TABLET;ORAL

**NYSTATIN**

<b>AA</b>	HERITAGE PHARMS INC	<u>500,000 UNITS</u>	<b>A062474 001</b>	Dec 22, 1983
<b>AA</b>	SUN PHARM INDUSTRIES	<u>500,000 UNITS</u>	<b>A062838 001</b>	Dec 22, 1988
<b>AA</b>	TEVA	<u>500,000 UNITS</u>	<b>A062506 001</b>	Jan 16, 1984

TABLET;VAGINAL

NYSTATIN

!	ODYSSEY PHARMS	100,000 UNITS		
---	----------------	---------------	--	--

A062615 001 Oct 17, 1985

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

**MYKACET**

<b>AT</b>	G AND W LABS INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A062367 001</b>	May 28, 1985
-----------	------------------	------------------------------	--------------------	--------------

**NYSTATIN AND TRIAMCINOLONE ACETONIDE**

<b>AT</b>	CROWN LABS INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A207730 001</b>	Dec 26, 2017
<b>AT</b>	DR REDDYS LABS LTD	<u>100,000 UNITS/GM;0.1%</u>	<b>A208326 001</b>	Oct 26, 2016
<b>AT</b>	FOUGERA PHARMS INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A062599 001</b>	Oct 08, 1985
<b>AT</b>	GLENMARK PHARMS LTD	<u>100,000 UNITS/GM;0.1%</u>	<b>A208136 001</b>	Oct 24, 2016
<b>AT</b>	PERRIGO UK FINCO	<u>100,000 UNITS/GM;0.1%</u>	<b>A208479 001</b>	Aug 14, 2017
<b>AT</b>	TARO	<u>100,000 UNITS/GM;0.1%</u>	<b>A062364 001</b>	Dec 22, 1987

OINTMENT;TOPICAL

**MYKACET**

<b>AT</b>	G AND W LABS INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A062733 001</b>	Mar 06, 1987
-----------	------------------	------------------------------	--------------------	--------------

**NYSTATIN AND TRIAMCINOLONE ACETONIDE**

<b>AT</b>	AKORN	<u>100,000 UNITS/GM;0.1%</u>	<b>A207217 001</b>	Aug 04, 2017
<b>AT</b>	CROWN LABS INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A207731 001</b>	Dec 26, 2017
<b>AT</b>	DR REDDYS LABS LTD	<u>100,000 UNITS/GM;0.1%</u>	<b>A207741 001</b>	Jan 31, 2017
<b>AT</b>	FOUGERA PHARMS INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A062602 001</b>	Oct 09, 1985
<b>AT</b>	GLENMARK PHARMS LTD	<u>100,000 UNITS/GM;0.1%</u>	<b>A208300 001</b>	Jun 23, 2016
<b>AT</b>	PERRIGO UK FINCO	<u>100,000 UNITS/GM;0.1%</u>	<b>A207380 001</b>	Dec 20, 2016
<b>AT</b>	RICONPHARMA LLC	<u>100,000 UNITS/GM;0.1%</u>	<b>A206785 001</b>	Dec 29, 2016
<b>AT</b>	TARO	<u>100,000 UNITS/GM;0.1%</u>	<b>A063305 001</b>	Mar 29, 1993
<b>AT</b>	TELIGENT PHARMA INC	<u>100,000 UNITS/GM;0.1%</u>	<b>A208287 001</b>	Dec 30, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-315 (of 436)

OBETICHOOLIC ACID

TABLET;ORAL

OCALEVA

+ INTERCEPT PHARMS INC	5MG	N207999 001	May 27, 2016
+!	10MG	N207999 002	May 27, 2016

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

**OCTREOTIDE ACETATE**

<b>AP</b>	FRESENIUS KABI USA	<u>EQ 0 .2MG BASE/ML</u>	<b>A077450 001</b>	Feb 10, 2006
<b>AP</b>		<u>EQ 1MG BASE/ML</u>	<b>A077450 002</b>	Feb 10, 2006
<b>AP</b>	SAGENT PHARMS	<u>EQ 0 .2MG BASE/ML</u>	<b>A091041 001</b>	Nov 12, 2013
<b>AP</b>		<u>EQ 1MG BASE/ML</u>	<b>A091041 002</b>	Nov 12, 2013
<b>AP</b>	SUN PHARM IND	<u>EQ 0 .05MG BASE/ML</u>	<b>A077372 001</b>	Aug 14, 2007
<b>AP</b>		<u>EQ 0 .1MG BASE/ML</u>	<b>A077372 002</b>	Aug 14, 2007
<b>AP</b>		<u>EQ 0 .2MG BASE/ML</u>	<b>A077373 001</b>	Aug 14, 2007
<b>AP</b>		<u>EQ 0 .5MG BASE/ML</u>	<b>A077372 003</b>	Aug 14, 2007
<b>AP</b>		<u>EQ 1MG BASE/ML</u>	<b>A077373 002</b>	Aug 14, 2007
<b>AP</b>	TEVA PHARMS USA	<u>EQ 0 .05MG BASE/ML</u>	<b>A075957 001</b>	Oct 03, 2005
<b>AP</b>		<u>EQ 0 .1MG BASE/ML</u>	<b>A075957 002</b>	Oct 03, 2005
<b>AP</b>		<u>EQ 0 .2MG BASE/ML</u>	<b>A075959 001</b>	Nov 21, 2005
<b>AP</b>		<u>EQ 0 .5MG BASE/ML</u>	<b>A075957 003</b>	Oct 03, 2005
<b>AP</b>		<u>EQ 1MG BASE/ML</u>	<b>A075959 002</b>	Nov 21, 2005
<b>AP</b>	! WEST-WARD PHARMS INT	<u>EQ 0 .2MG BASE/ML</u>	<b>A076330 001</b>	Apr 08, 2005
<b>AP</b>		<u>EQ 1MG BASE/ML</u>	<b>A076330 002</b>	Apr 08, 2005
<b>OCTREOTIDE ACETATE (PRESERVATIVE FREE)</b>				
<b>AP</b>	FRESENIUS KABI USA	<u>EQ 0 .05MG BASE/ML</u>	<b>A077457 001</b>	Feb 10, 2006
<b>AP</b>		<u>EQ 0 .1MG BASE/ML</u>	<b>A077457 002</b>	Feb 10, 2006
<b>AP</b>		<u>EQ 0 .5MG BASE/ML</u>	<b>A077457 003</b>	Feb 10, 2006
<b>AP</b>	MYLAN INSTITUTIONAL	<u>EQ 0 .05MG BASE/ML</u>	<b>A079198 001</b>	Feb 10, 2011
<b>AP</b>		<u>EQ 0 .1MG BASE/ML</u>	<b>A079198 002</b>	Feb 10, 2011
<b>AP</b>		<u>EQ 0 .5MG BASE/ML</u>	<b>A079198 003</b>	Feb 10, 2011
<b>AP</b>	SAGENT PHARMS	<u>EQ 0 .05MG BASE/ML</u>	<b>A090834 001</b>	Nov 12, 2013
<b>AP</b>		<u>EQ 0 .1MG BASE/ML</u>	<b>A090834 002</b>	Nov 12, 2013
<b>AP</b>		<u>EQ 0 .5MG BASE/ML</u>	<b>A090834 003</b>	Nov 12, 2013
<b>AP</b>	! WEST-WARD PHARMS INT	<u>EQ 0 .05MG BASE/ML</u>	<b>A076313 001</b>	Mar 28, 2005
<b>AP</b>		<u>EQ 0 .1MG BASE/ML</u>	<b>A076313 003</b>	Mar 28, 2005
<b>AP</b>		<u>EQ 0 .5MG BASE/ML</u>	<b>A076313 002</b>	Mar 28, 2005
<b>SANDOSTATIN</b>				
<b>AP</b>	+! NOVARTIS	<u>EQ 0 .05MG BASE/ML</u>	<b>N019667 001</b>	Oct 21, 1988
<b>AP</b>	+!	<u>EQ 0 .1MG BASE/ML</u>	<b>N019667 002</b>	Oct 21, 1988
<b>AP</b>	+!	<u>EQ 0 .2MG BASE/ML</u>	<b>N019667 004</b>	Jun 12, 1991
<b>AP</b>	+!	<u>EQ 0 .5MG BASE/ML</u>	<b>N019667 003</b>	Oct 21, 1988
<b>AP</b>	+!	<u>EQ 1MG BASE/ML</u>	<b>N019667 005</b>	Jun 12, 1991
SANDOSTATIN LAR				
	+ NOVARTIS	EQ 10MG BASE/VIAL	N021008 001	Nov 25, 1998
	+	EQ 20MG BASE/VIAL	N021008 002	Nov 25, 1998
	!+	EQ 30MG BASE/VIAL	N021008 003	Nov 25, 1998
<b>OFLOXACIN</b>				
SOLUTION/DROPS;OPHTHALMIC				
<b>OCUFLOX</b>				
<b>AT</b>	+! ALLERGAN	<u>0 .3%</u>	<b>N019921 001</b>	Jul 30, 1993
<b>OFLOXACIN</b>				
<b>AT</b>	AKORN	<u>0 .3%</u>	<b>A076407 001</b>	Apr 15, 2008
<b>AT</b>	ALTAIRE PHARMS INC	<u>0 .3%</u>	<b>A202692 001</b>	Apr 29, 2013
<b>AT</b>	ALVOGEN	<u>0 .3%</u>	<b>A076830 001</b>	Aug 31, 2004
<b>AT</b>	BAUSCH AND LOMB	<u>0 .3%</u>	<b>A076622 001</b>	May 14, 2004
<b>AT</b>	FDC LTD	<u>0 .3%</u>	<b>A078559 001</b>	Feb 25, 2009
<b>AT</b>	HI TECH PHARMA	<u>0 .3%</u>	<b>A076615 001</b>	May 14, 2004
<b>AT</b>	SANDOZ INC	<u>0 .3%</u>	<b>A076231 001</b>	May 14, 2004
SOLUTION/DROPS;OTIC				
<b>OFLOXACIN</b>				
<b>AT</b>	ALVOGEN	<u>0 .3%</u>	<b>A090395 001</b>	Aug 11, 2009
<b>AT</b>	APOTEX INC	<u>0 .3%</u>	<b>A076527 001</b>	Sep 28, 2007
<b>AT</b>	! BAUSCH AND LOMB	<u>0 .3%</u>	<b>A076128 001</b>	Mar 17, 2008
<b>AT</b>	HI TECH PHARMA	<u>0 .3%</u>	<b>A076616 001</b>	Mar 17, 2008
<b>AT</b>	SANDOZ INC	<u>0 .3%</u>	<b>A078222 001</b>	Mar 17, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-316 (of 436)

OFLOXACIN

TABLET;ORAL

OFLOXACIN

<u>AB</u>	CADILA PHARMS LTD	<u>200MG</u>	<u>A091656 001</u>	Sep 18, 2014
<u>AB</u>		<u>300MG</u>	<u>A091656 002</u>	Sep 18, 2014
<u>AB</u>		<u>400MG</u>	<u>A091656 003</u>	Sep 18, 2014
<u>AB</u>	DR REDDYS LABS LTD	<u>200MG</u>	<u>A077098 001</u>	Feb 10, 2006
<u>AB</u>		<u>300MG</u>	<u>A077098 002</u>	Feb 10, 2006
<u>AB</u>		<u>400MG</u>	<u>A077098 003</u>	Feb 10, 2006
<u>AB</u>	LARKEN LABS	<u>400MG</u>	<u>A076093 003</u>	Sep 02, 2003
<u>AB</u>	TEVA	<u>200MG</u>	<u>A076182 001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>A076182 002</u>	Sep 02, 2003
<u>AB</u>	!	<u>400MG</u>	<u>A076182 003</u>	Sep 02, 2003

OLANZAPINE

INJECTABLE;INTRAMUSCULAR

OLANZAPINE

<u>AP</u>	LUITPOLD	<u>10MG/VIAL</u>	<u>A201741 001</u>	Mar 20, 2012
<u>AP</u>	SANDOZ INC	<u>10MG/VIAL</u>	<u>A201588 001</u>	Oct 24, 2011
<u>AP</u>	ZYPREXA			

<u>AP</u>	+! LILLY	<u>10MG/VIAL</u>	<u>N021253 001</u>	Mar 29, 2004
-----------	----------	------------------	--------------------	--------------

TABLET;ORAL

OLANZAPINE

<u>AB</u>	ALKEM LABS LTD	<u>2.5MG</u>	<u>A202295 001</u>	Oct 20, 2015
<u>AB</u>		<u>5MG</u>	<u>A202295 002</u>	Oct 20, 2015
<u>AB</u>		<u>7.5MG</u>	<u>A202295 003</u>	Oct 20, 2015
<u>AB</u>		<u>10MG</u>	<u>A202295 004</u>	Oct 20, 2015
<u>AB</u>		<u>15MG</u>	<u>A202295 005</u>	Oct 20, 2015
<u>AB</u>		<u>20MG</u>	<u>A202295 006</u>	Oct 20, 2015
<u>AB</u>	APOTEX INC	<u>2.5MG</u>	<u>A090798 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A090798 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A090798 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A090798 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A090798 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A090798 006</u>	Apr 23, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A202050 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A202050 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A202050 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A202050 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A202050 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A202050 006</u>	Apr 23, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>2.5MG</u>	<u>A076255 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A076255 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A076255 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A076255 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A076133 001</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A076133 002</u>	Oct 24, 2011
<u>AB</u>	HIKMA PHARMS	<u>2.5MG</u>	<u>A204866 001</u>	Jun 16, 2017
<u>AB</u>		<u>5MG</u>	<u>A204866 002</u>	Jun 16, 2017
<u>AB</u>		<u>7.5MG</u>	<u>A204866 003</u>	Jun 16, 2017
<u>AB</u>		<u>10MG</u>	<u>A204866 004</u>	Jun 16, 2017
<u>AB</u>		<u>15MG</u>	<u>A204866 005</u>	Jun 16, 2017
<u>AB</u>		<u>20MG</u>	<u>A204866 006</u>	Jun 16, 2017
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A203333 001</u>	Mar 15, 2016
<u>AB</u>		<u>5MG</u>	<u>A203333 002</u>	Mar 15, 2016
<u>AB</u>		<u>7.5MG</u>	<u>A203333 003</u>	Mar 15, 2016
<u>AB</u>		<u>10MG</u>	<u>A203333 004</u>	Mar 15, 2016
<u>AB</u>		<u>15MG</u>	<u>A203333 005</u>	Mar 15, 2016
<u>AB</u>		<u>20MG</u>	<u>A203333 006</u>	Mar 15, 2016
<u>AB</u>	IVAX PHARMS INC	<u>20MG</u>	<u>A077301 001</u>	Apr 29, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A202862 001</u>	Aug 15, 2014
<u>AB</u>		<u>5MG</u>	<u>A202862 002</u>	Aug 15, 2014
<u>AB</u>		<u>7.5MG</u>	<u>A202862 003</u>	Aug 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A202862 004</u>	Aug 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A202862 005</u>	Aug 15, 2014
<u>AB</u>		<u>20MG</u>	<u>A202862 006</u>	Aug 15, 2014
<u>AB</u>	ORCHID HLTHCARE	<u>2.5MG</u>	<u>A202287 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A202287 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A202287 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A202287 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A202287 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A202287 006</u>	Apr 23, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-317 (of 436)

OLANZAPINE

TABLET;ORAL

OLANZAPINE

<u>AB</u>	QILU PHARM CO LTD	<u>2.5MG</u>	<u>A204319 001</u>	Jan 27, 2016
<u>AB</u>		<u>5MG</u>	<u>A204319 002</u>	Jan 27, 2016
<u>AB</u>		<u>7.5MG</u>	<u>A204319 003</u>	Jan 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A204319 004</u>	Jan 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A204319 005</u>	Jan 27, 2016
<u>AB</u>		<u>20MG</u>	<u>A204319 006</u>	Jan 27, 2016
<u>AB</u>	SUN PHARM INDs	<u>2.5MG</u>	<u>A091038 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A091038 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A091038 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A091038 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A091038 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A091038 006</u>	Apr 23, 2012
<u>AB</u>	TEVA PHARMS	<u>2.5MG</u>	<u>A076000 001</u>	Oct 24, 2011
<u>AB</u>		<u>5MG</u>	<u>A076000 002</u>	Oct 24, 2011
<u>AB</u>		<u>7.5MG</u>	<u>A076000 003</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A076000 004</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076000 005</u>	Oct 24, 2011
<u>AB</u>	TORRENT PHARMS LTD	<u>2.5MG</u>	<u>A091434 001</u>	Apr 23, 2012
<u>AB</u>		<u>5MG</u>	<u>A091434 002</u>	Apr 23, 2012
<u>AB</u>		<u>7.5MG</u>	<u>A091434 003</u>	Apr 23, 2012
<u>AB</u>		<u>10MG</u>	<u>A091434 004</u>	Apr 23, 2012
<u>AB</u>		<u>15MG</u>	<u>A091434 005</u>	Apr 23, 2012
<u>AB</u>		<u>20MG</u>	<u>A091434 006</u>	Apr 23, 2012

ZYPREXA

<u>AB</u>	+ LILLY	<u>2.5MG</u>	<u>N020592 001</u>	Sep 30, 1996
<u>AB</u>	+!	<u>5MG</u>	<u>N020592 002</u>	Sep 30, 1996
<u>AB</u>	+	<u>7.5MG</u>	<u>N020592 003</u>	Sep 30, 1996
<u>AB</u>	+	<u>10MG</u>	<u>N020592 004</u>	Sep 30, 1996
<u>AB</u>	+	<u>15MG</u>	<u>N020592 005</u>	Sep 09, 1997
<u>AB</u>	+	<u>20MG</u>	<u>N020592 006</u>	Sep 09, 1997

TABLET, ORALLY DISINTEGRATING;ORAL

OLANZAPINE

<u>AB</u>	AJANTA PHARMA LTD	<u>5MG</u>	<u>A204320 001</u>	May 30, 2017
<u>AB</u>		<u>10MG</u>	<u>A204320 002</u>	May 30, 2017
<u>AB</u>		<u>15MG</u>	<u>A204320 003</u>	May 30, 2017
<u>AB</u>		<u>20MG</u>	<u>A204320 004</u>	May 30, 2017
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A091265 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A091265 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A091265 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A091265 004</u>	Oct 24, 2011
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A203708 001</u>	May 15, 2014
<u>AB</u>		<u>10MG</u>	<u>A203708 002</u>	May 15, 2014
<u>AB</u>		<u>15MG</u>	<u>A203708 003</u>	May 15, 2014
<u>AB</u>		<u>20MG</u>	<u>A203708 004</u>	May 15, 2014
<u>AB</u>	BARR LABS INC	<u>5MG</u>	<u>A077243 001</u>	Jan 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077243 002</u>	Jan 30, 2012
<u>AB</u>		<u>15MG</u>	<u>A077243 003</u>	Jan 30, 2012
<u>AB</u>		<u>20MG</u>	<u>A077243 004</u>	Jan 30, 2012
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG</u>	<u>A076534 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A076534 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A076534 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A076534 004</u>	Oct 24, 2011
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A203456 001</u>	Mar 16, 2016
<u>AB</u>		<u>10MG</u>	<u>A203456 002</u>	Mar 16, 2016
<u>AB</u>		<u>15MG</u>	<u>A203456 003</u>	Mar 16, 2016
<u>AB</u>		<u>20MG</u>	<u>A203456 004</u>	Mar 16, 2016
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A200221 001</u>	Sep 12, 2012
<u>AB</u>		<u>10MG</u>	<u>A200221 002</u>	Sep 12, 2012
<u>AB</u>		<u>15MG</u>	<u>A200221 003</u>	Sep 12, 2012
<u>AB</u>		<u>20MG</u>	<u>A200221 004</u>	Sep 12, 2012
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A203044 001</u>	Feb 20, 2015
<u>AB</u>		<u>10MG</u>	<u>A203044 002</u>	Feb 20, 2015
<u>AB</u>		<u>15MG</u>	<u>A203044 003</u>	Feb 20, 2015
<u>AB</u>		<u>20MG</u>	<u>A203044 004</u>	Feb 20, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A202285 001</u>	May 12, 2014
<u>AB</u>		<u>10MG</u>	<u>A202285 002</u>	May 12, 2014
<u>AB</u>		<u>15MG</u>	<u>A202285 003</u>	May 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A202285 004</u>	May 12, 2014
<u>AB</u>	ORCHID HLTHCARE	<u>5MG</u>	<u>A202937 001</u>	Mar 02, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-318 (of 436)

OLANZAPINE

TABLET, ORALLY DISINTEGRATING;ORAL  
OLANZAPINE

<u>AB</u>		<u>10MG</u>	<u>A202937 002</u>	Mar 02, 2015
<u>AB</u>		<u>15MG</u>	<u>A202937 003</u>	Mar 02, 2015
<u>AB</u>		<u>20MG</u>	<u>A202937 004</u>	Mar 02, 2015
<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>A078109 001</u>	Oct 24, 2011
<u>AB</u>		<u>10MG</u>	<u>A078109 002</u>	Oct 24, 2011
<u>AB</u>		<u>15MG</u>	<u>A078109 003</u>	Oct 24, 2011
<u>AB</u>		<u>20MG</u>	<u>A078109 004</u>	Oct 24, 2011
<u>AB</u>	SUN PHARM INDS	<u>5MG</u>	<u>A090881 001</u>	Feb 28, 2012
<u>AB</u>		<u>10MG</u>	<u>A090881 002</u>	Feb 28, 2012
<u>AB</u>		<u>15MG</u>	<u>A090881 003</u>	Feb 28, 2012
<u>AB</u>		<u>20MG</u>	<u>A090881 004</u>	Feb 28, 2012
<u>AB</u>	TORRENT PHARMS LLC	<u>5MG</u>	<u>A091415 001</u>	Oct 25, 2011
<u>AB</u>		<u>10MG</u>	<u>A091415 002</u>	Oct 25, 2011
<u>AB</u>		<u>15MG</u>	<u>A091415 003</u>	Oct 25, 2011
<u>AB</u>		<u>20MG</u>	<u>A091415 004</u>	Oct 25, 2011
<b>ZYPREXA ZYDIS</b>				
<u>AB</u>	+! LILLY	<u>5MG</u>	<u>N021086 001</u>	Apr 06, 2000
<u>AB</u>	+	<u>10MG</u>	<u>N021086 002</u>	Apr 06, 2000
<u>AB</u>	+	<u>15MG</u>	<u>N021086 003</u>	Apr 06, 2000
<u>AB</u>	+	<u>20MG</u>	<u>N021086 004</u>	Apr 06, 2000

OLANZAPINE PAMOATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

ZYPREXA RELPREVV

+ ELI LILLY CO	EQ 210MG BASE/VIAL	N022173 001	Dec 11, 2009
+	EQ 300MG BASE/VIAL	N022173 002	Dec 11, 2009
+	EQ 405MG BASE/VIAL	N022173 003	Dec 11, 2009

OLAPARIB

CAPSULE;ORAL

LYNPARZA			
+!	ASTRAZENECA PHARMS	50MG	N206162 001
TABLET;ORAL			Dec 19, 2014
LYNPARZA			
+	ASTRAZENECA PHARMS	100MG	N208558 001
+		150MG	Aug 17, 2017
			N208558 002
			Aug 17, 2017

OLMESARTAN MEDOXOMIL

TABLET;ORAL

BENICAR

<u>AB</u>	+ DAIICHI SANKYO	<u>5MG</u>	<u>N021286 001</u>	Apr 25, 2002
<u>AB</u>	+	<u>20MG</u>	<u>N021286 003</u>	Apr 25, 2002
<u>AB</u>	+!	<u>40MG</u>	<u>N021286 004</u>	Apr 25, 2002

OLMESARTAN MEDOXOMIL

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A207662 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A207662 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A207662 003</u>	Apr 24, 2017
<u>AB</u>	ALEMBIC PHARMS LTD	<u>5MG</u>	<u>A203012 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A203012 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A203012 003</u>	Apr 24, 2017
<u>AB</u>	ALKEM LABS LTD	<u>5MG</u>	<u>A206763 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A206763 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A206763 003</u>	Apr 24, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A204798 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A204798 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A204798 003</u>	Apr 24, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>5MG</u>	<u>A203281 001</u>	May 25, 2017
<u>AB</u>		<u>20MG</u>	<u>A203281 002</u>	May 25, 2017
<u>AB</u>		<u>40MG</u>	<u>A203281 003</u>	May 25, 2017
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A205482 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A205482 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A205482 003</u>	Apr 24, 2017
<u>AB</u>	LUPIN LTD	<u>5MG</u>	<u>A206631 001</u>	Apr 27, 2017
<u>AB</u>		<u>20MG</u>	<u>A206631 002</u>	Apr 27, 2017
<u>AB</u>		<u>40MG</u>	<u>A206631 003</u>	Apr 27, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>5MG</u>	<u>A204814 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A204814 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A204814 003</u>	Apr 24, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A078276 001</u>	Oct 26, 2016
<u>AB</u>		<u>20MG</u>	<u>A078276 002</u>	Oct 26, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-319 (of 436)

OLMESARTAN MEDOXOMIL

TABLET;ORAL

OLMESARTAN MEDOXOMIL

<u>AB</u>		<u>40MG</u>	<u>A078276 003</u>	Oct 26, 2016
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A091079 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A091079 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A091079 003</u>	Apr 24, 2017
<u>AB</u>	TORRENT PHARMS LTD	<u>5MG</u>	<u>A202375 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A202375 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A202375 003</u>	Apr 24, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A205192 001</u>	Apr 24, 2017
<u>AB</u>		<u>20MG</u>	<u>A205192 002</u>	Apr 24, 2017
<u>AB</u>		<u>40MG</u>	<u>A205192 003</u>	Apr 24, 2017

OLODATEROL HYDROCHLORIDE

SPRAY, METERED;INHALATION

STRIVERDI RESPIMAT

+! BOEHRINGER  
INGELHEIM

EQ 0.0025MG BASE/INH

N203108 001 Jul 31, 2014

OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE

SPRAY, METERED;INHALATION

STIOLTO RESPIMAT

+! BOEHRINGER  
INGELHEIM

EQ 0.0025MG BASE/INH;EQ 0.0025MG  
BASE/INH

N206756 001 May 21, 2015

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

OLOPATADINE HYDROCHLORIDE

<u>AT</u>	AKORN	<u>EQ 0.2% BASE</u>	<u>A204723 001</u>	Dec 05, 2017
<u>AT</u>	AKORN INC	<u>EQ 0.1% BASE</u>	<u>A204532 001</u>	Jan 10, 2017
<u>AT</u>	APOTEX INC	<u>EQ 0.1% BASE</u>	<u>A078350 001</u>	Dec 07, 2015
<u>AT</u>		<u>EQ 0.2% BASE</u>	<u>A090918 001</u>	Dec 05, 2017
<u>AT</u>	AUROBINDO PHARMA LTD	<u>EQ 0.1% BASE</u>	<u>A204812 001</u>	Dec 18, 2015
<u>AT</u>	BARR LABS INC	<u>EQ 0.2% BASE</u>	<u>A090848 001</u>	Jul 13, 2015
<u>AT</u>	CIPLA LTD	<u>EQ 0.1% BASE</u>	<u>A206046 001</u>	Jul 26, 2017
<u>AT</u>		<u>EQ 0.2% BASE</u>	<u>A206087 001</u>	Dec 05, 2017
<u>AT</u>	SOMERSET THERAPS LLC	<u>EQ 0.1% BASE</u>	<u>A206306 001</u>	Dec 07, 2015
<u>AT</u>	USV NORTH AMERICA	<u>EQ 0.1% BASE</u>	<u>A203152 001</u>	Dec 07, 2015
<u>AT</u>	WOCKHARDT LTD	<u>EQ 0.1% BASE</u>	<u>A200810 001</u>	Jun 28, 2017
<u>AT</u>	ZAMBON SPA	<u>EQ 0.1% BASE</u>	<u>A204706 001</u>	Dec 07, 2015

PATADAY

<u>AT</u>	+! NOVARTIS PHARMS CORP	<u>EQ 0.2% BASE</u>	<u>N021545 001</u>	Dec 22, 2004
-----------	-------------------------	---------------------	--------------------	--------------

PATANOL

<u>AT</u>	+! NOVARTIS PHARMS CORP PAZEO	<u>EQ 0.1% BASE</u>	<u>N020688 001</u>	Dec 18, 1996
	+! NOVARTIS PHARMS CORP	EQ 0.7% BASE		N206276 001 Jan 30, 2015

SPRAY, METERED;NASAL

OLOPATADINE HYDROCHLORIDE

<u>AB</u>	APOTEX INC	<u>0.665MG/SPRAY</u>	<u>A091572 001</u>	Oct 08, 2014
<u>AB</u>	PERRIGO ISRAEL	<u>0.665MG/SPRAY</u>	<u>A202853 001</u>	Jan 31, 2017

PATANASE

<u>AB</u>	+! NOVARTIS PHARMS CORP	<u>0.665MG/SPRAY</u>	<u>N021861 001</u>	Apr 15, 2008
-----------	-------------------------	----------------------	--------------------	--------------

OLSALAZINE SODIUM

CAPSULE;ORAL

DIPENTUM

+! MYLAN SPECIALITY LP 250MG

N019715 001 Jul 31, 1990

OMACETAXINE MEPESUCCINATE

POWDER;SUBCUTANEOUS

SYNRIBO

+! TEVA PHARMS INTL 3.5MG/VIAL

N203585 001 Oct 26, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-320 (of 436)

OMBITASVIR; PARITAPREVIR; RITONAVIR

TABLET;ORAL

TECHNIVIE

+! ABBVIE INC

12.5MG;75MG;50MG

N207931 001 Jul 24, 2015

OMEGA-3-ACID ETHYL ESTERS

CAPSULE;ORAL

LOVAZA

AB +! SMITHKLINE BEECHAM

1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

N021654 001 Nov 10, 2004

OMEGA-3-ACID ETHYL ESTERS

AB AMNEAL PHARMS

1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A204940 001 Nov 27, 2015

AB APOTEX INC

1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A090973 001 Sep 30, 2014

AB PAR PHARM INC

1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A091018 001 Jun 24, 2014

AB STRIDES PHARMA

1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A203893 001 Sep 19, 2017

AB TEVA PHARMS USA

1GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

A091028 001 Apr 07, 2014

OMEGA-3-CARBOXYLIC ACIDS

CAPSULE;ORAL

EPANOVA

+! ASTRAZENECA PHARMS

1GM CONTAINS AT LEAST 850MG OF POLYUNSATURATED FATTY ACIDS

N205060 001 May 05, 2014

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS;ORAL

OMEPRAZOLE

AB ACTAVIS LABS FL INC 10MG

A075347 001 May 30, 2008

AB 20MG

A075347 002 May 30, 2008

AB 40MG

A075347 003 May 30, 2008

AB APOTEX 10MG

A076048 001 Oct 22, 2007

AB 20MG

A076048 002 Oct 22, 2007

AB 40MG

A076048 003 Jan 21, 2009

AB AUROBINDO PHARMA LTD 10MG

A203270 001 Aug 19, 2015

AB 20MG

A203270 002 Aug 19, 2015

AB 40MG

A203270 003 Aug 19, 2015

AB BRECKENRIDGE PHARM 10MG

A203481 001 Jul 03, 2017

AB 20MG

A203481 002 Jul 03, 2017

AB 40MG

A203481 003 Jul 03, 2017

AB DR REDDYS LABS LTD 10MG

A075576 003 Oct 22, 2007

AB 10MG

A078490 002 Mar 16, 2009

AB 20MG

A075576 002 Oct 22, 2007

AB 20MG

A078490 003 Mar 16, 2009

AB 40MG

A075576 001 Jan 21, 2009

AB 40MG

A078490 001 Apr 17, 2009

AB GLENMARK GENERICS 10MG

A091672 001 Oct 31, 2014

AB 20MG

A091672 002 Oct 31, 2014

AB 40MG

A091672 003 Oct 31, 2014

AB IMPAX LABS 10MG

A075785 001 Oct 22, 2007

AB 20MG

A075785 002 Oct 22, 2007

AB 40MG

A075785 003 Jan 21, 2009

AB KREMERS URBAN PHARMS 10MG

A075410 001 Nov 01, 2002

AB 20MG

A075410 002 Nov 01, 2002

AB 40MG

A075410 003 Jan 23, 2009

AB LUPIN LTD 40MG

A202384 001 Aug 25, 2015

AB MYLAN 10MG

A075876 001 May 29, 2003

AB 20MG

A075876 002 May 29, 2003

AB 40MG

A075876 003 Jan 21, 2009

AB SANDOZ 10MG

A075757 001 Jan 28, 2003

AB ! 20MG

A075757 002 Jan 28, 2003

AB ! 40MG

A076515 001 Jan 21, 2009

AB TEVA PHARMS USA 20MG

A204661 001 Jun 13, 2017

AB 40MG

A204661 002 Jun 13, 2017

AB ZYDUS PHARMS USA INC 10MG

A091352 001 Nov 19, 2012

AB 20MG

A091352 002 Nov 19, 2012

AB 40MG

A091352 003 Nov 19, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-321 (of 436)

OMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL  
 PRILOSEC

+ COVIS PHARMA BV EQ 2.5MG BASE/PACKET  
 +! EQ 10MG BASE/PACKET

N022056 001 Mar 20, 2008  
 N022056 002 Mar 20, 2008

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG;1.1GM</u>
<u>AB</u>		<u>40MG;1.1GM</u>
<u>AB</u>	AUROLIFE PHARMA LLC	<u>20MG;1.1GM</u>
<u>AB</u>		<u>40MG;1.1GM</u>
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG;1.1GM</u>
<u>AB</u>		<u>40MG;1.1GM</u>
<u>AB</u>	PAR PHARM	<u>20MG;1.1GM</u>
<u>AB</u>		<u>40MG;1.1GM</u>
<u>AB</u>	SCIEGEN PHARMS INC	<u>20MG;1.1GM</u>
<u>AB</u>		<u>40MG;1.1GM</u>

<u>A204228 001</u>	Jul 15, 2016
<u>A204228 002</u>	Jul 15, 2016
<u>A204922 001</u>	Aug 19, 2016
<u>A204922 002</u>	Aug 19, 2016
<u>A204068 001</u>	Jul 15, 2016
<u>A204068 002</u>	Jul 15, 2016
<u>A078966 001</u>	May 25, 2010
<u>A078966 002</u>	May 25, 2010
<u>A207476 001</u>	Dec 06, 2016
<u>A207476 002</u>	Dec 06, 2016

ZEGERID

<u>AB</u> + SANTARUS INC	<u>20MG;1.1GM</u>
<u>AB</u> +!	<u>40MG;1.1GM</u>

<u>N021849 001</u>	Feb 27, 2006
<u>N021849 002</u>	Feb 27, 2006

FOR SUSPENSION;ORAL

OMEPRAZOLE AND SODIUM BICARBONATE

<u>AB</u>	AJANTA PHARMA LTD	<u>20MG/PACKET;1.68GM/PACKET</u>
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>
<u>AB</u>	PAR PHARM	<u>20MG/PACKET;1.68GM/PACKET</u>
<u>AB</u>		<u>40MG/PACKET;1.68GM/PACKET</u>

<u>A205545 001</u>	Jul 27, 2016
<u>A205545 002</u>	Jul 27, 2016
<u>A079182 001</u>	Apr 19, 2013
<u>A079182 002</u>	Apr 19, 2013

ZEGERID

<u>AB</u> + SANTARUS INC	<u>20MG/PACKET;1.68GM/PACKET</u>
<u>AB</u> +!	<u>40MG/PACKET;1.68GM/PACKET</u>

<u>N021636 001</u>	Jun 15, 2004
<u>N021636 002</u>	Dec 21, 2004

ONDANSETRON

FILM;ORAL

ZUPLENZ

+ MEDIATECH PHARMA US 4MG  
 +! 8MG

N022524 001 Jul 02, 2010  
 N022524 002 Jul 02, 2010

TABLET, ORALLY DISINTEGRATING;ORAL

ONDANSETRON

<u>AB</u>	AUROBINDO PHARMA	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	BARR	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	GLENMARK GENERICS	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	MYLAN	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	SANDOZ	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	SUN PHARM INDNS	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	SUN PHARM INDNS LTD	<u>4MG</u>
<u>AB</u>		<u>8MG</u>
<u>AB</u>	TEVA	<u>4MG</u>
<u>AB</u>		<u>8MG</u>

<u>A090469 001</u>	Apr 12, 2010
<u>A090469 002</u>	Apr 12, 2010
<u>A076693 001</u>	Jun 25, 2007
<u>A076693 002</u>	Jun 25, 2007
<u>A078152 001</u>	Jun 27, 2007
<u>A078152 002</u>	Jun 27, 2007
<u>A078139 001</u>	Jun 25, 2007
<u>A078139 002</u>	Jun 25, 2007
<u>A078050 001</u>	Aug 13, 2007
<u>A078050 002</u>	Aug 13, 2007
<u>A077557 001</u>	Aug 02, 2007
<u>A077557 002</u>	Aug 02, 2007
<u>A078602 001</u>	Feb 24, 2011
<u>A078602 002</u>	Feb 24, 2011
<u>A076810 001</u>	Jun 25, 2007
<u>A076810 002</u>	Jun 25, 2007

ZOFRAN ODT

<u>AB</u> + NOVARTIS PHARMS CORP	<u>4MG</u>
<u>AB</u> +!	<u>8MG</u>

<u>N020781 001</u>	Jan 27, 1999
<u>N020781 002</u>	Jan 27, 1999

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 2MG BASE/ML</u>
<u>AP</u> ! BAXTER HLTHCARE CORP		<u>EQ 2MG BASE/ML</u>
<u>AP</u>	EMCURE PHARMS	<u>EQ 2MG BASE/ML</u>
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 2MG BASE/ML</u>
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>

<u>A206846 001</u>	Jul 13, 2015
<u>A202599 001</u>	Dec 21, 2012
<u>A078288 001</u>	Feb 22, 2013
<u>A090424 001</u>	Apr 16, 2010
<u>A076974 001</u>	Dec 26, 2006
<u>A079224 001</u>	Sep 25, 2009
<u>A090648 001</u>	Jun 15, 2012
<u>A076781 001</u>	Dec 26, 2006
<u>A077473 001</u>	Dec 26, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-322 (of 436)

## ONDANSETRON HYDROCHLORIDE

## INJECTABLE; INJECTION

## ONDANSETRON HYDROCHLORIDE

<u>AP</u>	LUITPOLD	<u>EQ 2MG BASE/ML</u>	<u>A077840 001</u>	Jan 19, 2007
<u>AP</u>	MYLAN LABS LTD	<u>EQ 2MG BASE/ML</u>	<u>A079039 001</u>	Nov 18, 2008
<u>AP</u>	QILU PHARM CO LTD	<u>EQ 2MG BASE/ML</u>	<u>A204906 001</u>	Jul 31, 2017
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A203711 001</u>	Sep 08, 2014
<u>AP</u>	TEVA	<u>EQ 2MG BASE/ML</u>	<u>A077430 001</u>	Jun 27, 2007
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 2MG BASE/ML</u>	<u>A076876 001</u>	Nov 22, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A076967 001</u>	Dec 26, 2006
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077365 001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077577 001</u>	Dec 26, 2006
<b><u>ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE</u></b>				
<u>AP</u>	ACCORD HLTHCARE	<u>EQ 2MG BASE/ML</u>	<u>A206845 001</u>	Mar 10, 2016
<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 2MG BASE/ML</u>	<u>A202600 001</u>	Dec 21, 2012
<u>AP</u>	! BAXTER HLTHCARE CORP	<u>EQ 2MG BASE/ML</u>	<u>A078287 001</u>	Feb 22, 2013
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 2MG BASE/ML</u>	<u>A078945 001</u>	Jan 03, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 2MG BASE/ML</u>	<u>A076972 001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A202253 001</u>	Jul 19, 2013
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 2MG BASE/ML</u>	<u>A076780 001</u>	Dec 26, 2006
<u>AP</u>	HOSPIRA	<u>EQ 2MG BASE/ML</u>	<u>A077548 001</u>	Dec 26, 2006
<u>AP</u>	LUITPOLD	<u>EQ 2MG BASE/ML</u>	<u>A079032 001</u>	Nov 18, 2008
<u>AP</u>	SANDOZ INC	<u>EQ 2MG BASE/ML</u>	<u>A077551 001</u>	Jun 27, 2007
<u>AP</u>	SUN PHARM INDs LTD	<u>EQ 2MG BASE/ML</u>	<u>A077173 001</u>	Dec 26, 2006
<u>AP</u>	TEVA	<u>EQ 2MG BASE/ML</u>	<u>A076759 001</u>	Nov 22, 2006
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 2MG BASE/ML</u>	<u>A077011 001</u>	Dec 26, 2006
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>A077541 001</u>	Dec 26, 2006
<u>AP</u>	WOCKHARDT	<u>EQ 2MG BASE/ML</u>	<u>A077716 001</u>	Dec 26, 2006

## SOLUTION; ORAL

## ONDANSETRON HYDROCHLORIDE

<b>AA</b>	AMNEAL PHARMS	<b>EQ 4MG BASE/5ML</b>	<b>A091483 001</b>	Jan 31, 2011
<b>AA</b>	APOTEX	<b>EQ 4MG BASE/5ML</b>	<b>A078127 001</b>	Jun 25, 2007
<b>AA</b>	AUROBINDO PHARMA	<b>EQ 4MG BASE/5ML</b>	<b>A078776 001</b>	Nov 28, 2007
<b>AA</b>	SILARX	<b>EQ 4MG BASE/5ML</b>	<b>A091342 001</b>	Jan 27, 2011
<b>AA</b>	TARO	<b>EQ 4MG BASE/5ML</b>	<b>A077009 001</b>	Nov 30, 2007
<b>AA</b>	WEST-WARD PHARMS	<b>EQ 4MG BASE/5ML</b>	<b>A076960 001</b>	Dec 26, 2006
	TNT			

ZOFRAN

**AA** +! NOVARTIS PHARMS CORP **EQ 4MG BASE/5ML**

N020605 001 Jan 24, 1997

**TABLET; ORAL**

#### **ONDANSETRON HYDROCHLORIDE**

<u><a href="#">AB</a></u>	APOTEX	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A077306 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	AUROBINDO PHARMA	<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A077306 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A078539 001</a></u>	Jul 31, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A078539 002</a></u>	Jul 31, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 24MG BASE</a></u>	<u><a href="#">A078539 003</a></u>	Jul 31, 2007
<u><a href="#">AB</a></u>	DR REDDYS LABS LTD	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A076183 003</a></u>	Dec 26, 2006
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A076183 002</a></u>	Dec 26, 2006
<u><a href="#">AB</a></u>		<u><a href="#">EO 24MG BASE</a></u>	<u><a href="#">A076183 001</a></u>	Dec 26, 2006
<u><a href="#">AB</a></u>	GLENMARK GENERICS	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A077535 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A077535 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 24MG BASE</a></u>	<u><a href="#">A077535 003</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	IPCA LABS LTD	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A203761 001</a></u>	Jan 23, 2014
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A203761 002</a></u>	Jan 23, 2014
<u><a href="#">AB</a></u>	MYLAN	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A076930 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A076930 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 24MG BASE</a></u>	<u><a href="#">A076930 004</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	NATCO PHARMA LTD	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A077851 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A077851 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	PLIVA HRVATSKA DOO	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A077112 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A077112 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 24MG BASE</a></u>	<u><a href="#">A077112 003</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	SANDOZ	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A077517 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A077517 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 24MG BASE</a></u>	<u><a href="#">A077517 003</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	SUN PHARM INDs (IN)	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A077050 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A077050 002</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>	TEVA	<u><a href="#">EO 4MG BASE</a></u>	<u><a href="#">A076252 001</a></u>	Jun 25, 2007
<u><a href="#">AB</a></u>		<u><a href="#">EO 8MG BASE</a></u>	<u><a href="#">A076252 002</a></u>	Jun 25, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-323 (of 436)

ONDANSETRON HYDROCHLORIDE

TABLET;ORAL

ONDANSETRON HYDROCHLORIDE

<u>AB</u>		<u>EQ 24MG BASE</u>	<u>A076252 003</u>	Jun 25, 2007
<u>ZOFRAN</u>				
<u>AB</u>	+	NOVARTIS PHARMS CORP	<u>EQ 4MG BASE</u>	<u>N020103 001</u> Dec 31, 1992
<u>AB</u>	+		<u>EQ 8MG BASE</u>	<u>N020103 002</u> Dec 31, 1992
<u>AB</u>	+!	ONDANSETRON HYDROCHLORIDE DR REDDYS LABS LTD	<u>EQ 24MG BASE</u>	<u>N020103 003</u> Aug 27, 1999
				EQ 16MG BASE
				A076183 004 Dec 26, 2006

ORITAVANCIN DIPHOSPHATE

POWDER;IV (INFUSION)

ORBACTIV

+! THE MEDICINES CO

EQ 400MG BASE/VIAL

N206334 001 Aug 06, 2014

ORLISTAT

CAPSULE;ORAL

XENICAL

+! CHEPLAPHARM

120MG

N020766 001 Apr 23, 1999

ORPHENADRINE CITRATE

INJECTABLE;INJECTION

ORPHENADRINE CITRATE

<u>AP</u>	!	AKORN	<u>30MG/ML</u>	<u>A040484 001</u> May 24, 2006
<u>AP</u>		SAGENT PHARMS	<u>30MG/ML</u>	<u>A090585 001</u> Aug 30, 2011
<u>AP</u>		WATSON LABS	<u>30MG/ML</u>	<u>A084779 001</u> Mar 15, 1982
<u>AP</u>		WEST-WARD PHARMS INT	<u>30MG/ML</u>	<u>A040463 001</u> Mar 04, 2003

TABLET, EXTENDED RELEASE;ORAL

ORPHENADRINE CITRATE

<u>AB</u>		GAVIS PHARMS	<u>100MG</u>	<u>A040284 001</u> Jun 19, 1998
<u>AB</u>		IMPAX PHARMS	<u>100MG</u>	<u>A040368 001</u> Jun 23, 2000
<u>AB</u>		INVAGEN PHARMS	<u>100MG</u>	<u>A091158 001</u> Jul 27, 2012
<u>AB</u>	!	SANDOZ	<u>100MG</u>	<u>A040327 001</u> Feb 15, 2000
<u>AB</u>		TEDOR PHARMA INC	<u>100MG</u>	<u>A040249 001</u> Jan 29, 1999

OSELTAMIVIR PHOSPHATE

CAPSULE;ORAL

OSELTAMIVIR PHOSPHATE

<u>AB</u>		AMNEAL PHARMS	<u>EQ 30MG BASE</u>	<u>A209093 001</u> May 17, 2017
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A209093 002</u> May 17, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A209093 003</u> May 17, 2017
<u>AB</u>		LUPIN ATLANTIS	<u>EQ 30MG BASE</u>	<u>A208348 001</u> Jan 09, 2018
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A208348 002</u> Jan 09, 2018
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A208348 003</u> Jan 09, 2018
<u>AB</u>		MACLEODS PHARMS LTD	<u>EQ 30MG BASE</u>	<u>A207211 001</u> Sep 14, 2017
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A207211 002</u> Sep 14, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A207211 003</u> Sep 14, 2017
<u>AB</u>		NATCO PHARMA LTD	<u>EQ 30MG BASE</u>	<u>A202595 001</u> Aug 03, 2016
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A202595 002</u> Aug 03, 2016
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A202595 003</u> Aug 03, 2016
<u>AB</u>		NESHER PHARMS	<u>EQ 30MG BASE</u>	<u>A208578 001</u> Feb 24, 2017
<u>AB</u>			<u>EQ 45MG BASE</u>	<u>A208578 002</u> Feb 24, 2017
<u>AB</u>			<u>EQ 75MG BASE</u>	<u>A208578 003</u> Feb 24, 2017

TAMIFLU

<u>AB</u>	+	ROCHE	<u>EQ 30MG BASE</u>
<u>AB</u>	+		<u>EQ 45MG BASE</u>
<u>AB</u>	+!		<u>EQ 75MG BASE</u>

FOR SUSPENSION;ORAL

OSELTAMIVIR PHOSPHATE

<u>AB</u>		ALVOGEN PINE BROOK	<u>EQ 6MG BASE/ML</u>	<u>A208823 001</u> Oct 31, 2017
<u>AB</u>		NESHER PHARMS	<u>EQ 6MG BASE/ML</u>	<u>A209113 001</u> Sep 14, 2017

TAMIFLU

<u>AB</u>	+!	ROCHE	<u>EQ 6MG BASE/ML</u>
-----------	----	-------	-----------------------

N021087 003 Jul 02, 2007

N021087 002 Jul 02, 2007

N021087 001 Oct 27, 1999

OSIMERTINIB MESYLATE

TABLET;ORAL

TAGRISSO

+	ASTRAZENECA PHARMS	<u>EQ 40MG BASE</u>
+		<u>EQ 80MG BASE</u>

N208065 001 Nov 13, 2015

N208065 002 Nov 13, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-324 (of 436)

OSPEMIFENE

TABLET;ORAL

OSPHENA

+! DUCHESNAY

60MG

N203505 001 Feb 26, 2013

OXACILLIN SODIUM

INJECTABLE; INJECTION

OXACILLIN SODIUM

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 1GM BASE/VIAL</u>	<u>A201539 001</u>	Jan 18, 2013
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A201539 002</u>	Jan 18, 2013
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A201538 001</u>	Jan 18, 2013
<u>AP</u>	HOSPIRA INC	<u>EQ 1GM BASE/VIAL</u>	<u>A203950 001</u>	Dec 11, 2015
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A203950 002</u>	Dec 11, 2015
<u>AP</u>	RENAISSANCE SSA LLC	<u>EQ 1GM BASE/VIAL</u>	<u>A206681 001</u>	Sep 11, 2017
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A206681 002</u>	Sep 11, 2017
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A206760 001</u>	Oct 26, 2017
<u>AP</u>	SAGENT PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>A091246 001</u>	Mar 30, 2012
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A091246 002</u>	Mar 30, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A091245 001</u>	Mar 30, 2012
<u>AP</u>	! SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A061490 003</u>	
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A061490 004</u>	
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A061490 006</u>	May 09, 1991
<u>AP</u>	WOCKHARDT BIO AG	<u>EQ 1GM BASE/VIAL</u>	<u>A207147 001</u>	Jul 31, 2017
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>A207147 002</u>	Jul 31, 2017
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A207148 001</u>	Nov 24, 2017
	BACTOCILL IN PLASTIC CONTAINER			
	+! BAXTER HLTHCARE	EQ 20MG BASE/ML	N050640 001	Oct 26, 1989
	+!	EQ 40MG BASE/ML	N050640 002	Oct 26, 1989
	POWDER; INTRAVENOUS			
	<u>OXACILLIN SODIUM</u>			
<u>AP</u>	! SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>A062737 001</u>	Dec 23, 1986
<u>AP</u>	!	<u>EQ 2GM BASE/VIAL</u>	<u>A062737 002</u>	Dec 23, 1986

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

<u>AP</u>	+! SANOFI AVENTIS US	<u>50MG/10ML (5MG/ML)</u>	<u>N021759 001</u>	Jan 31, 2005
<u>AP</u>	+!	<u>100MG/20ML (5MG/ML)</u>	<u>N021759 002</u>	Jan 31, 2005

OXALIPLATIN

<u>AP</u>	ACCORD HLTHCARE	<u>50MG/10ML (5MG/ML)</u>	<u>A207474 001</u>	Mar 21, 2017
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A207474 002</u>	Mar 21, 2017
<u>AP</u>		<u>200MG/40ML (5MG/ML)</u>	<u>A207474 003</u>	Mar 21, 2017
<u>AP</u>	ACTAVIS TOTOWA	<u>50MG/VIAL</u>	<u>A078803 001</u>	Aug 08, 2012
<u>AP</u>		<u>100MG/VIAL</u>	<u>A078803 002</u>	Aug 08, 2012
<u>AP</u>	CIPILA LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A208523 001</u>	Feb 10, 2017
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A208523 002</u>	Feb 10, 2017
<u>AP</u>	FRESENIUS KABI ONCOL	<u>50MG/10ML (5MG/ML)</u>	<u>A078811 001</u>	Jun 10, 2010
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A078811 002</u>	Jun 10, 2010
<u>AP</u>		<u>50MG/VIAL</u>	<u>A078810 001</u>	Aug 07, 2009
<u>AP</u>		<u>100MG/VIAL</u>	<u>A078810 002</u>	Aug 07, 2009
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/VIAL</u>	<u>A078819 001</u>	Jun 02, 2010
<u>AP</u>		<u>50MG/10ML (5MG/ML)</u>	<u>A090030 001</u>	Jan 31, 2017
<u>AP</u>		<u>100MG/VIAL</u>	<u>A078819 002</u>	Jun 02, 2010
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A090030 002</u>	Jan 31, 2017
<u>AP</u>		<u>200MG/40ML (5MG/ML)</u>	<u>A090030 003</u>	Jan 31, 2017
<u>AP</u>	GLAND PHARMA LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A207325 001</u>	Feb 10, 2017
<u>AP</u>		<u>50MG/VIAL</u>	<u>A207385 001</u>	May 23, 2017
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A207325 002</u>	Feb 10, 2017
<u>AP</u>		<u>100MG/VIAL</u>	<u>A207385 002</u>	May 23, 2017
<u>AP</u>		<u>200MG/40ML (5MG/ML)</u>	<u>A207325 003</u>	Oct 18, 2017
<u>AP</u>	HOSPIRA INC	<u>50MG/VIAL</u>	<u>A078815 001</u>	Sep 30, 2009
<u>AP</u>		<u>100MG/VIAL</u>	<u>A078815 002</u>	Sep 30, 2009
<u>AP</u>	HOSPIRA WORLDWIDE	<u>50MG/10ML (5MG/ML)</u>	<u>A078813 001</u>	Aug 07, 2009
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A078813 002</u>	Aug 07, 2009
<u>AP</u>	JIANGSU HENGRIUI MED	<u>50MG/10ML (5MG/ML)</u>	<u>A203869 001</u>	Jun 18, 2014
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A203869 002</u>	Jun 18, 2014
<u>AP</u>	LUITPOLD PHARMS INC	<u>50MG/10ML (5MG/ML)</u>	<u>A204378 001</u>	May 12, 2017
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A204378 002</u>	May 12, 2017
<u>AP</u>	MYLAN LABS LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A091358 001</u>	Aug 07, 2012
<u>AP</u>		<u>50MG/VIAL</u>	<u>A200979 001</u>	Aug 08, 2012
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A091358 002</u>	Aug 07, 2012
<u>AP</u>		<u>100MG/VIAL</u>	<u>A200979 002</u>	Aug 08, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-325 (of 436)

OXALIPLATIN

INJECTABLE; IV (INFUSION)

OXALIPLATIN

<u>AP</u>		<u>200MG/40ML (5MG/ML)</u>	<u>A091358 003</u>	Nov 14, 2017
<u>AP</u>	QILU PHARM CO LTD	<u>50MG/10ML (5MG/ML)</u>	<u>A204368 001</u>	Jun 07, 2016
<u>AP</u>		<u>50MG/VIAL</u>	<u>A204616 001</u>	May 11, 2016
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A204368 002</u>	Jun 07, 2016
<u>AP</u>		<u>100MG/VIAL</u>	<u>A204616 002</u>	May 11, 2016
<u>AP</u>	!	<u>200MG/40ML (5MG/ML)</u>	<u>A204368 003</u>	Jun 07, 2016
<u>AP</u>	SANDOZ	<u>50MG/10ML (5MG/ML)</u>	<u>A078817 001</u>	Jan 24, 2011
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A078817 002</u>	Jan 24, 2011
<u>AP</u>	SANJA PHARMS CO	<u>50MG/10ML (5MG/ML)</u>	<u>A205529 001</u>	Sep 06, 2017
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A205529 002</u>	Sep 06, 2017
<u>AP</u>	! SUN PHARMA GLOBAL	<u>50MG/VIAL</u>	<u>A078818 001</u>	Aug 07, 2009
<u>AP</u>		<u>50MG/10ML (5MG/ML)</u>	<u>A202922 001</u>	Apr 08, 2014
<u>AP</u>	!	<u>100MG/VIAL</u>	<u>A078818 002</u>	Aug 07, 2009
<u>AP</u>		<u>100MG/20ML (5MG/ML)</u>	<u>A202922 002</u>	Apr 08, 2014
<u>AP</u>	+! TEVA PHARMS	<u>50MG/10ML (5MG/ML)</u>	<u>N022160 001</u>	Aug 07, 2009
<u>AP</u>	+!	<u>100MG/20ML (5MG/ML)</u>	<u>N022160 002</u>	Aug 07, 2009

OXANDROLONE

TABLET; ORAL

OXANDRIN

<u>AB</u>	+ GEMINI LABS LLC	<u>2.5MG</u>	<u>N013718 001</u>	
<u>AB</u>	+!	<u>10MG</u>	<u>N013718 002</u>	Nov 05, 2001
	<u>OXANDROLONE</u>			
<u>AB</u>	PAR PHARM	<u>2.5MG</u>	<u>A077827 001</u>	Jun 22, 2007
<u>AB</u>		<u>10MG</u>	<u>A077827 002</u>	Jun 22, 2007
<u>AB</u>	UPSHER-SMITH LABS	<u>2.5MG</u>	<u>A076761 001</u>	Dec 01, 2006
<u>AB</u>		<u>10MG</u>	<u>A078033 001</u>	Mar 22, 2007

OXAPROZIN

TABLET; ORAL

DAYPRO

<u>AB</u>	+! GD SEARLE	<u>600MG</u>	<u>N018841 004</u>	Oct 29, 1992
	<u>OXAPROZIN</u>			
<u>AB</u>	AMNEAL PHARMS CO	<u>600MG</u>	<u>A208633 001</u>	May 04, 2017
<u>AB</u>	APOTEX INC	<u>600MG</u>	<u>A075987 001</u>	Sep 02, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>600MG</u>	<u>A075855 001</u>	Jan 31, 2001
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>600MG</u>	<u>A075846 001</u>	May 13, 2002
<u>AB</u>	SANDOZ	<u>600MG</u>	<u>A075845 001</u>	Jan 31, 2001
<u>AB</u>	SUN PHARM INDNS INC	<u>600MG</u>	<u>A075844 001</u>	Jan 03, 2002
<u>AB</u>	TEVA	<u>600MG</u>	<u>A075849 001</u>	Jul 03, 2002

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>10MG</u>	<u>A072253 002</u>	Apr 14, 1988
<u>AB</u>		<u>15MG</u>	<u>A072253 003</u>	Apr 14, 1988
<u>AB</u>	!	<u>30MG</u>	<u>A072253 001</u>	Apr 14, 1988
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A071813 001</u>	Apr 19, 1988
<u>AB</u>		<u>15MG</u>	<u>A071756 001</u>	Apr 19, 1988
<u>AB</u>		<u>30MG</u>	<u>A071814 001</u>	Apr 19, 1988

OXCARBAZEPINE

SUSPENSION; ORAL

OXCARBAZEPINE

<u>AB</u>	AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A202961 001</u>	Sep 17, 2012
<u>AB</u>	SUN PHARM INDNS LTD	<u>300MG/5ML</u>	<u>A078734 001</u>	Jun 26, 2009
<u>AB</u>	WEST-WARD PHARMS INT	<u>300MG/5ML</u>	<u>A201193 001</u>	Oct 03, 2012

TRILEPTAL

<u>AB</u>	+! NOVARTIS	<u>300MG/5ML</u>	<u>N021285 001</u>	May 25, 2001
-----------	-------------	------------------	--------------------	--------------

TABLET; ORAL

OXCARBAZEPINE

<u>AB</u>	ANI PHARMS INC	<u>150MG</u>	<u>A078005 001</u>	Dec 11, 2007
<u>AB</u>		<u>300MG</u>	<u>A078005 002</u>	Dec 11, 2007
<u>AB</u>		<u>600MG</u>	<u>A078005 003</u>	Dec 11, 2007
<u>AB</u>	APOTEX INC	<u>150MG</u>	<u>A077747 001</u>	Apr 09, 2008
<u>AB</u>		<u>300MG</u>	<u>A077747 002</u>	Apr 09, 2008
<u>AB</u>		<u>600MG</u>	<u>A077747 003</u>	Apr 09, 2008
<u>AB</u>	BRECKENRIDGE PHARM	<u>150MG</u>	<u>A078069 001</u>	Jan 11, 2008
<u>AB</u>		<u>300MG</u>	<u>A078069 002</u>	Jan 11, 2008
<u>AB</u>		<u>600MG</u>	<u>A078069 003</u>	Jan 11, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-326 (of 436)

OXCARBAZEPINE

TABLET;ORAL

**OXCARBAZEPINE**

<b>AB</b>	GLENMARK PHARMS LTD	<u>150MG</u>	<b>A077802 001</b>	Oct 09, 2007
<b>AB</b>		<u>300MG</u>	<b>A077802 002</b>	Oct 09, 2007
<b>AB</b>		<u>600MG</u>	<b>A077802 003</b>	Oct 09, 2007
<b>AB</b>	SUN PHARM INDNS	<u>150MG</u>	<b>A077794 001</b>	Oct 09, 2007
<b>AB</b>		<u>300MG</u>	<b>A077794 002</b>	Oct 09, 2007
<b>AB</b>		<u>600MG</u>	<b>A077794 003</b>	Oct 09, 2007
<b>AB</b>	TARO	<u>150MG</u>	<b>A077801 001</b>	Nov 15, 2007
<b>AB</b>		<u>300MG</u>	<b>A077801 002</b>	Nov 15, 2007
<b>AB</b>		<u>600MG</u>	<b>A077801 003</b>	Nov 15, 2007
<b>AB</b>	WEST-WARD PHARMS INT	<u>150MG</u>	<b>A077795 001</b>	Oct 09, 2007
<b>AB</b>		<u>300MG</u>	<b>A077795 002</b>	Oct 09, 2007
<b>AB</b>		<u>600MG</u>	<b>A077795 003</b>	Oct 09, 2007

**TRILEPTAL**

<b>AB</b>	+	NOVARTIS	<u>150MG</u>	<b>N021014 001</b>	Jan 14, 2000
<b>AB</b>	+		<u>300MG</u>	<b>N021014 002</b>	Jan 14, 2000
<b>AB</b>	+!		<u>600MG</u>	<b>N021014 003</b>	Jan 14, 2000

TABLET, EXTENDED RELEASE;ORAL  
 OXTELLAR XR

+	SUPERNUS PHARMS	150MG	N202810 001	Oct 19, 2012
+		300MG	N202810 002	Oct 19, 2012
+!		600MG	N202810 003	Oct 19, 2012

OXICONAZOLE NITRATE

CREAM;TOPICAL

**OXICONAZOLE NITRATE**

<b>AB</b>	TARO	<u>EQ 1% BASE</u>	<b>A205076 001</b>	Mar 07, 2016	
<b>AB</b>	+!	FOUGERA PHARMS	<u>EQ 1% BASE</u>	<b>N019828 001</b>	Dec 30, 1988
LOTION;TOPICAL					
OXISTAT					
+!	FOUGERA PHARMS	EQ 1% BASE	N020209 001	Sep 30, 1992	

OXTRIPTYLLINE

TABLET, EXTENDED RELEASE;ORAL  
 CHOLEDYL SA

!	WARNER CHILCOTT LLC	400MG	A087863 001	May 24, 1983
---	---------------------	-------	-------------	--------------

OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL

**OXYBUTYNIN**

<b>AB</b>	BARR LABS DIV TEVA	<u>3.9MG/24HR</u>	<b>A090526 001</b>	Mar 04, 2014	
<b>AB</b>	+!	ALLERGAN SALES LLC	<u>3.9MG/24HR</u>	<b>N021351 002</b>	Feb 26, 2003

OXYBUTYNIN CHLORIDE

GEL;TRANSDERMAL

GELNIQUE

+!	ALLERGAN SALES LLC	10%(100MG/PACKET)	N022204 001	Jan 27, 2009
----	--------------------	-------------------	-------------	--------------

SYRUP;ORAL

**OXYBUTYNIN CHLORIDE**

<b>AA</b>	PHARM ASSOC	<u>5MG/5ML</u>	<b>A075137 001</b>	Dec 18, 1998
<b>AA</b>	SILARK	<u>5MG/5ML</u>	<b>A074520 001</b>	Mar 29, 1996
<b>AA</b>	VINTAGE PHARMS	<u>5MG/5ML</u>	<b>A076682 001</b>	Dec 28, 2004
<b>AA</b>	! WOCKHARDT BIO AG	<u>5MG/5ML</u>	<b>A074868 001</b>	Feb 12, 1997

TABLET;ORAL

**OXYBUTYNIN CHLORIDE**

<b>AB</b>	ABHAI LLC	<u>5MG</u>	<b>A209335 001</b>	Dec 22, 2017
<b>AB</b>	APPCO PHARMA LLC	<u>5MG</u>	<b>A209025 001</b>	Dec 21, 2017
<b>AB</b>	NOVITIUM PHARMA	<u>5MG</u>	<b>A209823 001</b>	Oct 23, 2017
<b>AB</b>	TEVA PHARMS USA	<u>5MG</u>	<b>A071655 001</b>	Nov 14, 1988
<b>AB</b>	UPSHER-SMITH LABS	<u>5MG</u>	<b>A074625 001</b>	Jul 31, 1996
<b>AB</b>	! VINTAGE PHARMS	<u>5MG</u>	<b>A075079 001</b>	Oct 31, 1997

TABLET, EXTENDED RELEASE;ORAL

**DITROPAN XL**

<b>AB</b>	+	JANSSEN PHARMS	<u>5MG</u>	<b>N020897 001</b>	Dec 16, 1998
<b>AB</b>	+		<u>10MG</u>	<b>N020897 002</b>	Dec 16, 1998
<b>AB</b>	+!		<u>15MG</u>	<b>N020897 003</b>	Jun 22, 1999

**OXYBUTYNIN CHLORIDE**

<b>AB</b>	ACCORD HLTHCARE	<u>5MG</u>	<b>A207138 001</b>	Feb 29, 2016
<b>AB</b>		<u>10MG</u>	<b>A207138 002</b>	Feb 29, 2016
<b>AB</b>		<u>15MG</u>	<b>A207138 003</b>	Feb 29, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-327 (of 436)

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE; ORAL

OXYBUTYNIN CHLORIDE

<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A204010 001</u>	Nov 23, 2015
<u>AB</u>		<u>10MG</u>	<u>A204010 002</u>	Nov 23, 2015
<u>AB</u>		<u>15MG</u>	<u>A204010 003</u>	Nov 23, 2015
<u>AB</u>	IMPAX PHARMS	<u>5MG</u>	<u>A076745 002</u>	May 09, 2007
<u>AB</u>		<u>10MG</u>	<u>A076745 003</u>	May 09, 2007
<u>AB</u>		<u>15MG</u>	<u>A076745 001</u>	Nov 09, 2006
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076702 001</u>	Nov 09, 2006
<u>AB</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A076644 001</u>	Nov 09, 2006
<u>AB</u>		<u>15MG</u>	<u>A076644 002</u>	May 10, 2007
<u>AB</u>	OSMOTICA PHARM US	<u>5MG</u>	<u>A078503 001</u>	Feb 04, 2009
<u>AB</u>		<u>10MG</u>	<u>A078503 002</u>	Feb 04, 2009
<u>AB</u>		<u>15MG</u>	<u>A078503 003</u>	Feb 04, 2009
<u>AB</u>	UNIQUE PHARM LABS	<u>5MG</u>	<u>A206121 001</u>	May 27, 2016
<u>AB</u>		<u>10MG</u>	<u>A206121 002</u>	May 27, 2016
<u>AB</u>		<u>15MG</u>	<u>A206121 003</u>	May 27, 2016
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A202332 001</u>	Jun 26, 2017
<u>AB</u>		<u>10MG</u>	<u>A202332 002</u>	Jun 26, 2017
<u>AB</u>		<u>15MG</u>	<u>A202332 003</u>	Jun 26, 2017

OXYCODONE

CAPSULE, EXTENDED RELEASE; ORAL

XTAMPZA ER

+	COLLEGIUM PHARM INC	9MG
+		13.5MG
+		18MG
+		27MG
+!		36MG

N208090 001	Apr 26, 2016
N208090 002	Apr 26, 2016
N208090 003	Apr 26, 2016
N208090 004	Apr 26, 2016
N208090 005	Apr 26, 2016

OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS INC	<u>5MG</u>	<u>A205177 001</u>	Mar 31, 2016	
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A202773 001</u>	Aug 17, 2015	
<u>AB</u>	LANNETT HOLDINGS INC	<u>5MG</u>	<u>A203823 001</u>	Aug 01, 2014	
<u>AB</u>	+!	LEHIGH VALLEY	<u>5MG</u>	<u>N200534 001</u>	Oct 20, 2010
<u>AB</u>	MAYNE PHARMA INC	<u>5MG</u>	<u>A203107 001</u>	Jul 26, 2012	
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204752 001</u>	Aug 24, 2015	

SOLUTION; ORAL

OXYCODONE HYDROCHLORIDE

<u>AA</u>	ABHAI LLC	<u>5MG/5ML</u>	<u>A208593 001</u>	Jul 21, 2017	
<u>AA</u>		<u>100MG/5ML</u>	<u>A208593 002</u>	Jul 21, 2017	
<u>AA</u>	ANI PHARMS INC	<u>5MG/5ML</u>	<u>A204979 001</u>	Jun 01, 2015	
<u>AA</u>		<u>100MG/5ML</u>	<u>A203447 001</u>	Aug 30, 2017	
<u>AA</u>	ASCENT PHARMS INC	<u>5MG/5ML</u>	<u>A209021 001</u>	Nov 09, 2017	
<u>AA</u>		<u>100MG/5ML</u>	<u>A209021 002</u>	Nov 09, 2017	
<u>AA</u>	HI-TECH PHARMACAL	<u>5MG/5ML</u>	<u>A208817 001</u>	Aug 10, 2017	
<u>AA</u>		<u>100MG/5ML</u>	<u>A208795 001</u>	Aug 07, 2017	
<u>AA</u>	LANNETT HOLDINGS INC	<u>100MG/5ML</u>	<u>A204085 001</u>	Sep 09, 2014	
<u>AA</u>	+ LEHIGH VALLEY	<u>5MG/5ML</u>	<u>N200535 002</u>	Aug 22, 2013	
<u>AA</u>	+!	<u>100MG/5ML</u>	<u>N200535 001</u>	Oct 20, 2010	
<u>AA</u>	MAYNE PHARMA INC	<u>100MG/5ML</u>	<u>A204092 001</u>	Jun 05, 2014	
<u>AA</u>	NOVEL LABS INC	<u>100MG/5ML</u>	<u>A204603 001</u>	Apr 29, 2015	
<u>AA</u>	PHARM ASSOC	<u>100MG/5ML</u>	<u>A206822 001</u>	Aug 15, 2017	
<u>AA</u>	+!	VISTAPHARM	<u>5MG/5ML</u>	<u>N201194 001</u>	Jan 12, 2012
<u>AA</u>		<u>100MG/5ML</u>	<u>A202537 001</u>	Jul 30, 2012	
<u>AA</u>	WES PHARMA INC	<u>5MG/5ML</u>	<u>A207511 001</u>	Nov 23, 2016	
<u>AA</u>		<u>100MG/5ML</u>	<u>A209897 001</u>	Sep 06, 2017	
<u>AA</u>	WEST-WARD PHARMS INT	<u>5MG/5ML</u>	<u>A204037 001</u>	Jul 15, 2013	
<u>AA</u>		<u>100MG/5ML</u>	<u>A203208 001</u>	Jul 12, 2013	
<u>AA</u>	WOCKHARDT BIO AG	<u>5MG/5ML</u>	<u>A206456 001</u>	Jun 16, 2015	

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>5MG</u>	<u>A076636 003</u>	Apr 07, 2015
<u>AB</u>		<u>15MG</u>	<u>A076636 001</u>	Feb 06, 2004
<u>AB</u>		<u>30MG</u>	<u>A076636 002</u>	Feb 06, 2004
<u>AB</u>	ALVOGEN MALTA	<u>5MG</u>	<u>A202116 001</u>	Dec 30, 2011
<u>AB</u>		<u>15MG</u>	<u>A202116 002</u>	Dec 30, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-328 (of 436)

OXYCODONE HYDROCHLORIDE

TABLET;ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>		<u>30MG</u>	<u>A202116 003</u>	Dec 30, 2011
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A203638 001</u>	Jun 03, 2014
<u>AB</u>		<u>10MG</u>	<u>A203638 002</u>	Jun 03, 2014
<u>AB</u>		<u>15MG</u>	<u>A203638 003</u>	Jun 03, 2014
<u>AB</u>		<u>20MG</u>	<u>A203638 004</u>	Jun 03, 2014
<u>AB</u>		<u>30MG</u>	<u>A203638 005</u>	Jun 03, 2014
<u>AB</u>	ASCENT PHARMS INC	<u>15MG</u>	<u>A207418 001</u>	Aug 07, 2017
<u>AB</u>		<u>30MG</u>	<u>A207418 002</u>	Aug 07, 2017
<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A202160 001</u>	Nov 19, 2012
<u>AB</u>		<u>15MG</u>	<u>A202160 002</u>	Nov 19, 2012
<u>AB</u>		<u>30MG</u>	<u>A202160 003</u>	Nov 19, 2012
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A091393 001</u>	Aug 31, 2009
<u>AB</u>		<u>10MG</u>	<u>A091393 002</u>	Aug 31, 2009
<u>AB</u>		<u>15MG</u>	<u>A091393 003</u>	Aug 31, 2009
<u>AB</u>		<u>20MG</u>	<u>A091393 004</u>	Aug 31, 2009
<u>AB</u>		<u>30MG</u>	<u>A091393 005</u>	Aug 31, 2009
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A090895 001</u>	Aug 24, 2009
<u>AB</u>		<u>5MG</u>	<u>A202662 001</u>	Sep 22, 2015
<u>AB</u>		<u>10MG</u>	<u>A202662 002</u>	Sep 22, 2015
<u>AB</u>		<u>15MG</u>	<u>A090895 002</u>	Aug 24, 2009
<u>AB</u>		<u>15MG</u>	<u>A202662 003</u>	Sep 22, 2015
<u>AB</u>		<u>30MG</u>	<u>A090895 003</u>	Aug 24, 2009
<u>AB</u>		<u>30MG</u>	<u>A202662 004</u>	Sep 22, 2015
<u>AB</u>	KEN LIFESCIENCE	<u>5MG</u>	<u>A207119 001</u>	Apr 12, 2016
<u>AB</u>		<u>10MG</u>	<u>A207119 002</u>	Apr 12, 2016
<u>AB</u>		<u>15MG</u>	<u>A207119 003</u>	Apr 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A207119 004</u>	Apr 12, 2016
<u>AB</u>		<u>30MG</u>	<u>A207119 005</u>	Apr 12, 2016
<u>AB</u>	MAYNE PHARMA INC	<u>5MG</u>	<u>A091313 001</u>	Feb 18, 2011
<u>AB</u>		<u>10MG</u>	<u>A091313 004</u>	Apr 29, 2016
<u>AB</u>		<u>15MG</u>	<u>A091313 002</u>	Feb 18, 2011
<u>AB</u>		<u>20MG</u>	<u>A091313 005</u>	Apr 29, 2016
<u>AB</u>		<u>30MG</u>	<u>A091313 003</u>	Feb 18, 2011
<u>AB</u>	NESHER PHARMS	<u>5MG</u>	<u>A077290 001</u>	Dec 08, 2005
<u>AB</u>		<u>10MG</u>	<u>A077290 002</u>	Dec 08, 2005
<u>AB</u>		<u>15MG</u>	<u>A077290 003</u>	Dec 08, 2005
<u>AB</u>		<u>20MG</u>	<u>A077290 004</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>A077290 005</u>	Dec 08, 2005
<u>AB</u>	NOVEL LABS INC	<u>5MG</u>	<u>A204021 001</u>	Jun 12, 2017
<u>AB</u>		<u>10MG</u>	<u>A204021 002</u>	Jun 12, 2017
<u>AB</u>		<u>15MG</u>	<u>A204021 003</u>	Jun 12, 2017
<u>AB</u>		<u>20MG</u>	<u>A204021 004</u>	Jun 12, 2017
<u>AB</u>		<u>30MG</u>	<u>A204021 005</u>	Jun 12, 2017
<u>AB</u>	RHODES PHARMS	<u>5MG</u>	<u>A091490 001</u>	Mar 09, 2011
<u>AB</u>		<u>10MG</u>	<u>A091490 002</u>	Mar 09, 2011
<u>AB</u>		<u>15MG</u>	<u>A091490 003</u>	Mar 09, 2011
<u>AB</u>		<u>20MG</u>	<u>A091490 004</u>	Mar 09, 2011
<u>AB</u>		<u>30MG</u>	<u>A091490 005</u>	Mar 09, 2011
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A076758 003</u>	Mar 19, 2007
<u>AB</u>		<u>15MG</u>	<u>A076758 001</u>	Jun 30, 2004
<u>AB</u>		<u>30MG</u>	<u>A076758 002</u>	Jun 30, 2004
<u>AB</u>	SUN PHARM INDs INC	<u>5MG</u>	<u>A090659 001</u>	Apr 10, 2009
<u>AB</u>		<u>10MG</u>	<u>A090659 005</u>	Nov 06, 2012
<u>AB</u>		<u>15MG</u>	<u>A090659 002</u>	Apr 10, 2009
<u>AB</u>		<u>20MG</u>	<u>A090659 004</u>	Nov 06, 2012
<u>AB</u>		<u>30MG</u>	<u>A090659 003</u>	Apr 10, 2009
<u>AB</u>	VINTAGE PHARMS	<u>5MG</u>	<u>A077712 003</u>	Mar 02, 2009
<u>AB</u>		<u>10MG</u>	<u>A077712 004</u>	Apr 13, 2015
<u>AB</u>		<u>15MG</u>	<u>A077712 001</u>	Jan 31, 2007
<u>AB</u>		<u>20MG</u>	<u>A077712 005</u>	Apr 13, 2015
<u>AB</u>		<u>30MG</u>	<u>A077712 002</u>	Jan 31, 2007
<b><u>ROXICODONE</u></b>				
<u>AB</u>	+ SPECGX LLC	<u>5MG</u>	<u>N021011 003</u>	May 15, 2009
<u>AB</u>	+!	<u>15MG</u>	<u>N021011 001</u>	Aug 31, 2000
<u>AB</u>	+	<u>30MG</u>	<u>N021011 002</u>	Aug 31, 2000
OXAYDO				
	EGALET US INC	5MG	N202080 001	Jun 17, 2011
		7.5MG	N202080 002	Jun 17, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-329 (of 436)

OXYCODONE HYDROCHLORIDE

TABLET;ORAL

ROXYBOND

DAIICHI SANKYO INC	5MG	N209777	001	Apr 20,	2017
	15MG	N209777	002	Apr 20,	2017
	30MG	N209777	003	Apr 20,	2017

TABLET, EXTENDED RELEASE;ORAL

OXYCONTIN

+	PURDUE PHARMA LP	10MG	N022272	001	Apr 05,	2010
+		15MG	N022272	002	Apr 05,	2010
+		20MG	N022272	003	Apr 05,	2010
+		30MG	N022272	004	Apr 05,	2010
+		40MG	N022272	005	Apr 05,	2010
+		60MG	N022272	006	Apr 05,	2010
+		80MG	N022272	007	Apr 05,	2010

OXYMETAZOLINE HYDROCHLORIDE

CREAM;TOPICAL

RHOFADE

+	ALLERGAN INC	1%	N208552	001	Jan 18,	2017
---	--------------	----	---------	-----	---------	------

OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE

SPRAY, METERED;NASAL

KOVANAZE

+	ST RENATUS	0.1MG/SPRAY; 6MG/SPRAY	N208032	001	Jun 29,	2016
---	------------	------------------------	---------	-----	---------	------

OXYMETHOLONE

TABLET;ORAL

ANADROL-50

+	MYLAN SPECIALITY LP	50MG	N016848	001		
---	---------------------	------	---------	-----	--	--

OXYMORPHONE HYDROCHLORIDE

TABLET;ORAL

OPANA

<u>AB</u>	+	ENDO PHARMS	<u>5MG</u>	<u>N021611</u>	<u>001</u>	Jun 22,	2006
<u>AB</u>	+		<u>10MG</u>	<u>N021611</u>	<u>002</u>	Jun 22,	2006

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	AUROLIFE PHARMA LLC	<u>5MG</u>	<u>A204459</u>	<u>001</u>	Apr 26,	2016
<u>AB</u>		<u>10MG</u>	<u>A204459</u>	<u>002</u>	Apr 26,	2016
<u>AB</u>	AVANTHI INC	<u>5MG</u>	<u>A203601</u>	<u>001</u>	Jan 30,	2013
<u>AB</u>		<u>10MG</u>	<u>A203601</u>	<u>002</u>	Jan 30,	2013
<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A201187</u>	<u>001</u>	Dec 15,	2014
<u>AB</u>		<u>10MG</u>	<u>A201187</u>	<u>002</u>	Dec 15,	2014
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A202321</u>	<u>001</u>	Apr 25,	2013
<u>AB</u>		<u>10MG</u>	<u>A202321</u>	<u>002</u>	Apr 25,	2013
<u>AB</u>	TEVA	<u>5MG</u>	<u>A091443</u>	<u>002</u>	Feb 15,	2011
<u>AB</u>		<u>10MG</u>	<u>A091443</u>	<u>001</u>	Feb 15,	2011
<u>AB</u>	WEST-WARD PHARMS INT	<u>5MG</u>	<u>A090964</u>	<u>001</u>	Sep 27,	2010
<u>AB</u>		<u>10MG</u>	<u>A090964</u>	<u>002</u>	Sep 27,	2010

TABLET, EXTENDED RELEASE;ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>5MG</u>	<u>A079046</u>	<u>003</u>	Jul 11,	2013
<u>AB</u>		<u>7.5MG</u>	<u>A079046</u>	<u>001</u>	Dec 13,	2010
<u>AB</u>		<u>10MG</u>	<u>A079046</u>	<u>004</u>	Jul 11,	2013
<u>AB</u>		<u>15MG</u>	<u>A079046</u>	<u>002</u>	Dec 13,	2010
<u>AB</u>		<u>20MG</u>	<u>A079046</u>	<u>005</u>	Jul 11,	2013
<u>AB</u>		<u>30MG</u>	<u>A079046</u>	<u>006</u>	Jul 11,	2013
<u>AB</u>		<u>40MG</u>	<u>A079046</u>	<u>007</u>	Jul 11,	2013
<u>AB</u>	IMPAX LABS	<u>5MG</u>	<u>A079087</u>	<u>001</u>	Jun 14,	2010
<u>AB</u>		<u>7.5MG</u>	<u>A079087</u>	<u>002</u>	Dec 21,	2010
<u>AB</u>		<u>10MG</u>	<u>A079087</u>	<u>003</u>	Jun 14,	2010
<u>AB</u>		<u>15MG</u>	<u>A079087</u>	<u>004</u>	Dec 21,	2010
<u>AB</u>		<u>20MG</u>	<u>A079087</u>	<u>005</u>	Jun 14,	2010
<u>AB</u>		<u>30MG</u>	<u>A079087</u>	<u>006</u>	Jul 22,	2010
<u>AB</u>		<u>40MG</u>	<u>A079087</u>	<u>007</u>	Jun 14,	2010
<u>AB</u>	SPECGX LLC	<u>5MG</u>	<u>A202946</u>	<u>001</u>	Jun 27,	2014
<u>AB</u>		<u>7.5MG</u>	<u>A202946</u>	<u>002</u>	Jun 27,	2014
<u>AB</u>		<u>10MG</u>	<u>A202946</u>	<u>003</u>	Jun 27,	2014
<u>AB</u>		<u>15MG</u>	<u>A202946</u>	<u>004</u>	Jun 27,	2014
<u>AB</u>		<u>20MG</u>	<u>A202946</u>	<u>005</u>	Jun 27,	2014
<u>AB</u>		<u>30MG</u>	<u>A202946</u>	<u>006</u>	Jun 27,	2014
<u>AB</u>		<u>40MG</u>	<u>A202946</u>	<u>007</u>	Jun 27,	2014
<u>AB</u>	SUN PHARM INDs LTD	<u>5MG</u>	<u>A203506</u>	<u>001</u>	Apr 24,	2015
<u>AB</u>		<u>7.5MG</u>	<u>A203506</u>	<u>002</u>	Apr 24,	2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-330 (of 436)

OXYMORPHONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

OXYMORPHONE HYDROCHLORIDE

<u>AB</u>	<u>10MG</u>	<u>A203506 003</u>	Apr 24, 2015
<u>AB</u>	<u>15MG</u>	<u>A203506 004</u>	Apr 24, 2015
<u>AB</u>	<u>20MG</u>	<u>A203506 005</u>	Apr 24, 2015
<u>AB</u>	<u>30MG</u>	<u>A203506 006</u>	Apr 24, 2015
<u>AB</u>	<u>40MG</u>	<u>A203506 007</u>	Apr 24, 2015
<u>AB</u>	WEST-WARD PHARMS INT	<u>5MG</u>	<u>A200822 002</u> Jul 15, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A200822 003</u> Jul 15, 2013
<u>AB</u>		<u>10MG</u>	<u>A200822 004</u> Jul 15, 2013
<u>AB</u>		<u>15MG</u>	<u>A200822 005</u> Jul 15, 2013
<u>AB</u>		<u>20MG</u>	<u>A200822 006</u> Jul 15, 2013
<u>AB</u>		<u>30MG</u>	<u>A200822 007</u> Jul 15, 2013
<u>AB</u>		<u>40MG</u>	<u>A200822 001</u> Jul 15, 2013

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

TERRAMYCIN W/ POLYMYXIN B SULFATE

! CASPER PHARMA LLC EQ 5MG BASE/GM; 10,000 UNITS/GM

A061015 001

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN

<u>AP</u> +!	FRESENIUS KABI USA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018248 001</u>
<u>AP</u> +!		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018248 002</u>
<u>AP</u>	HIKMA FARMACEUTICA	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>A200219 001</u> Feb 13, 2013
<u>AP</u> +!	WEST-WARD PHARMS INT	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018243 001</u>
<u>AP</u> +!		<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018243 002</u> Jan 10, 2007
	PITOCIN		
<u>AP</u> +!	PAR STERILE PRODUCTS	<u>10USP UNITS/ML (10USP UNITS/ML)</u>	<u>N018261 001</u>
<u>AP</u> +	OXYTOCIN	<u>100USP UNITS/10ML (10USP UNITS/ML)</u>	<u>N018261 002</u> Jul 27, 2007
+!	FRESENIUS KABI USA	300USP UNITS/30ML (10USP UNITS/ML)	N018248 003 Jul 27, 2007
PITOCIN			
+	PAR STERILE PRODUCTS	500USP UNITS/50ML (10USP UNITS/ML)	N018261 003 Sep 05, 2012

OZENOXACIN

CREAM; TOPICAL

XEPI

+! FERRER INTERNACIONAL 1%

N208945 001 Dec 11, 2017

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

ABRAXANE

+! ABRAXIS BIOSCIENCE 100MG/VIAL

N021660 001 Jan 07, 2005

INJECTABLE; INJECTION

PACITAXEL

<u>AP</u>	GLAND PHARMA LTD	<u>6MG/ML</u>	<u>A207326 001</u> Aug 23, 2016
	<u>PACLITAXEL</u>		
<u>AP</u>	ACTAVIS TOTOWA	<u>6MG/ML</u>	<u>A090130 001</u> Dec 09, 2009
<u>AP</u>	FRESENIUS KABI ONCOL	<u>6MG/ML</u>	<u>A077574 001</u> Nov 27, 2006
<u>AP</u> !	HOSPIRA	<u>6MG/ML</u>	<u>A076131 001</u> May 08, 2002
<u>AP</u>	MYLAN LABS LTD	<u>6MG/ML</u>	<u>A091540 001</u> Sep 29, 2011
<u>AP</u>	SANDOZ INC	<u>6MG/ML</u>	<u>A078167 001</u> Dec 26, 2007
<u>AP</u>	TEVA PHARMS	<u>6MG/ML</u>	<u>A075184 001</u> Jan 25, 2002
<u>AP</u>	WEST-WARD PHARMS INT	<u>6MG/ML</u>	<u>A075190 001</u> Jan 28, 2002
	<u>TAXOL</u>		
<u>AP</u> +	HQ SPCLT PHARMA	<u>6MG/ML</u>	<u>N020262 001</u> Dec 29, 1992

PALBOCICLIB

CAPSULE; ORAL

IBRANCE

+ PFIZER INC 75MG  
+ 100MG  
+! 125MG

N207103 001 Feb 03, 2015  
N207103 002 Feb 03, 2015  
N207103 003 Feb 03, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-331 (of 436)

PALIPIPERIDONE

TABLET, EXTENDED RELEASE;ORAL

INVEGA

<u>AB</u>	+	JANSSEN PHARMS	<u>1.5MG</u>	<u>N021999 006</u>	Aug 26, 2008
<u>AB</u>	+		<u>3MG</u>	<u>N021999 001</u>	Dec 19, 2006
<u>AB</u>	+!		<u>6MG</u>	<u>N021999 002</u>	Dec 19, 2006
<u>AB</u>	+		<u>9MG</u>	<u>N021999 003</u>	Dec 19, 2006

PALIPIPERIDONE

<u>AB</u>		ACTAVIS LABS FL INC	<u>1.5MG</u>	<u>A202645 001</u>	Aug 03, 2015
<u>AB</u>			<u>3MG</u>	<u>A202645 002</u>	Aug 03, 2015
<u>AB</u>			<u>6MG</u>	<u>A202645 003</u>	Aug 03, 2015
<u>AB</u>			<u>9MG</u>	<u>A202645 004</u>	Aug 03, 2015
<u>AB</u>		MYLAN PHARMS INC	<u>1.5MG</u>	<u>A203802 001</u>	Sep 24, 2015
<u>AB</u>			<u>3MG</u>	<u>A203802 002</u>	Sep 24, 2015
<u>AB</u>			<u>6MG</u>	<u>A203802 003</u>	Sep 24, 2015
<u>AB</u>			<u>9MG</u>	<u>A203802 004</u>	Sep 24, 2015

PALIPIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

INVEGA SUSTENNA

+	JANSSEN PHARMS	39MG/0.25ML (39MG/0.25ML)	N022264 001	Jul 31, 2009
+		78MG/0.5ML (78MG/0.5ML)	N022264 002	Jul 31, 2009
+		117MG/0.75ML (117MG/0.75ML)	N022264 003	Jul 31, 2009
+!		156MG/ML (156MG/ML)	N022264 004	Jul 31, 2009
+		234MG/1.5ML (156MG/ML)	N022264 005	Jul 31, 2009

INVEGA TRINZA

+	JANSSEN PHARMS	273MG/0.875ML (273MG/0.875ML)	N207946 001	May 18, 2015
+		410MG/1.315ML (311.79MG/ML)	N207946 002	May 18, 2015
+		546MG/1.75ML (312MG/ML)	N207946 003	May 18, 2015
+!		819MG/2.625ML (312MG/ML)	N207946 004	May 18, 2015

PALONOSETRON HYDROCHLORIDE

INJECTABLE;INTRAVENOUS

ALOXI

<u>AP</u>	+!	HELSINN HLTHCARE	<u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u>	<u>N021372 002</u>	Feb 29, 2008
<u>AP</u>	+!		<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>N021372 001</u>	Jul 25, 2003

PALONOSETRON HYDROCHLORIDE

<u>AP</u>		DR REDDYS LABS LTD	<u>EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)</u>	<u>A201533 001</u>	Apr 21, 2016
<u>AP</u>			<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A201533 002</u>	Apr 21, 2016
<u>AP</u>			<u>EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)</u>	<u>A202521 001</u>	Oct 13, 2015

SOLUTION;INTRAVENOUS

PALONOSETRON HYDROCHLORIDE

EXELA PHARMA SCIENCE	EQ 0.25MG BASE/2ML (EQ 0.125MG BASE/ML)	N207963 001	Aug 22, 2016
+ FRESENIUS KABI USA	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N208109 001	Nov 21, 2017

PAMIDRONATE DISODIUM

INJECTABLE;INJECTION

PAMIDRONATE DISODIUM

<u>AP</u>		AREVA PHARMS	<u>30MG/VIAL</u>	<u>A077433 001</u>	Nov 26, 2008
<u>AP</u>			<u>90MG/VIAL</u>	<u>A077433 003</u>	Nov 26, 2008
<u>AP</u>		FRESENIUS KABI USA	<u>30MG/VIAL</u>	<u>A075773 001</u>	May 06, 2002
<u>AP</u>			<u>30MG/10ML (3MG/ML)</u>	<u>A076207 001</u>	May 17, 2002
<u>AP</u>			<u>90MG/VIAL</u>	<u>A075773 002</u>	May 06, 2002
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A076207 002</u>	May 17, 2002
<u>AP</u>	!	HOSPIRA	<u>30MG/10ML (3MG/ML)</u>	<u>A075841 001</u>	Jun 27, 2002
<u>AP</u>	!		<u>60MG/10ML (6MG/ML)</u>	<u>A075841 002</u>	Jun 27, 2002
<u>AP</u>	!		<u>90MG/10ML (9MG/ML)</u>	<u>A075841 003</u>	Jun 27, 2002
<u>AP</u>		LUITPOLD	<u>30MG/10ML (3MG/ML)</u>	<u>A078942 001</u>	Jul 25, 2008
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A078942 002</u>	Jul 25, 2008
<u>AP</u>		MYLAN LABS LTD	<u>30MG/10ML (3MG/ML)</u>	<u>A078520 001</u>	Oct 31, 2008
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A078520 002</u>	Oct 31, 2008
<u>AP</u>		PLIVA LACHEMA	<u>30MG/10ML (3MG/ML)</u>	<u>A078156 001</u>	Aug 19, 2008
<u>AP</u>			<u>60MG/10ML (6MG/ML)</u>	<u>A078156 002</u>	Aug 19, 2008
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A078156 003</u>	Aug 19, 2008
<u>AP</u>		SAGENT PHARMS	<u>30MG/10ML (3MG/ML)</u>	<u>A078373 001</u>	Dec 23, 2008
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A078373 002</u>	Dec 23, 2008
<u>AP</u>		SUN PHARMA GLOBAL	<u>30MG/VIAL</u>	<u>A077703 001</u>	Dec 24, 2008
<u>AP</u>			<u>90MG/VIAL</u>	<u>A077703 002</u>	Dec 24, 2008
<u>AP</u>		TEVA PHARMS USA	<u>30MG/10ML (3MG/ML)</u>	<u>A076153 001</u>	Mar 27, 2002
<u>AP</u>			<u>90MG/10ML (9MG/ML)</u>	<u>A076153 002</u>	Mar 27, 2002
<u>AP</u>		WEST-WARD PHARMS INT	<u>30MG/VIAL</u>	<u>A075290 001</u>	Apr 30, 2001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-332 (of 436)

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

**PAMIDRONATE DISODIUM**

<b>AP</b>	+!	<b>30MG/10ML (3MG/ML)</b>	<b>N021113 001</b>	Mar 04, 2002
<b>AP</b>		<b>90MG/VIAL</b>	<b>A075290 003</b>	Apr 30, 2001
<b>AP</b>	+!	<b>90MG/10ML (9MG/ML)</b>	<b>N021113 002</b>	Mar 04, 2002

AREVA PHARMS 60MG/VIAL A077433 002 Nov 26, 2008

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE, DELAYED RELEASE; ORAL

CREON

+	ABBVIE	60,000USP UNITS;12,000USP UNITS;38,000USP UNITS	N020725 002	Apr 30, 2009
+		15,000USP UNITS;3,000USP UNITS;9,500USP UNITS	N020725 004	Jul 12, 2011
+		30,000USP UNITS;6,000USP UNITS;19,000USP UNITS	N020725 001	Apr 30, 2009
+		180,000USP UNITS;36,000USP UNITS;114,000USP UNITS	N020725 005	Mar 14, 2013
+!		120,000USP UNITS;24,000USP UNITS;76,000USP UNITS	N020725 003	Apr 30, 2009

PANCREAZE

+	JANSSEN PHARMS	10,850USP UNITS;2,600USP UNITS;6,200USP UNITS	N022523 005	Mar 07, 2014
+		17,500USP/ UNITS;4,200USP/ UNITS;10,000USP/ UNITS	N022523 001	Apr 12, 2010
+		43,750USP/ UNITS;10,500USP/ UNITS;25,000USP/ UNITS	N022523 002	Apr 12, 2010
+!		61,000USP/ UNITS;21,000USP/ UNITS;37,000USP/ UNITS	N022523 003	Apr 12, 2010
+		70,000USP/ UNITS;16,800USP/ UNITS;40,000USP/ UNITS	N022523 004	Apr 12, 2010

PERTZYE

+	DIGESTIVE CARE INC	30,250USP UNITS;8,000USP UNITS;28,750USP UNITS	N022175 001	May 17, 2012
+!		60,500USP UNITS;16,000USP UNITS;57,500USP UNITS	N022175 002	May 17, 2012
+		15,125USP UNITS;4,000USP UNITS;14,375USP UNITS	N022175 003	Oct 06, 2016
+		90,750USP UNITS;24,000USP UNITS;86,250USP UNITS	N022175 004	Jul 13, 2017

ULTRESA

+	FOREST LABS INC	27,600USP UNITS;13,800USP UNITS;27,600USP UNITS	N022222 001	Mar 01, 2012
+		41,400USP UNITS;20,700USP UNITS;41,400USP UNITS	N022222 002	Mar 01, 2012
+!		46,000USP UNITS;23,000USP UNITS;46,000USP UNITS	N022222 003	Mar 01, 2012

ZENPEP

+	FOREST LABS INC	168,000USP UNITS;40,000USP UNITS;126,000USP UNITS	N022210 007	Mar 25, 2014
+		14,000USP UNITS;3,000USP UNITS;10,000USP UNITS	N022210 005	Jun 15, 2011
+		24,000USP UNITS;5,000USP UNITS;17,000USP UNITS	N022210 001	Aug 27, 2009
+		42,000USP UNITS;10,000USP UNITS;32,000USP UNITS	N022210 002	Aug 27, 2009
+		63,000USP UNITS;15,000USP UNITS;47,000USP UNITS	N022210 003	Aug 27, 2009
+		84,000USP UNITS;20,000USP UNITS;63,000USP UNITS	N022210 004	Aug 27, 2009
+!		105,000USP UNITS;25,000USP UNITS;79,000USP UNITS	N022210 006	Jul 13, 2011

TABLET; ORAL

VIOKACE

+	FOREST LABS INC	39,150USP UNITS;10,440USP UNITS;39,150USP UNITS	N022542 001	Mar 01, 2012
+!		78,300USP UNITS;20,880USP UNITS;78,300USP UNITS	N022542 002	Mar 01, 2012

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

**PANCURONIUM BROMIDE**

<b>AP</b>	HOSPIRA	<b>1MG/ML</b>	<b>A072320 001</b>	Jan 19, 1989
<b>AP</b>	! TEVA PHARMS USA	<b>1MG/ML</b>	<b>A072759 001</b>	Jul 31, 1990

!

2MG/ML A072760 001 Jul 31, 1990

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-333 (of 436)

PANOBINOSTAT LACTATE

CAPSULE;ORAL

FARYDAK

+ NOVARTIS PHARMS CORP	EQ 10MG BASE	N205353 001	Feb 23, 2015
+!	EQ 15MG BASE	N205353 002	Feb 23, 2015
	EQ 20MG BASE	N205353 003	Feb 23, 2015

PANTOPRAZOLE SODIUM

FOR SUSPENSION, DELAYED RELEASE;ORAL  
 PROTONIX

+! WYETH PHARMS INC	EQ 40MG BASE	N022020 001	Nov 14, 2007
INJECTABLE;IV (INFUSION)			

PANTOPRAZOLE SODIUM

<u>AP</u> AKORN INC	<u>EQ 40MG BASE/VIAL</u>	<u>A079197 001</u>	Nov 08, 2012
<u>AP</u> AUROBINDO PHARMA LTD	<u>EQ 40MG BASE/VIAL</u>	<u>A205675 001</u>	Mar 30, 2016
<u>AP</u> SANDOZ INC	<u>EQ 40MG BASE/VIAL</u>	<u>A090296 001</u>	Jul 14, 2015
<u>PROTONIX IV</u>			
<u>AP</u> +! WYETH PHARMS INC POWDER;IV (INFUSION)	<u>EQ 40MG BASE/VIAL</u>	<u>N020988 001</u>	Mar 22, 2001
PANTOPRAZOLE SODIUM EXELA PHARMA SCS LLC	EQ 40MG BASE/VIAL	N209463 001	Jun 30, 2017

TABLET, DELAYED RELEASE;ORAL

PANTOPRAZOLE SODIUM

<u>AB</u> ACTAVIS TOTOWA	<u>EQ 20MG BASE</u>	<u>A090797 001</u>	Feb 07, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A090797 002</u>	Feb 07, 2011
<u>AB</u> AMNEAL PHARMS	<u>EQ 20MG BASE</u>	<u>A205119 001</u>	Jan 26, 2016
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A205119 002</u>	Jan 26, 2016
<u>AB</u> APOTEX INC	<u>EQ 20MG BASE</u>	<u>A090807 001</u>	May 02, 2012
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A090807 002</u>	May 02, 2012
<u>AB</u> AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A202038 001</u>	Sep 28, 2012
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A202038 002</u>	Sep 28, 2012
<u>AB</u> DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A077619 001</u>	Jan 19, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A077619 002</u>	Jan 19, 2011
<u>AB</u> HETERO LABS LTD V	<u>EQ 20MG BASE</u>	<u>A202882 001</u>	Sep 10, 2014
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A202882 002</u>	Sep 10, 2014
<u>AB</u> JUBILANT GENERICS	<u>EQ 20MG BASE</u>	<u>A090901 001</u>	Aug 30, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A090901 002</u>	Aug 30, 2011
<u>AB</u> KREMERS URBAN PHARMS	<u>EQ 20MG BASE</u>	<u>A078281 001</u>	Jan 20, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A078281 002</u>	Jan 20, 2011
<u>AB</u> MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A200821 001</u>	Feb 16, 2012
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A200821 002</u>	Feb 16, 2012
<u>AB</u> MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A090970 001</u>	Jan 19, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A090970 002</u>	Jan 19, 2011
<u>AB</u> ORCHID HLTHCARE	<u>EQ 20MG BASE</u>	<u>A202052 001</u>	Dec 02, 2014
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A202052 002</u>	Dec 02, 2014
<u>AB</u> PERRIGO R AND D	<u>EQ 20MG BASE</u>	<u>A203024 001</u>	May 07, 2014
<u>AB</u> SUN PHARM INDs LTD	<u>EQ 20MG BASE</u>	<u>A200794 001</u>	May 02, 2012
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A200794 002</u>	May 02, 2012
<u>AB</u> TEVA	<u>EQ 20MG BASE</u>	<u>A077056 001</u>	Aug 02, 2007
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A077056 002</u>	Aug 02, 2007
<u>AB</u> TORRENT PHARMS	<u>EQ 20MG BASE</u>	<u>A090074 001</u>	Jan 19, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A090074 002</u>	Jan 19, 2011
<u>AB</u> WOCKHARDT	<u>EQ 20MG BASE</u>	<u>A091231 001</u>	Jan 19, 2011
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>A091231 002</u>	Jan 19, 2011
<u>PROTONIX</u>			
<u>AB</u> + WYETH PHARMS INC	<u>EQ 20MG BASE</u>	<u>N020987 002</u>	Jun 12, 2001
<u>AB</u> +!	<u>EQ 40MG BASE</u>	<u>N020987 001</u>	Feb 02, 2000

PARICALCITOL

CAPSULE;ORAL

PARICALCITOL

<u>AB</u> AMNEAL PHARMS	<u>1MCG</u>	<u>A204327 001</u>	Jan 13, 2016
<u>AB</u>	<u>2MCG</u>	<u>A204327 002</u>	Jan 13, 2016
<u>AB</u>	<u>4MCG</u>	<u>A204327 003</u>	Jan 13, 2016
<u>AB</u> AUROBINDO PHARMA LTD	<u>1MCG</u>	<u>A207672 001</u>	Jan 14, 2016
<u>AB</u>	<u>2MCG</u>	<u>A207672 002</u>	Jan 14, 2016
<u>AB</u>	<u>4MCG</u>	<u>A207672 003</u>	Jan 14, 2016
<u>AB</u> BIONPHARMA INC	<u>1MCG</u>	<u>A202539 001</u>	Mar 27, 2014
<u>AB</u>	<u>2MCG</u>	<u>A202539 002</u>	Mar 27, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-334 (of 436)

PARICALCITOL

CAPSULE;ORAL

PARICALCITOL

<u>AB</u>		<u>4MCG</u>	<u>A202539</u>	<u>003</u>	Mar 27, 2014
<u>AB</u>	DR REDDYS LABS LTD	<u>1MCG</u>	<u>A091412</u>	<u>001</u>	Jun 24, 2014
<u>AB</u>		<u>2MCG</u>	<u>A091412</u>	<u>002</u>	Jun 24, 2014
<u>AB</u>		<u>4MCG</u>	<u>A091412</u>	<u>003</u>	Jun 24, 2014
<u>AB</u>	ECI PHARMS LLC	<u>1MCG</u>	<u>A206710</u>	<u>001</u>	Feb 24, 2016
<u>AB</u>		<u>2MCG</u>	<u>A206710</u>	<u>002</u>	Feb 24, 2016
<u>AB</u>		<u>4MCG</u>	<u>A206710</u>	<u>003</u>	Feb 24, 2016
<u>AB</u>	MARKSANS PHARMA	<u>1MCG</u>	<u>A204948</u>	<u>001</u>	Oct 07, 2016
<u>AB</u>		<u>2MCG</u>	<u>A204948</u>	<u>002</u>	Oct 07, 2016
<u>AB</u>		<u>4MCG</u>	<u>A204948</u>	<u>003</u>	Oct 07, 2016
<u>AB</u>	RISING PHARMS INC	<u>1MCG</u>	<u>A202124</u>	<u>001</u>	Jun 24, 2014
<u>AB</u>		<u>2MCG</u>	<u>A202124</u>	<u>002</u>	Jun 24, 2014
<u>AB</u>		<u>4MCG</u>	<u>A202124</u>	<u>003</u>	Jun 24, 2014
<u>AB</u>	TEVA PHARMS USA	<u>1MCG</u>	<u>A090829</u>	<u>001</u>	Sep 27, 2013
<u>AB</u>		<u>2MCG</u>	<u>A090829</u>	<u>002</u>	Sep 27, 2013
<u>AB</u>	!	<u>4MCG</u>	<u>A090829</u>	<u>003</u>	Sep 27, 2013
<u>ZEMPLAR</u>					
<u>AB</u>	+ ABBVIE	<u>1MCG</u>	<u>N021606</u>	<u>001</u>	May 26, 2005
<u>AB</u>	+	<u>2MCG</u>	<u>N021606</u>	<u>002</u>	May 26, 2005
SOLUTION;INTRAVENOUS					
<u>PARICALCITOL</u>					
<u>AP</u>	ACCORD HLTHCARE	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N207174</u>	<u>001</u>	Feb 04, 2016
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N207174</u>	<u>002</u>	Feb 04, 2016
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N207174</u>	<u>003</u>	Feb 04, 2016
<u>AP</u>	AKORN	<u>0.005MG/ML (0.005MG/ML)</u>	<u>A207692</u>	<u>001</u>	Oct 16, 2017
<u>AP</u>	AMNEAL PHARMS CO	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A206699</u>	<u>001</u>	Mar 09, 2017
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A206699</u>	<u>002</u>	Mar 09, 2017
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A206699</u>	<u>003</u>	Mar 09, 2017
<u>AP</u>	DR REDDYS LABS LTD	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A204910</u>	<u>001</u>	Aug 17, 2016
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A204910</u>	<u>002</u>	Aug 17, 2016
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A204910</u>	<u>003</u>	Aug 17, 2016
<u>AP</u>	HIKMA PHARMS	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N205917</u>	<u>001</u>	Nov 18, 2014
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N205917</u>	<u>002</u>	Nov 18, 2014
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N205917</u>	<u>003</u>	Nov 18, 2014
<u>AP</u>	HOSPIRA INC	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N201657</u>	<u>001</u>	Oct 21, 2014
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>N201657</u>	<u>002</u>	Oct 21, 2014
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N201657</u>	<u>003</u>	Oct 21, 2014
<u>AP</u>	MYLAN LABS LTD	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A203897</u>	<u>001</u>	Nov 02, 2017
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A203897</u>	<u>002</u>	Nov 02, 2017
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A203897</u>	<u>003</u>	Nov 02, 2017
<u>AP</u>	SANDOZ INC	<u>0.002MG/ML (0.002MG/ML)</u>	<u>A091108</u>	<u>001</u>	Jul 27, 2011
<u>AP</u>		<u>0.005MG/ML (0.005MG/ML)</u>	<u>A091108</u>	<u>002</u>	Jul 27, 2011
<u>AP</u>		<u>0.01MG/2ML (0.005MG/ML)</u>	<u>A091108</u>	<u>003</u>	Jul 27, 2011
<u>ZEMPLAR</u>					
<u>AP</u>	+! ABBVIE	<u>0.002MG/ML (0.002MG/ML)</u>	<u>N020819</u>	<u>002</u>	Feb 01, 2000
<u>AP</u>	+	<u>0.005MG/ML (0.005MG/ML)</u>	<u>N020819</u>	<u>001</u>	Apr 17, 1998
<u>AP</u>	+	<u>0.01MG/2ML (0.005MG/ML)</u>	<u>N020819</u>	<u>003</u>	Feb 01, 2000
<u>PAROMOMYCIN SULFATE</u>					
CAPSULE;ORAL					
<u>PAROMOMYCIN SULFATE</u>					
<u>AA</u>	HERITAGE PHARMS INC	<u>EQ 250MG BASE</u>	<u>A065173</u>	<u>001</u>	Dec 14, 2007
<u>AA</u>	! SUN PHARM INDNS INC	<u>EQ 250MG BASE</u>	<u>A064171</u>	<u>001</u>	Jun 30, 1997
<u>PAROXETINE HYDROCHLORIDE</u>					
SUSPENSION;ORAL					
PAXIL					
	+! APOTEX TECHNOLOGIES	EQ 10MG BASE/5ML			
TABLET;ORAL					
<u>PAROXETINE</u>					
<u>AB</u>	PRINSTON INC	<u>EQ 10MG BASE</u>	<u>A203854</u>	<u>001</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203854</u>	<u>002</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203854</u>	<u>003</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203854</u>	<u>004</u>	Oct 31, 2014
<u>PAROXETINE HYDROCHLORIDE</u>					
<u>AB</u>	APOTEX	<u>EQ 10MG BASE</u>	<u>A075356</u>	<u>001</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A075356</u>	<u>002</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A075356</u>	<u>003</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A075356</u>	<u>004</u>	Jul 30, 2003
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078406</u>	<u>001</u>	Jul 25, 2007

PAROMOMYCIN SULFATE

CAPSULE;ORAL

PAROMOMYCIN SULFATE

<u>AA</u>	HERITAGE PHARMS INC	<u>EQ 250MG BASE</u>	<u>A065173</u>	<u>001</u>	Dec 14, 2007
<u>AA</u>	! SUN PHARM INDNS INC	<u>EQ 250MG BASE</u>	<u>A064171</u>	<u>001</u>	Jun 30, 1997

PAROXETINE HYDROCHLORIDE

SUSPENSION;ORAL

PAXIL

+! APOTEX TECHNOLOGIES EQ 10MG BASE/5ML

N020710 001 Jun 25, 1997

TABLET;ORAL

PAROXETINE

<u>AB</u>	PRINSTON INC	<u>EQ 10MG BASE</u>	<u>A203854</u>	<u>001</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A203854</u>	<u>002</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A203854</u>	<u>003</u>	Oct 31, 2014
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A203854</u>	<u>004</u>	Oct 31, 2014

PAROXETINE HYDROCHLORIDE

TABLET;ORAL

PAROXETINE

<u>AB</u>	APOTEX	<u>EQ 10MG BASE</u>	<u>A075356</u>	<u>001</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A075356</u>	<u>002</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A075356</u>	<u>003</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A075356</u>	<u>004</u>	Jul 30, 2003
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 10MG BASE</u>	<u>A078406</u>	<u>001</u>	Jul 25, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-335 (of 436)

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078406 002</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078406 003</u>	Jul 25, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078406 004</u>	Jul 25, 2007
<u>AB</u>	JUBILANT GENERICS	<u>EQ 10MG BASE</u>	<u>A205528 001</u>	Nov 27, 2015
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205528 002</u>	Nov 27, 2015
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A205528 003</u>	Nov 27, 2015
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205528 004</u>	Nov 27, 2015
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>A078902 001</u>	Mar 13, 2008
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078902 002</u>	Mar 13, 2008
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078902 003</u>	Mar 13, 2008
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078902 004</u>	Mar 13, 2008
<u>AB</u>	OXFORD PHARMS	<u>EQ 10MG BASE</u>	<u>A076968 001</u>	Jun 21, 2010
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076968 002</u>	Jun 21, 2010
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A076968 003</u>	Jun 21, 2010
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076968 004</u>	Jun 21, 2010
<u>AB</u>	SUN PHARM INDNS INC	<u>EQ 10MG BASE</u>	<u>A078194 001</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A078194 002</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A078194 003</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A078194 004</u>	Jun 29, 2007
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>A076618 001</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076618 002</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A076618 003</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076618 004</u>	Aug 15, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 10MG BASE</u>	<u>A077584 001</u>	Mar 07, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A077584 002</u>	Mar 07, 2007
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>A077584 003</u>	Mar 07, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A077584 004</u>	Mar 07, 2007

PAXIL

<u>AB</u>	+	APOTEX TECHNOLOGIES	<u>EQ 10MG BASE</u>	<u>N020031 001</u>	Dec 29, 1992
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N020031 002</u>	Dec 29, 1992
<u>AB</u>	+		<u>EQ 30MG BASE</u>	<u>N020031 003</u>	Dec 29, 1992
<u>AB</u>	!+		<u>EQ 40MG BASE</u>	<u>N020031 005</u>	Dec 29, 1992

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>	KREMERS URBAN PHARMS	<u>EQ 12.5MG BASE</u>	<u>A204744 001</u>	Jun 10, 2016
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204744 002</u>	Jun 10, 2016
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A204744 003</u>	Jun 10, 2016
<u>AB</u>	LUPIN LTD	<u>EQ 12.5MG BASE</u>	<u>A204134 001</u>	Jan 20, 2017
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A204134 002</u>	Jan 20, 2017
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A204134 003</u>	Jan 20, 2017
<u>AB</u>	MYLAN	<u>EQ 12.5MG BASE</u>	<u>A077873 001</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>A077873 002</u>	Jun 29, 2007
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A091427 001</u>	Apr 14, 2011

PAXIL CR

<u>AB</u>	+	APOTEX TECHNOLOGIES	<u>EQ 12.5MG BASE</u>	<u>N020936 001</u>	Feb 16, 1999
<u>AB</u>	+		<u>EQ 25MG BASE</u>	<u>N020936 002</u>	Feb 16, 1999
<u>AB</u>	!+		<u>EQ 37.5MG BASE</u>	<u>N020936 003</u>	Dec 06, 2000

PAROXETINE MESYLATE

CAPSULE; ORAL

BRISDELLE

<u>AB</u>	!+ SEBELA IRELAND LTD	<u>EQ 7.5MG BASE</u>	<u>N204516 001</u>	Jun 28, 2013
<u>AB</u>		<u>PAROXETINE MESYLATE</u>		
<u>AB</u>	ACTAVIS LABS FL INC	<u>EQ 7.5MG BASE</u>	<u>A207139 001</u>	Jun 20, 2017

TABLET; ORAL

PEXEVA

+	SEBELA IRELAND LTD	<u>EQ 10MG BASE</u>	N021299 001	Jul 03, 2003
+		<u>EQ 20MG BASE</u>	N021299 002	Jul 03, 2003
+		<u>EQ 30MG BASE</u>	N021299 003	Jul 03, 2003
!+		<u>EQ 40MG BASE</u>	N021299 004	Jul 03, 2003

PASIRETOIDE DIASPARTATE

SOLUTION; SUBCUTANEOUS

SIGNIFOR

+	NOVARTIS	<u>EQ 0.3MG BASE/ML (EQ 0.3MG BASE/ML)</u>	N200677 001	Dec 14, 2012
+		<u>EQ 0.6MG BASE/ML (EQ 0.6MG BASE/ML)</u>	N200677 002	Dec 14, 2012
!+		<u>EQ 0.9MG BASE/ML (EQ 0.9MG BASE/ML)</u>	N200677 003	Dec 14, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-336 (of 436)

**PASIREOTIDE PAMOATE**

POWDER;INTRAMUSCULAR

SIGNIFOR LAR

+ NOVARTIS PHARMS CORP	EQ 20MG BASE/VIAL	N203255 001 Dec 15, 2014
+!	EQ 40MG BASE/VIAL	N203255 002 Dec 15, 2014
+!	EQ 60MG BASE/VIAL	N203255 003 Dec 15, 2014

**PATIROMER SORBITEX CALCIUM**

POWDER;ORAL

VELTASSA

+ RELYPSA INC	EQ 8.4GM BASE/PACKET	N205739 001 Oct 21, 2015
+	EQ 16.8GM BASE/PACKET	N205739 002 Oct 21, 2015
+!	EQ 25.2GM BASE/PACKET	N205739 003 Oct 21, 2015

**PAZOPANIB HYDROCHLORIDE**

TABLET;ORAL

VOTRIENT

+! NOVARTIS PHARMS CORP	EQ 200MG BASE	N022465 001 Oct 19, 2009
-------------------------	---------------	--------------------------

**PEGADEMASE BOVINE**

INJECTABLE;INJECTION

ADAGEN

+! SIGMA TAU	250 UNITS/ML	N019818 001 Mar 21, 1990
--------------	--------------	--------------------------

**PEGAPTANIB SODIUM**

INJECTABLE;INTRAVITREAL

MACUGEN

+! VALEANT PHARMS LLC	EQ 0.3MG ACID/0.09ML	N021756 001 Dec 17, 2004
-----------------------	----------------------	--------------------------

**PEGVISOMANT**

INJECTABLE;SUBCUTANEOUS

SOMAVERT

+! PHARMACIA AND UPJOHN	10MG/VIAL	N021106 001 Mar 25, 2003
+!	15MG/VIAL	N021106 002 Mar 25, 2003
+!	20MG/VIAL	N021106 003 Mar 25, 2003
+!	25MG/VIAL	N021106 004 Jul 31, 2014
+!	30MG/VIAL	N021106 005 Jul 31, 2014

**PEMETREXED DISODIUM**

INJECTABLE;IV (INFUSION)

ALIMTA

+! LILLY	EQ 100MG BASE/VIAL	N021462 002 Sep 07, 2007
+!	EQ 500MG BASE/VIAL	N021462 001 Feb 04, 2004

**PENCICLOVIR**

CREAM;TOPICAL

DENAVIR

+! MYLAN PHARMS INC	1%	N020629 001 Sep 24, 1996
---------------------	----	--------------------------

**PENICILLAMINE**

CAPSULE;ORAL

CUPRIMINE

+! ATON	250MG	N019853 001
---------	-------	-------------

TABLET;ORAL

DEPEN

+! MYLAN SPECIALITY LP	250MG	N019854 001
------------------------	-------	-------------

**PENICILLIN G BENZATHINE**

INJECTABLE;INJECTION

BICILLIN L-A

BC +! KING PHARMS LLC	600,000 UNITS/ML	N050141 001
-----------------------	------------------	-------------

PERMAPEN

BC CASPER PHARMA LLC	600,000 UNITS/ML	A060014 001
----------------------	------------------	-------------

BICILLIN L-A

+! KING PHARMS LLC	300,000 UNITS/ML	N050141 003
--------------------	------------------	-------------

**PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE**

INJECTABLE;INJECTION

BICILLIN C-R

+! KING PHARMS LLC	150,000 UNITS/ML;150,000 UNITS/ML	N050138 002
--------------------	-----------------------------------	-------------

+!	300,000 UNITS/ML;300,000 UNITS/ML	N050138 001
----	-----------------------------------	-------------

BICILLIN C-R 900/300

+! KING PHARMS LLC	900,000 UNITS/2ML;300,000 UNITS/2ML	N050138 003
--------------------	-------------------------------------	-------------



38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-338 (of 436)

<u>PENTETATE ZINC TRISODIUM</u>			
SOLUTION;INHALATION, INTRAVENOUS	PENTETATE ZINC TRISODIUM	+! HAMELN PHARMA PLUS	EQ 1GM BASE/5ML (EQ 200MG BASE/ML) N021751 001 Aug 11, 2004
<u>PENTOBARBITAL SODIUM</u>			
INJECTABLE;INJECTION	NEMBUTAL SODIUM	AP ! OAK PHARMS	<u>50MG/ML</u> A083246 001
<u>PENTOBARBITAL SODIUM</u>			
AP CUSTOPHARM INC	<u>50MG/ML</u>	A203619 001	Nov 13, 2017
AP RENAISSANCE SSA LLC	<u>50MG/ML</u>	A206677 001	Nov 27, 2017
AP SAGENT PHARMS	<u>50MG/ML</u>	A206404 001	May 23, 2016
<u>PENTOSAN POLYSULFATE SODIUM</u>			
CAPSULE;ORAL	ELMIRON	+! JANSSEN PHARMS	100MG N020193 001 Sep 26, 1996
<u>PENTOSTATIN</u>			
INJECTABLE;INJECTION	NIPENT	AP +! HOSPIRA INC	<u>10MG/VIAL</u> N020122 001 Oct 11, 1991
<u>PENTOSTATIN</u>			
AP MYLAN INSTITUTIONAL	<u>10MG/VIAL</u>	A203554 001	Sep 19, 2014
AP WEST-WARD PHARMS INT	<u>10MG/VIAL</u>	A077841 001	Aug 07, 2007
<u>PENTOXIFYLLINE</u>			
TABLET, EXTENDED RELEASE;ORAL	PENTOXIFYLLINE	AB ! APOTEX	<u>400MG</u> A075191 001 Jun 09, 1999
AB MYLAN	<u>400MG</u>	A074425 001	Jul 08, 1997
AB VALEANT PHARMS	<u>400MG</u>	A075028 001	Jul 20, 1998
<u>PENTOXIL</u>			
AB UPSHER-SMITH LABS	<u>400MG</u>	A074962 001	Mar 31, 1999
<u>PERAMIVIR</u>			
SOLUTION;IV (INFUSION)	RAPIVAB	+! BIOCRYST	200MG/20ML (10MG/ML) N206426 001 Dec 19, 2014
<u>PERAMPANEL</u>			
SUSPENSION;ORAL	FYCOMPA	+! EISAI INC	0.5MG/ML N208277 001 Apr 29, 2016
TABLET;ORAL			
FYCOMPA	+ EISAI INC	2MG	N202834 001 Oct 22, 2012
	+	4MG	N202834 002 Oct 22, 2012
	+	6MG	N202834 003 Oct 22, 2012
	+	8MG	N202834 004 Oct 22, 2012
	+	10MG	N202834 005 Oct 22, 2012
	+!	12MG	N202834 006 Oct 22, 2012
<u>PERFLUTREN</u>			
INJECTABLE;INTRAVENOUS	DEFINITY	+! LANTHEUS MEDCL	6.52MG/ML N021064 001 Jul 31, 2001
<u>PERINDOPRIL ERBUMINE</u>			
TABLET;ORAL	PERINDOPRIL ERBUMINE	AB ANI PHARMS INC	<u>2MG</u> A078138 001 Nov 10, 2009
		AB	<u>4MG</u> A078138 002 Nov 10, 2009
		AB	<u>8MG</u> A078138 003 Nov 10, 2009
	APOTEX	AB	<u>2MG</u> A090463 001 Aug 30, 2010
		AB	<u>4MG</u> A090463 002 Aug 30, 2010
		AB	<u>8MG</u> A090463 003 Aug 30, 2010
	! AUROBINDO PHARMA	AB	<u>2MG</u> A079070 001 Nov 10, 2009
		AB	<u>4MG</u> A079070 002 Nov 10, 2009
		AB	<u>8MG</u> A079070 003 Nov 10, 2009
	WEST-WARD PHARMS INT	AB	<u>2MG</u> A090072 001 Nov 10, 2009
		AB	<u>4MG</u> A090072 002 Nov 10, 2009
		AB	<u>8MG</u> A090072 003 Nov 10, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-339 (of 436)

PERMETHRIN

CREAM;TOPICAL

ELIMITE

AB +! MYLAN PHARMS INC 5%

N019855 001 Aug 25, 1989

PERMETHRIN

AB ACTAVIS LABS 5%

A074806 001 Jan 23, 1998

AB PERRIGO NEW YORK 5%

A076369 001 Apr 21, 2003

PERPHENAZINE

TABLET;ORAL

PERPHENAZINE

AB MYLAN PHARMS INC 2MG

A206691 001 Apr 14, 2017

AB 4MG

A206691 002 Apr 14, 2017

AB 8MG

A206691 003 Apr 14, 2017

AB 16MG

A206691 004 Apr 14, 2017

AB SANDOZ 2MG

A089685 002 Dec 08, 1988

AB 4MG

A089685 003 Dec 08, 1988

AB 8MG

A089685 001 Dec 08, 1988

AB ! 16MG

A089685 004 Dec 08, 1988

AB VINTAGE PHARMS 2MG

A040226 001 Dec 31, 1998

AB 4MG

A040226 002 Dec 31, 1998

AB 8MG

A040226 003 Dec 31, 1998

AB 16MG

A040226 004 Dec 31, 1998

AB WATSON LABS INC 2MG

A207582 001 Oct 17, 2016

AB 4MG

A207582 002 Oct 17, 2016

AB 8MG

A207582 003 Oct 17, 2016

AB 16MG

A207582 004 Oct 17, 2016

AB WILSHIRE PHARMS INC 2MG

A205973 001 Dec 17, 2015

AB 4MG

A205973 002 Dec 17, 2015

AB 8MG

A205973 003 Dec 17, 2015

AB 16MG

A205973 004 Dec 17, 2015

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

PHENDIMETRAZINE TARTRATE

+! SANDOZ 105MG

N018074 001

TABLET;ORAL

BONTRIL PDM

AA ! VALEANT 35MG

A085272 001

PHENDIMETRAZINE TARTRATE

AA ELITE LABS INC 35MG

A040762 001 Jan 28, 2008

AA 35MG

A203600 001 Dec 27, 2017

AA KVK TECH 35MG

A091042 001 Aug 31, 2010

AA MIKART 35MG

A089452 001 Oct 30, 1991

AA SANDOZ 35MG

A085588 001

PHENELZINE SULFATE

TABLET;ORAL

NARDIL

AB +! PARKE DAVIS EQ 15MG BASE

N011909 002

PHENELZINE SULFATE

AB NOVEL LABS INC EQ 15MG BASE

A200181 001 Dec 08, 2010

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE;ORAL

DIBENZYLINE

AB +! CONCORDIA PHARMS 10MG

N008708 001

INC

PHENOXYBENZAMINE HYDROCHLORIDE

AB PAR PHARM INC 10MG

A204522 001 Jan 24, 2017

AB WEST-WARD PHARMS 10MG

A201050 001 Jul 16, 2012

INT

PHENTERMINE HYDROCHLORIDE

CAPSULE;ORAL

ADIPEX-P

AA ! TEVA 37.5MG

A088023 001 Aug 02, 1983

PHENTERMINE HYDROCHLORIDE

AA AUROLIFE PHARMA LLC 15MG

A204318 001 Nov 09, 2016

AA 30MG

A204318 002 Nov 09, 2016

AA BARR 15MG

A090591 001 Mar 18, 2010

AA 30MG

A090591 002 Mar 18, 2010

AA ELITE LABS 15MG

A202248 001 Sep 28, 2012

AA 30MG

A202248 002 Sep 28, 2012

AA ELITE LABS INC 37.5MG

A040228 001 Jun 19, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-340 (of 436)

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

<u>AA</u>	INVAGEN PHARMS	<u>15MG</u>	<u>A202858 001</u>	Feb 14, 2014
<u>AA</u>		<u>30MG</u>	<u>A202858 002</u>	Feb 14, 2014
<u>AA</u>		<u>30MG</u>	<u>A204414 001</u>	May 05, 2014
<u>AA</u>		<u>37.5MG</u>	<u>A202846 001</u>	Feb 05, 2014
<u>AA</u>	KEN LIFESCIENCE	<u>15MG</u>	<u>A205019 001</u>	Dec 05, 2014
<u>AA</u>		<u>30MG</u>	<u>A205019 002</u>	Dec 05, 2014
<u>AA</u>		<u>37.5MG</u>	<u>A205017 001</u>	Sep 25, 2014
<u>AA</u>	KVK TECH	<u>15MG</u>	<u>A040886 002</u>	Mar 31, 2008
<u>AA</u>		<u>30MG</u>	<u>A040875 001</u>	Mar 21, 2008
<u>AA</u>		<u>30MG</u>	<u>A040886 001</u>	Mar 31, 2008
<u>AA</u>		<u>37.5MG</u>	<u>A040887 001</u>	Apr 24, 2008
<u>AA</u>	LANNETT	<u>15MG</u>	<u>A087022 002</u>	Jan 20, 2012
<u>AA</u>		<u>30MG</u>	<u>A087022 001</u>	Feb 03, 1983
<u>AA</u>		<u>30MG</u>	<u>A091359 001</u>	Jul 16, 2010
<u>AA</u>	LANNETT HOLDINGS INC	<u>37.5MG</u>	<u>A201961 001</u>	Jul 20, 2011
<u>AA</u> !	SANDOZ	<u>15MG</u>	<u>A087190 002</u>	
<u>AA</u> !		<u>30MG</u>	<u>A086945 001</u>	Jul 20, 1983
<u>AA</u> !		<u>30MG</u>	<u>A087190 001</u>	
<u>AA</u>	SUN PHARM INDUSTRIES	<u>30MG</u>	<u>A040525 001</u>	Oct 23, 2003

TABLET; ORAL

ADIPEX-P

<u>AA</u> !	TEVA	<u>37.5MG</u>	<u>A085128 001</u>	
	<u>LOMAIR</u>			
<u>AA</u> !	AVANTHI INC	<u>8MG</u>	<u>A203495 001</u>	Sep 12, 2016
	<u>PHENTERMINE HYDROCHLORIDE</u>			
<u>AA</u>	AUROLIFE PHARMA LLC	<u>37.5MG</u>	<u>A203068 001</u>	Aug 06, 2014
<u>AA</u>	BARR	<u>37.5MG</u>	<u>A090470 001</u>	Aug 31, 2009
<u>AA</u>	ELITE LABS	<u>37.5MG</u>	<u>A200272 001</u>	Jan 31, 2011
<u>AA</u>	ELITE LABS INC	<u>37.5MG</u>	<u>A040190 001</u>	May 30, 1997
<u>AA</u>	INGENUS PHARMS NJ	<u>37.5MG</u>	<u>A091451 001</u>	Sep 21, 2012
<u>AA</u>	INVAGEN PHARMS	<u>37.5MG</u>	<u>A202942 001</u>	Feb 05, 2014
<u>AA</u>	KEN LIFESCIENCE	<u>37.5MG</u>	<u>A205008 001</u>	Sep 25, 2014
<u>AA</u>	KVK TECH	<u>37.5MG</u>	<u>A040876 001</u>	Mar 31, 2008
<u>AA</u>	KVK TECH INC	<u>8MG</u>	<u>A203436 001</u>	Mar 17, 2017
<u>AA</u>	LANNETT	<u>37.5MG</u>	<u>A040555 001</u>	Apr 15, 2005
<u>AA</u>	POLYGEN PHARMS	<u>37.5MG</u>	<u>A206342 001</u>	Nov 18, 2016
<u>AA</u>	PRINSTON INC	<u>37.5MG</u>	<u>A040377 001</u>	Jan 04, 2002
<u>AA</u>	SUN PHARM INDS INC	<u>37.5MG</u>	<u>A040790 001</u>	Aug 21, 2007
<u>AA</u>	SUN PHARM INDUSTRIES	<u>37.5MG</u>	<u>A040526 001</u>	Oct 23, 2003

TABLET, ORALLY DISINTEGRATING; ORAL  
 PHENTERMINE HYDROCHLORIDE

ZYDUS PHARMS USA INC	15MG	A204663 001	Jun 28, 2017
	30MG	A204663 002	Jun 28, 2017
	37.5MG	A204663 003	Jun 28, 2017

PHENTERMINE HYDROCHLORIDE; TOPIRAMATE

CAPSULE, EXTENDED RELEASE; ORAL

QSYMIA

+	VIVUS	EQ 3.75MG BASE; 23MG	N022580 001	Jul 17, 2012
+		EQ 7.5MG BASE; 46MG	N022580 002	Jul 17, 2012
+		EQ 11.25MG BASE; 69MG	N022580 003	Jul 17, 2012
++!		EQ 15MG BASE; 92MG	N022580 004	Jul 17, 2012

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

<u>AP</u>	PRECISION DOSE INC	<u>5MG/VIAL</u>	<u>A207686 001</u>	Jul 14, 2017
<u>AP</u> !	WEST-WARD PHARMS INT	<u>5MG/VIAL</u>	<u>A040235 001</u>	Mar 11, 1998
	ORAVERSE			
+!	SEPTODONT HOLDING	0.4MG/1.7ML	N022159 001	May 09, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-341 (of 436)

PHENYLEPHRINE HYDROCHLORIDE

SOLUTION;IV (INFUSION)

PHENYLEPHRINE HYDROCHLORIDE

+! WEST WARD PHARM CORP

VAZCULEP

+ AVADEL LEGACY 10MG/ML (10MG/ML)  
+ 50MG/5ML (10MG/ML)  
+! 100MG/10ML (10MG/ML)

N203826 001 Dec 20, 2012

N204300 001 Jun 27, 2014  
N204300 002 Jun 27, 2014  
N204300 003 Jun 27, 2014

SOLUTION/DROPS;OPHTHALMIC

PHENYLEPHRINE HYDROCHLORIDE

+! AKORN INC 2.5%  
+! 10%  
+! PARAGON BIOTECK 2.5%  
+! 10%

N207926 001 Jan 15, 2015  
N207926 002 Jan 15, 2015  
N203510 001 Mar 21, 2013  
N203510 002 Mar 21, 2013

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

AA HI-TECH PHARMACAL 5MG/5ML;6.25MG/5ML

A040675 001 Dec 23, 2014

AA VINTAGE 5MG/5ML;6.25MG/5ML

A040654 001 Dec 07, 2006

PROMETH VC PLAIN

AA ! G AND W LABS INC 5MG/5ML;6.25MG/5ML

A088761 001 Nov 08, 1984

PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE

AA AMNEAL PHARMS 5MG/5ML;6.25MG/5ML

A040902 001 Aug 25, 2009

PHENYTOIN

SUSPENSION;ORAL

DILANTIN-125

AB +! PARKE DAVIS 125MG/5ML

N008762 001

PHENYTOIN

AB TARO 125MG/5ML

A040521 001 Mar 08, 2004

AB VISTAPHARM 125MG/5ML

A040342 001 Jan 31, 2001

AB 125MG/5ML

A040610 001 Aug 18, 2005

AB WOCKHARDT EU OPERATN 125MG/5ML

A040420 001 Apr 19, 2002

TABLET, CHEWABLE;ORAL

DILANTIN

AB ! PFIZER 50MG

A084427 001

PHENYTOIN

AB EPIC PHARMA LLC 50MG

A040884 001 Nov 28, 2014

AB MYLAN PHARMS INC 50MG

A200691 001 Dec 26, 2012

AB TARO 50MG

A200565 001 Apr 17, 2014

PHENYTOIN SODIUM

CAPSULE;ORAL

DILANTIN

AB ! PARKE-DAVIS 100MG EXTENDED

A084349 002

EXTENDED PHENYTOIN SODIUM

AB AMNEAL PHARMS NY 100MG EXTENDED

A040765 001 Nov 12, 2008

AB MYLAN 100MG EXTENDED

A040298 001 Dec 28, 1998

AB SUN PHARM INDs 200MG EXTENDED

A040731 001 Jun 30, 2008

AB 300MG EXTENDED

A040731 002 Jun 30, 2008

AB SUN PHARM INDs (IN) 100MG EXTENDED

A040621 001 Dec 11, 2006

AB TARO 100MG EXTENDED

A040684 001 Sep 05, 2006

PHENYTEK

AB MYLAN 200MG EXTENDED

A040298 002 Dec 06, 2001

AB ! 300MG EXTENDED

A040298 003 Dec 06, 2001

PHENYTOIN SODIUM

AB AUROBINDO PHARMA LTD 100MG EXTENDED

A204309 001 Jun 10, 2015

DILANTIN

! PARKE-DAVIS 30MG EXTENDED

A084349 001

INJECTABLE;INJECTION

PHENYTOIN SODIUM

AP ACELLA PHARMS LLC 50MG/ML

A040573 001 Sep 13, 2006

AP LUITPOLD 50MG/ML

A040781 001 Dec 04, 2007

AP ! WEST-WARD PHARMS INT 50MG/ML

A084307 001

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-342 (of 436)

PHYTONADIONE

INJECTABLE; INJECTION  
 PHYTONADIONE  
 BP ! INTL MEDICATION 1MG/0.5ML A083722 001  
 VITAMIN K1  
 BP ! HOSPIRA 1MG/0.5ML A087954 001 Jul 25, 1983  
 ! 10MG/ML A087955 001 Jul 25, 1983  
 TABLET; ORAL  
 MEPHYTON  
 +! VALEANT PHARMS 5MG N010104 003

PILOCARPINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC  
IISOPTO CARPINE  
AT + NOVARTIS PHARMS 1% N200890 001 Jun 22, 2010  
 CORP  
AT + 2% N200890 002 Jun 22, 2010  
AT +! 4% N200890 003 Jun 22, 2010  
PILOCARPINE HYDROCHLORIDE  
AT AKORN INC 1% A204398 001 Sep 27, 2017  
AT 2% A204398 002 Sep 27, 2017  
AT 4% A204398 003 Sep 27, 2017

TABLET; ORAL  
PILOCARPINE HYDROCHLORIDE  
AB IMPAX LABS 5MG A077248 001 Mar 31, 2006  
AB 7.5MG A077248 002 Mar 31, 2006  
AB INNOGENIX 5MG A076963 001 Dec 22, 2004  
AB 7.5MG A076963 002 Feb 27, 2007  
AB LANNETT 5MG A077220 001 Oct 14, 2005  
AB 7.5MG A077220 002 May 06, 2009  
AB PERRIGO PHARMA INTL 5MG A076746 001 Nov 16, 2004  
SALAGEN  
AB + EISAI INC 5MG N020237 001 Mar 22, 1994  
AB +! 7.5MG N020237 002 Apr 18, 2003

PIMAVANSERIN TARTRATE

TABLET; ORAL  
 NUPLAZID  
 +! ACADIA PHARMS INC EQ 17MG BASE N207318 001 Apr 29, 2016

PIMECROLIMUS

CREAM; TOPICAL  
 ELIDEL  
 +! VALEANT BERMUDA 1% N021302 001 Dec 13, 2001

PIMOZIDE

TABLET; ORAL  
ORAP  
AB + TEVA 1MG N017473 003 Aug 27, 1997  
AB +! 2MG N017473 001 Jul 31, 1984  
PIMOZIDE  
AB PAR PHARM 1MG A204521 001 Sep 28, 2015  
AB 2MG A204521 002 Sep 28, 2015

PINDOLOL

TABLET; ORAL  
PINDOLOL  
AB IDT AUSTRALIA LTD 5MG A073608 001 Mar 29, 1993  
AB 10MG A073609 001 Mar 29, 1993  
AB MYLAN PHARMS INC 5MG A074019 001 Sep 03, 1992  
AB ! 10MG A074019 002 Sep 03, 1992  
AB NOSTRUM LABS INC 5MG A205415 001 Jan 13, 2016  
AB 10MG A205415 002 Jan 13, 2016  
AB SUN PHARM INDUSTRIES 5MG A074063 001 Jan 27, 1994  
AB 10MG A074063 002 Jan 27, 1994  
AB ZYDUS PHARMS USA 5MG A209866 001 Aug 18, 2017  
AB INC 10MG A209866 002 Aug 18, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-343 (of 436)

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

**ACTOS**

<b>AB</b>	+	TAKEDA PHARMS USA	<u>EQ 15MG BASE</u>	<b>N021073 001</b>	Jul 15, 1999
<b>AB</b>	+		<u>EQ 30MG BASE</u>	<b>N021073 002</b>	Jul 15, 1999
<b>AB</b>	+!		<u>EQ 45MG BASE</u>	<b>N021073 003</b>	Jul 15, 1999
<b>PIOGLITAZONE HYDROCHLORIDE</b>					
<b>AB</b>		ACCORD HLTHCARE	<u>EQ 15MG BASE</u>	<b>A200044 001</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A200044 002</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A200044 003</b>	Feb 13, 2013
<b>AB</b>		AUROBINDO PHARMA LTD	<u>EQ 15MG BASE</u>	<b>A200268 001</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A200268 002</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A200268 003</b>	Feb 13, 2013
<b>AB</b>		BRECKENRIDGE PHARM	<u>EQ 15MG BASE</u>	<b>A078472 001</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A078472 002</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A078472 003</b>	Feb 13, 2013
<b>AB</b>		CIPLA LTD	<u>EQ 15MG BASE</u>	<b>A076798 001</b>	Oct 26, 2012
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A076798 002</b>	Oct 26, 2012
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A076798 003</b>	Oct 26, 2012
<b>AB</b>		DR REDDYS LABS LTD	<u>EQ 15MG BASE</u>	<b>A078383 001</b>	Mar 12, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A078383 002</b>	Mar 12, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A078383 003</b>	Mar 12, 2013
<b>AB</b>		LUPIN LTD	<u>EQ 15MG BASE</u>	<b>A204133 001</b>	Apr 07, 2014
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A204133 002</b>	Apr 07, 2014
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A204133 003</b>	Apr 07, 2014
<b>AB</b>		MACLEODS PHARMS LTD	<u>EQ 15MG BASE</u>	<b>A202467 001</b>	Feb 06, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A202467 002</b>	Feb 06, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A202467 003</b>	Feb 06, 2013
<b>AB</b>		MYLAN PHARMS INC	<u>EQ 15MG BASE</u>	<b>A076801 001</b>	Aug 17, 2012
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A076801 002</b>	Aug 17, 2012
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A076801 003</b>	Aug 17, 2012
<b>AB</b>		PURACAP PHARM LLC	<u>EQ 15MG BASE</u>	<b>A206738 001</b>	Oct 06, 2017
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A206738 002</b>	Oct 06, 2017
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A206738 003</b>	Oct 06, 2017
<b>AB</b>		SANDOZ	<u>EQ 15MG BASE</u>	<b>A078670 001</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A078670 002</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A078670 003</b>	Feb 13, 2013
<b>AB</b>		TEVA PHARMS USA	<u>EQ 15MG BASE</u>	<b>A077210 001</b>	Jan 10, 2014
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A077210 002</b>	Jan 10, 2014
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A077210 003</b>	Jan 10, 2014
<b>AB</b>		TORRENT PHARMS LTD	<u>EQ 15MG BASE</u>	<b>A091298 001</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A091298 002</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A091298 003</b>	Feb 13, 2013
<b>AB</b>		ZYDUS PHARMS USA INC	<u>EQ 15MG BASE</u>	<b>A202456 001</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 30MG BASE</u>	<b>A202456 002</b>	Feb 13, 2013
<b>AB</b>			<u>EQ 45MG BASE</u>	<b>A202456 003</b>	Feb 13, 2013

PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPERACILLIN

!	ISTITUTO BIO ITA SPA	<u>EQ 2GM BASE/VIAL</u>	<b>A065114 001</b>	Nov 14, 2003
!		<u>EQ 3GM BASE/VIAL</u>	<b>A065114 002</b>	Nov 14, 2003
!		<u>EQ 4GM BASE/VIAL</u>	<b>A065114 003</b>	Nov 14, 2003
!		<u>EQ 40GM BASE/VIAL</u>	<b>A065157 001</b>	Jul 12, 2004

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

**PIPERACILLIN AND TAZOBACTAM**

<b>AP</b>	Apollo PHARMS INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<b>A207847 001</b>	Jan 13, 2017
<b>AP</b>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<b>A207847 002</b>	Jan 13, 2017
<b>AP</b>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<b>A207848 002</b>	Jan 13, 2017
<b>AP</b>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<b>A207847 003</b>	Jan 13, 2017
<b>AP</b>	AUROBINDO PHARMA LTD	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<b>A065498 001</b>	May 23, 2011
<b>AP</b>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<b>A065498 002</b>	May 23, 2011
<b>AP</b>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<b>A065498 003</b>	May 23, 2011
<b>AP</b>	HOSPIRA INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<b>A065386 001</b>	Sep 15, 2009
<b>AP</b>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<b>A065386 002</b>	Sep 15, 2009
<b>AP</b>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<b>A065386 003</b>	Sep 15, 2009
<b>AP</b>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<b>A065446 001</b>	Sep 15, 2009
<b>AP</b>	ISTITUTO BIO ITA	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<b>A065523 001</b>	May 31, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-344 (of 436)

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

**PIPERACILLIN AND TAZOBACTAM**

SPA

<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065523 002</u>	May 31, 2011	
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065523 003</u>	May 31, 2011	
<u>AP</u>		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>A090498 001</u>	May 31, 2011	
<u>AP</u>	MYLAN LABS LTD	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065458 001</u>	Aug 15, 2014	
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065458 002</u>	Aug 15, 2014	
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065458 003</u>	Aug 15, 2014	
<u>AP</u>	SANDOZ	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065362 001</u>	Oct 21, 2010	
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A065363 001</u>	Oct 21, 2010	
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065362 002</u>	Oct 21, 2010	
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A065363 002</u>	Oct 21, 2010	
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065362 003</u>	Oct 21, 2010	
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A065363 003</u>	Oct 21, 2010	
<u>AP</u>	WOCKHARDT BIO AG	<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A207146 001</u>	Mar 17, 2017	
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>A206996 001</u>	Mar 22, 2017	
<u>AP</u>		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>A206996 002</u>	Mar 22, 2017	
<u>AP</u>		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>A206996 003</u>	Mar 22, 2017	
<b>ZOSYN</b>					
<u>AP</u>	+!	WYETH PHARMS INC	<u>EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL</u>	<u>N050684 001</u>	Oct 22, 1993
<u>AP</u>	+!		<u>EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL</u>	<u>N050684 002</u>	Oct 22, 1993
<u>AP</u>	+!		<u>EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N050684 003</u>	Oct 22, 1993
<u>AP</u>	+!		<u>EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL</u>	<u>N050684 004</u>	Oct 22, 1993
ZOSYN IN PLASTIC CONTAINER					
	+!	WYETH PHARMS INC	EQ 40MG BASE/ML;EQ 5MG BASE/ML	N050750 001	Feb 24, 1998
	+!		EQ 60MG BASE/ML;EQ 7.5MG BASE/ML	N050750 002	Feb 24, 1998
	+!		EQ 4GM BASE/100ML;EQ 500MG BASE/100ML	N050750 003	Feb 24, 1998
INJECTABLE; IV (INFUSION)					
<b>PIPERACILLIN AND TAZOBACTAM</b>					
SANDOZ INC					
		EQ 12GM BASE/VIAL;EQ 1.5GM BASE/VIAL	A203557 001	Oct 29, 2014	

PIRFENIDONE

CAPSULE; ORAL

ESBRIET

+! GENENTECH INC

267MG

N022535 001 Oct 15, 2014

TABLET; ORAL

ESBRIET

+ GENENTECH INC

267MG

N208780 001 Jan 11, 2017

+!

801MG

N208780 003 Jan 11, 2017

PIROXICAM

CAPSULE; ORAL

**FELDENE**

AB + PFIZER

10MG

N018147 002 Apr 06, 1982

AB +!

20MG

N018147 003 Apr 06, 1982

**PIROXICAM**

AB FLAMINGO PHARMS

10MG

A207938 001 Sep 09, 2016

AB HIKMA PHARMS

20MG

A207938 002 Sep 09, 2016

AB MICROLABS

10MG

A209256 001 Aug 11, 2017

AB +

20MG

A209256 002 Aug 11, 2017

AB MYLAN IRELAND LTD

10MG

A206152 001 Dec 29, 2017

AB +

20MG

A206152 002 Dec 29, 2017

AB PII

10MG

A074116 001 Jun 15, 1993

AB +

20MG

A074118 001 Jun 15, 1993

AB SUN PHARM INDUSTRIES

10MG

A206136 001 Jun 20, 2017

AB TEVA

20MG

A206136 002 Jun 20, 2017

AB UNICHEM LABS LTD

10MG

A073536 002 Jan 23, 2008

AB +

20MG

A073536 001 Mar 12, 1993

AB TEVA

10MG

A074131 001 Dec 11, 1992

AB +

20MG

A074131 002 Dec 11, 1992

AB UNICHEM LABS LTD

10MG

A208340 001 Apr 13, 2017

AB +

20MG

A208340 002 Apr 13, 2017

PITAVASTATIN CALCIUM

TABLET; ORAL

**LIVALO**

AB + KOWA CO

EQ 1MG BASE

N022363 001 Aug 03, 2009

AB +

EQ 2MG BASE

N022363 002 Aug 03, 2009

AB +!

EQ 4MG BASE

N022363 003 Aug 03, 2009

**PITAVASTATIN CALCIUM**

AB AUROBINDO PHARMA

EQ 1MG BASE

A206015 001 Dec 20, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-345 (of 436)

PITAVASTATIN CALCIUM

TABLET;ORAL

**PITAVASTATIN CALCIUM**

LTD

<b>AB</b>	<b>EQ 2MG BASE</b>	<b>A206015 002</b>	Dec 20, 2016
<b>AB</b>	<b>EQ 4MG BASE</b>	<b>A206015 003</b>	Dec 20, 2016
<b>AB</b>	<b>ORIENT PHARMA CO LTD</b>	<b>A205932 001</b>	Feb 03, 2017
<b>AB</b>	<b>EQ 2MG BASE</b>	<b>A205932 002</b>	Feb 03, 2017
<b>AB</b>	<b>EQ 4MG BASE</b>	<b>A205932 003</b>	Feb 03, 2017
<b>AB</b>	<b>SAWAI USA</b>	<b>A205955 001</b>	Feb 03, 2017
<b>AB</b>	<b>EQ 1MG BASE</b>	<b>A205955 002</b>	Feb 03, 2017
<b>AB</b>	<b>EQ 2MG BASE</b>	<b>A205955 003</b>	Feb 03, 2017
<b>AB</b>	<b>EQ 4MG BASE</b>		

PITAVASTATIN MAGNESIUM

TABLET;ORAL

ZYPITAMAG

+ ZYDUS PHARMS USA INC	EQ 1MG BASE	N208379 001	Jul 14, 2017
+!	EQ 2MG BASE	N208379 002	Jul 14, 2017
+!	EQ 4MG BASE	N208379 003	Jul 14, 2017

PLECANATIDE

TABLET;ORAL

TRULANCE

+! SYNERGY PHARMS	3MG	N208745 001	Jan 19, 2017
-------------------	-----	-------------	--------------

PLERIXAFOR

SOLUTION;SUBCUTANEOUS

MOZOBIL

+! GENZYME	24MG/1.2ML (20MG/ML)	N022311 001	Dec 15, 2008
------------	----------------------	-------------	--------------

PODOFILOX

GEL;TOPICAL

CONDYLOX

+! ALLERGAN SALES LLC	0.5%	N020529 001	Mar 13, 1997
-----------------------	------	-------------	--------------

SOLUTION;TOPICAL

CONDYLOX

<b>AT</b> +! ALLERGAN SALES LLC	<b>0.5%</b>	<b>N019795 001</b>	Dec 13, 1990
<b>AT</b> PADDOCK LLC	<b>0.5%</b>	<b>A075600 001</b>	Jan 29, 2002

POLIDOCANOL

SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS

VARITHENA

+! PROVENSIS	77.5MG/7.75ML (10MG/ML)	N205098 002	Dec 21, 2017
--------------	-------------------------	-------------	--------------

SOLUTION;INTRAVENOUS

ASCLERA

+ CHEMISCH FBRK KRSSLR	10MG/2ML (5MG/ML)	N021201 001	Mar 30, 2010
+!	20MG/2ML (10MG/ML)	N021201 002	Mar 30, 2010

VARITHENA

+! PROVENSIS	180MG/18ML (10MG/ML)	N205098 001	Nov 25, 2013
--------------	----------------------	-------------	--------------

POLYETHYLENE GLYCOL 3350

FOR SOLUTION;ORAL

GLYCOLAX

<b>AA</b> KREMERS URBAN PHARMS	<b>17GM/SCOOPFUL</b>	<b>A076652 001</b>	Jul 02, 2004
<b>AA</b> BRECKENRIDGE PHARM	<b>17GM/SCOOPFUL</b>	<b>A077736 001</b>	May 26, 2006

<b>AA</b> NEXGEN PHARMA INC	<b>17GM/SCOOPFUL</b>	<b>A077706 001</b>	Sep 27, 2006
<b>AA</b> PADDOCK LLC	<b>17GM/SCOOPFUL</b>	<b>A077893 001</b>	May 26, 2006

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION;ORAL

LAX-LYTE WITH FLAVOR PACKS

<b>AA</b> PADDOCK LLC	<b>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</b>	<b>A079232 001</b>	Feb 25, 2010
<b>AA</b> +! BRAINTREE	<b>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</b>	<b>N019797 001</b>	Apr 22, 1991

NULYTLEY

<b>AA</b> +! BRAINTREE	<b>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</b>	<b>N019797 002</b>	Nov 18, 1994
<b>AA</b> NOVEL LABS INC	<b>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</b>	<b>A090019 001</b>	May 27, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-346 (of 436)

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE  
FOR SOLUTION;ORAL

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE  
GM/BOT  
AA STRIDES PHARMA 420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT A204559 001 Apr 13, 2015  
PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE  
AA BRECKENRIDGE PHARM 420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT A202060 001 Mar 08, 2017  
TRILYTE  
AA MYLAN PHARMS INC 420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT A076491 001 Feb 05, 2004

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE  
ANHYDROUS

FOR SOLUTION;ORAL  
COLYTE WITH FLAVOR PACKS  
AA MYLAN SPECIALITY LP 240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT N018983 012 Oct 08, 1998  
GOLYTELY  
AA +! BRAINTREE 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT N019011 001 Jul 13, 1984  
PEG 3350 AND ELECTROLYTES  
AA NOVEL LABS INC 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT A090231 001 Jun 01, 2009  
AA 240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT A090186 001 Jun 01, 2009  
GOLYTELY  
+! BRAINTREE 227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET N019011 002 Jun 02, 1992

POLYMYXIN B SULFATE

INJECTABLE;INJECTION

POLYMYCIN B SULFATE  
AP SAGENT STRIDES EQ 500,000 UNITS BASE/VIAL A090110 001 Jun 29, 2011  
POLYMYXIN B SULFATE  
AP AUROBINDO PHARMA LTD EQ 500,000 UNITS BASE/VIAL A206589 001 Apr 04, 2016  
AP FRESENIUS KABI USA EQ 500,000 UNITS BASE/VIAL A065372 001 Jan 10, 2008  
AP GLAND PHARMA LTD EQ 500,000 UNITS BASE/VIAL A207322 001 Apr 14, 2016  
AP ! WEST-WARD PHARMS INT EQ 500,000 UNITS BASE/VIAL A060716 001  
AP X GEN PHARMS EQ 500,000 UNITS BASE/VIAL A063000 001 Sep 30, 1994  
AP XELLIA PHARMS APS EQ 500,000 UNITS BASE/VIAL A202766 001 Jan 15, 2014

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS;OPHTHALMIC

POLYTRIM  
AT +! ALLERGAN 10,000 UNITS/ML;EQ 1MG BASE/ML N050567 001 Oct 20, 1988  
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE  
AT AKORN INC 10,000 UNITS/ML;EQ 1MG BASE/ML A065006 001 Dec 17, 1998  
AT BAUSCH AND LOMB 10,000 UNITS/ML;EQ 1MG BASE/ML A064120 001 Feb 14, 1997  
AT SANDOZ INC 10,000 UNITS/ML;EQ 1MG BASE/ML A064211 001 Apr 13, 1998

POMALIDOMIDE

CAPSULE;ORAL

POMALYST  
+ CELGENE 1MG N204026 001 Feb 08, 2013  
+ 2MG N204026 002 Feb 08, 2013  
+ 3MG N204026 003 Feb 08, 2013  
+! 4MG N204026 004 Feb 08, 2013

PONATINIB HYDROCHLORIDE

TABLET;ORAL

ICLUSIG  
+ ARIAD EQ 15MG BASE N203469 001 Dec 14, 2012  
+ EQ 30MG BASE N203469 003 Apr 23, 2015  
+! EQ 45MG BASE N203469 002 Dec 14, 2012

PORACTANT ALFA

SUSPENSION;INTRATRACHEAL

CUROSURF  
+! CHIESI USA INC 80MG/ML N020744 001 Nov 18, 1999

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-347 (of 436)

PORFIMER SODIUM

INJECTABLE; INJECTION  
 PHOTOFRIN  
 CONCORDIA LABS INC 75MG/VIAL N020451 001 Dec 27, 1995

POSACONAZOLE

SOLUTION; IV (INFUSION)  
 NOXAFL  
 +! MERCK SHARP DOHME 300MG/16.7ML (18MG/ML) N205596 001 Mar 13, 2014  
 SUSPENSION; ORAL  
 NOXAFL  
 +! SCHERING 40MG/ML N022003 001 Sep 15, 2006  
 TABLET, DELAYED RELEASE; ORAL  
 NOXAFL  
 +! MERCK SHARP DOHME 100MG N205053 001 Nov 25, 2013

POTASSIUM ACETATE

INJECTABLE; INJECTION  
POTASSIUM ACETATE  
AP EXELA PHARMA SCS 2MEO/ML A206203 001 Dec 29, 2015  
 LLC  
AP +! HOSPIRA 2MEO/ML N018896 001 Jul 20, 1984

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL  
KLOR-CON  
AB UPSHER-SMITH LABS 8MEO A203106 001 Jul 10, 2015  
AB 10MEO A203106 002 Jul 10, 2015  
MICRO-K  
AB + NESHER PHARMS 8MEO N018238 001  
MICRO-K 10  
AB + NESHER PHARMS 10MEO N018238 002 May 14, 1984  
POTASSIUM CHLORIDE  
AB ACTAVIS LABS FL INC 8MEO A077419 001 Jun 02, 2008  
AB ! 10MEO A077419 002 Jun 02, 2008  
AB ADARE PHARMS INC 8MEO A208864 001 Mar 17, 2017  
AB 10MEO A208864 002 Mar 17, 2017  
AB AMNEAL PHARMS 10MEO A202128 001 Feb 22, 2013  
AB ANCHEN PHARMS 8MEO A202886 001 Dec 26, 2013  
AB 10MEO A202886 002 Dec 26, 2013  
AB GLENMARK PHARMS LTD 10MEO A202868 001 Jan 19, 2016  
AB KREMERS URBAN 8MEO A204210 001 Mar 28, 2016  
 PHARMS  
AB 10MEO A204210 002 Mar 28, 2016  
AB LUPIN LTD 8MEO A203002 001 Dec 18, 2015  
AB 10MEO A203002 002 Dec 18, 2015  
AB NOVEL LABS INC 8MEO A204828 001 Aug 16, 2016  
AB 10MEO A204828 002 Aug 16, 2016  
AB PADDOCK LLC 8MEO A200185 001 May 18, 2011  
AB 10MEO A200185 002 May 18, 2011  
AB PII 8MEO A205549 001 Dec 08, 2015  
AB 10MEO A205549 002 Dec 08, 2015  
AB TRIS PHARMA INC 8MEO A201944 001 Mar 04, 2016  
AB 10MEO A201944 002 Mar 04, 2016

FOR SOLUTION; ORAL

KLOR-CON  
AA UPSHER-SMITH LABS 20MEO A209662 001 Oct 23, 2017  
POTASSIUM CHLORIDE  
AA +! PHARMA RES SOFTWARE 20MEO N208019 001 Aug 19, 2015

INJECTABLE; INJECTION

POTASSIUM CHLORIDE  
AP B BRAUN 2MEO/ML A085870 001  
AP FRESENIUS KABI USA 2MEO/ML A080225 001  
AP ! HOSPIRA 2MEO/ML A080205 001  
POTASSIUM CHLORIDE 10MEO IN PLASTIC CONTAINER  
AP +! BAXTER HLTHCARE 14.9MG/ML N019904 001 Dec 26, 1989  
AP +! 746MG/100ML N019904 005 Dec 17, 1990  
AP + ICU MEDICAL INC 14.9MG/ML N020161 005 Nov 30, 1992  
AP + 745MG/100ML N020161 001 Nov 30, 1992  
POTASSIUM CHLORIDE 20MEO IN PLASTIC CONTAINER  
AP +! BAXTER HLTHCARE 29.8MG/ML N019904 002 Dec 26, 1989  
AP +! 1.49GM/100ML N019904 006 Dec 17, 1990  
AP +! ICU MEDICAL INC 29.8MG/ML N020161 006 Aug 11, 1998  
AP + 1.49GM/100ML N020161 002 Nov 30, 1992

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-348 (of 436)

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

<u>AP</u>	+!	BAXTER HLTHCARE	<u>2.24GM/100ML</u>	<u>N019904 003</u>	Dec 26, 1989
<u>AP</u>	+!	ICU MEDICAL INC	<u>2.24GM/100ML</u>	<u>N020161 003</u>	Aug 11, 1998

POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

<u>AP</u>	+!	BAXTER HLTHCARE	<u>2.98GM/100ML</u>	<u>N019904 004</u>	Dec 26, 1989
<u>AP</u>	+!	ICU MEDICAL INC	<u>2.98GM/100ML</u>	<u>N020161 004</u>	Aug 11, 1998

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	FRESENIUS KABI USA	<u>2MEO/ML</u>	<u>A088901 001</u>	Jan 25, 1985
<u>AP</u>		<u>2MEO/ML</u>	<u>A088908 001</u>	Jan 25, 1985

POTASSIUM CHLORIDE

! FRESENIUS KABI USA 3MEO/ML

A080225 003

SOLUTION; ORAL

POTASSIUM CHLORIDE

+ GENUS LIFESCIENCES 20MEO/15ML  
 +! 40MEO/15ML

N206814 001 Dec 22, 2014  
 N206814 002 Dec 22, 2014

TABLET, EXTENDED RELEASE; ORAL

KLOR-CON M10

<u>AB1</u>	UPSHER-SMITH LABS	<u>10MEO</u>	<u>A074726 002</u>	Aug 09, 2000
------------	-------------------	--------------	--------------------	--------------

KLOR-CON M20

<u>AB1</u>	! UPSHER-SMITH LABS	<u>20MEO</u>	<u>A074726 001</u>	Nov 20, 1998
------------	---------------------	--------------	--------------------	--------------

POTASSIUM CHLORIDE

<u>AB1</u>	ACTAVIS LABS FL INC	<u>10MEO</u>	<u>A075604 001</u>	Apr 10, 2002
<u>AB1</u>		<u>20MEO</u>	<u>A075604 002</u>	Apr 10, 2002
<u>AB1</u>	ADARE PHARMS INC	<u>20MEO</u>	<u>A076368 001</u>	Aug 18, 2004
<u>AB1</u>	GLENMARK PHARMS LTD	<u>10MEO</u>	<u>A203562 001</u>	Jul 26, 2016
<u>AB1</u>		<u>20MEO</u>	<u>A203562 002</u>	Jul 26, 2016
<u>AB1</u>	NOVEL LABS INC	<u>10MEO</u>	<u>A206347 001</u>	Jan 21, 2016
<u>AB1</u>		<u>20MEO</u>	<u>A206347 002</u>	Jan 21, 2016

KLOR-CON

<u>AB2</u>	+ UPSHER-SMITH LABS	<u>8MEO</u>	<u>N019123 001</u>	Apr 17, 1986
<u>AB2</u>	+!	<u>10MEO</u>	<u>N019123 002</u>	Apr 17, 1986

POTASSIUM CHLORIDE

<u>AB2</u>	MYLAN PHARMS INC	<u>8MEO</u>	<u>A204662 001</u>	Aug 21, 2014
<u>AB2</u>		<u>10MEO</u>	<u>A204662 002</u>	Aug 21, 2014
<u>AB2</u>	NOVEL LABS INC	<u>8MEO</u>	<u>A206759 001</u>	Aug 09, 2016
<u>AB2</u>		<u>10MEO</u>	<u>A206759 002</u>	Aug 09, 2016
<u>AB2</u>	PADDOCK LLC	<u>8MEO</u>	<u>A205993 001</u>	Nov 05, 2015
<u>AB2</u>		<u>10MEO</u>	<u>A205993 002</u>	Nov 05, 2015
<u>AB2</u>	SIGMAPHARM LABS LLC	<u>8MEO</u>	<u>A207528 001</u>	Aug 19, 2016
<u>AB2</u>		<u>10MEO</u>	<u>A207528 002</u>	Aug 19, 2016

K-TAB

BC	+	ABBVIE	8MEO	N018279 002 Aug 01, 1988
BC	+		10MEO	N018279 001
BC	!+		20MEO	N018279 003 Nov 25, 2013

KLOR-CON M15

UPSHER-SMITH LABS 15MEO A074726 003 Jun 06, 2003

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AP</u>	ICU MEDICAL INC	<u>149MG/100ML;450MG/100ML</u>	<u>A078446 001</u>	Sep 10, 2008
-----------	-----------------	--------------------------------	--------------------	--------------

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AP</u>	+!	BAXTER HLTHCARE	<u>150MG/100ML;450MG/100ML</u>	<u>N017648 005</u>	Nov 26, 2002
-----------	----	-----------------	--------------------------------	--------------------	--------------

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>150MG/100ML;900MG/100ML</u>	<u>N019708 004</u>	Sep 29, 1989
-----------	---------	--------------------------------	--------------------	--------------

<u>AP</u>	+	BAXTER HLTHCARE	<u>150MG/100ML;900MG/100ML</u>	<u>N017648 001</u>
-----------	---	-----------------	--------------------------------	--------------------

POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>300MG/100ML;900MG/100ML</u>	<u>N017648 002</u>
-----------	---	-----------------	--------------------------------	--------------------

POTASSIUM CHLORIDE 20MEO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	ICU MEDICAL INC	<u>149MG/100ML;900MG/100ML</u>	<u>N019686 001</u>	Oct 17, 1988
-----------	-----------------	--------------------------------	--------------------	--------------

POTASSIUM CHLORIDE 40MEO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	ICU MEDICAL INC	<u>298MG/100ML;900MG/100ML</u>	<u>N019686 002</u>	Oct 17, 1988
-----------	-----------------	--------------------------------	--------------------	--------------

POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9%

+ BAXTER HLTHCARE 224MG/100ML;900MG/100ML

N017648 003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-349 (of 436)

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL

**POTASSIUM CITRATE**

<b>AB</b>	COREPHARMA	<u>5MEO</u>	<b>A077440 001</b>	Jun 09, 2006
<b>AB</b>		<u>10MEO</u>	<b>A077440 002</b>	Jun 09, 2006
<b>AB</b>	STRIDES PHARMA	<u>5MEO</u>	<b>A206813 001</b>	Sep 11, 2017
<b>AB</b>		<u>10MEO</u>	<b>A206813 002</b>	Sep 11, 2017
<b>AB</b>		<u>15MEO</u>	<b>A206813 003</b>	Sep 11, 2017
<b>AB</b>	ZYDUS PHARMS USA INC	<u>5MEO</u>	<b>A203546 001</b>	Aug 06, 2014
<b>AB</b>		<u>10MEO</u>	<b>A203546 002</b>	Aug 06, 2014
<b>AB</b>		<u>15MEO</u>	<b>A203546 003</b>	Aug 06, 2014
<b>UROCIT-K</b>				
<b>AB</b>	+ MISSION PHARMA	<u>5MEO</u>	<b>N019071 001</b>	Aug 30, 1985
<b>AB</b>	+!	<u>10MEO</u>	<b>N019071 002</b>	Aug 31, 1992
<b>AB</b>	+!	<u>15MEO</u>	<b>N019071 003</b>	Dec 30, 2009

POVIDONE-IODINE

SOLUTION/DROPS; OPHTHALMIC

BETADINE

+! ALCON PHARMS LTD

5%

N018634 001 Dec 17, 1986

PRALATREXATE

SOLUTION; INTRAVENOUS

FOLOTYN

+! ALLOS

20MG/ML (20MG/ML)

N022468 001 Sep 24, 2009

+!

40MG/2ML (20MG/ML)

N022468 002 Sep 24, 2009

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE

+! MERIDIAN MEDCL TECHN

300MG/ML

N018986 001 Apr 26, 1983

PROTOPAM CHLORIDE

+! BAXTER HLTHCARE CORP

1GM/VIAL

N014134 001

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

**MIRAPEX**

<b>AB</b>	+ BOEHRINGER INGELHEIM	<u>0.125MG</u>	<b>N020667 001</b>	Jul 01, 1997
<b>AB</b>	+!	<u>0.25MG</u>	<b>N020667 002</b>	Jul 01, 1997
<b>AB</b>	+	<u>0.5MG</u>	<b>N020667 006</b>	Feb 12, 1998
<b>AB</b>	+	<u>0.75MG</u>	<b>N020667 007</b>	Jul 30, 2007
<b>AB</b>	+	<u>1MG</u>	<b>N020667 003</b>	Jul 01, 1997
<b>AB</b>	+	<u>1.5MG</u>	<b>N020667 005</b>	Jul 01, 1997
<b>PRAMIPEXOLE DIHYDROCHLORIDE</b>				
<b>AB</b>	ALEMBIC PHARMS LTD	<u>0.125MG</u>	<b>A078894 001</b>	Oct 08, 2010
<b>AB</b>		<u>0.25MG</u>	<b>A078894 002</b>	Oct 08, 2010
<b>AB</b>		<u>0.5MG</u>	<b>A078894 003</b>	Oct 08, 2010
<b>AB</b>		<u>1MG</u>	<b>A078894 004</b>	Oct 08, 2010
<b>AB</b>		<u>1.5MG</u>	<b>A078894 005</b>	Oct 08, 2010
<b>AB</b>	APOTEX INC	<u>0.125MG</u>	<b>A090151 001</b>	Apr 30, 2012
<b>AB</b>		<u>0.25MG</u>	<b>A090151 002</b>	Apr 30, 2012
<b>AB</b>		<u>0.5MG</u>	<b>A090151 003</b>	Apr 30, 2012
<b>AB</b>		<u>0.75MG</u>	<b>A090151 006</b>	Apr 30, 2012
<b>AB</b>		<u>1MG</u>	<b>A090151 004</b>	Apr 30, 2012
<b>AB</b>		<u>1.5MG</u>	<b>A090151 005</b>	Apr 30, 2012
<b>AB</b>	AUROBINDO PHARMA LTD	<u>0.125MG</u>	<b>A202633 001</b>	Oct 26, 2012
<b>AB</b>		<u>0.25MG</u>	<b>A202633 002</b>	Oct 26, 2012
<b>AB</b>		<u>0.5MG</u>	<b>A202633 003</b>	Oct 26, 2012
<b>AB</b>		<u>0.75MG</u>	<b>A202633 004</b>	Oct 26, 2012
<b>AB</b>		<u>1MG</u>	<b>A202633 005</b>	Oct 26, 2012
<b>AB</b>		<u>1.5MG</u>	<b>A202633 006</b>	Oct 26, 2012
<b>AB</b>	BARR	<u>0.125MG</u>	<b>A077724 001</b>	Feb 19, 2008
<b>AB</b>		<u>0.25MG</u>	<b>A077724 002</b>	Feb 19, 2008
<b>AB</b>		<u>0.5MG</u>	<b>A077724 003</b>	Feb 19, 2008
<b>AB</b>		<u>1MG</u>	<b>A077724 004</b>	Feb 19, 2008
<b>AB</b>		<u>1.5MG</u>	<b>A077724 005</b>	Feb 19, 2008
<b>AB</b>	BRECKENRIDGE PHARM	<u>0.125MG</u>	<b>A091450 001</b>	Oct 08, 2010
<b>AB</b>		<u>0.25MG</u>	<b>A091450 002</b>	Oct 08, 2010
<b>AB</b>		<u>0.5MG</u>	<b>A091450 003</b>	Oct 08, 2010
<b>AB</b>		<u>1MG</u>	<b>A091450 004</b>	Oct 08, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-350 (of 436)

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>		<u>1.5MG</u>	<u>A091450</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>	GLENMARK GENERICS	<u>0.125MG</u>	<u>A090781</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A090781</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090781</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090781</u>	<u>006</u>	Sep 11, 2015
<u>AB</u>		<u>1MG</u>	<u>A090781</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090781</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>	MACLEODS PHARMS LTD	<u>0.125MG</u>	<u>A202164</u>	<u>001</u>	Sep 20, 2012
<u>AB</u>		<u>0.25MG</u>	<u>A202164</u>	<u>002</u>	Sep 20, 2012
<u>AB</u>		<u>0.5MG</u>	<u>A202164</u>	<u>003</u>	Sep 20, 2012
<u>AB</u>		<u>1MG</u>	<u>A202164</u>	<u>004</u>	Sep 20, 2012
<u>AB</u>		<u>1.5MG</u>	<u>A202164</u>	<u>005</u>	Sep 20, 2012
<u>AB</u>	MYLAN	<u>0.125MG</u>	<u>A077854</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A077854</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A077854</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090764</u>	<u>001</u>	Apr 09, 2010
<u>AB</u>		<u>1MG</u>	<u>A077854</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A077854</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>	SCIEGEN PHARMS INC	<u>0.125MG</u>	<u>A203855</u>	<u>001</u>	Oct 28, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A203855</u>	<u>002</u>	Oct 28, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A203855</u>	<u>003</u>	Oct 28, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A203855</u>	<u>004</u>	Oct 28, 2014
<u>AB</u>		<u>1MG</u>	<u>A203855</u>	<u>005</u>	Oct 28, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A203855</u>	<u>006</u>	Oct 28, 2014
<u>AB</u>	STRIDES PHARMA	<u>0.125MG</u>	<u>A202702</u>	<u>001</u>	Jun 03, 2014
<u>AB</u>		<u>0.25MG</u>	<u>A202702</u>	<u>002</u>	Jun 03, 2014
<u>AB</u>		<u>0.5MG</u>	<u>A202702</u>	<u>003</u>	Jun 03, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202702</u>	<u>004</u>	Jun 03, 2014
<u>AB</u>		<u>1MG</u>	<u>A202702</u>	<u>005</u>	Jun 03, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202702</u>	<u>006</u>	Jun 03, 2014
<u>AB</u>	SUN PHARM INDs INC	<u>0.125MG</u>	<u>A091683</u>	<u>001</u>	Mar 27, 2013
<u>AB</u>		<u>0.25MG</u>	<u>A091683</u>	<u>002</u>	Mar 27, 2013
<u>AB</u>		<u>0.5MG</u>	<u>A091683</u>	<u>003</u>	Mar 27, 2013
<u>AB</u>		<u>0.75MG</u>	<u>A091683</u>	<u>004</u>	Mar 27, 2013
<u>AB</u>		<u>1MG</u>	<u>A091683</u>	<u>005</u>	Mar 27, 2013
<u>AB</u>		<u>1.5MG</u>	<u>A091683</u>	<u>006</u>	Mar 27, 2013
<u>AB</u>	TEVA PHARMS	<u>0.125MG</u>	<u>A090241</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A090241</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090241</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090241</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1MG</u>	<u>A090241</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090241</u>	<u>006</u>	Oct 08, 2010
<u>AB</u>	TORRENT PHARMS	<u>0.125MG</u>	<u>A090865</u>	<u>001</u>	Oct 08, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A090865</u>	<u>002</u>	Oct 08, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A090865</u>	<u>003</u>	Oct 08, 2010
<u>AB</u>		<u>0.75MG</u>	<u>A090865</u>	<u>004</u>	Oct 08, 2010
<u>AB</u>		<u>1MG</u>	<u>A090865</u>	<u>005</u>	Oct 08, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A090865</u>	<u>006</u>	Oct 08, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.125MG</u>	<u>A078920</u>	<u>001</u>	Jul 06, 2010
<u>AB</u>		<u>0.25MG</u>	<u>A078920</u>	<u>002</u>	Jul 06, 2010
<u>AB</u>		<u>0.5MG</u>	<u>A078920</u>	<u>003</u>	Jul 06, 2010
<u>AB</u>		<u>1MG</u>	<u>A078920</u>	<u>004</u>	Jul 06, 2010
<u>AB</u>		<u>1.5MG</u>	<u>A078920</u>	<u>005</u>	Jul 06, 2010

TABLET, EXTENDED RELEASE; ORAL

MIRAPEX ER

<u>AB</u>	+!	BOEHRINGER INGELHEIM	<u>0.375MG</u>	<u>N022421</u>	<u>001</u>	Feb 19, 2010
<u>AB</u>	+		<u>0.75MG</u>	<u>N022421</u>	<u>002</u>	Feb 19, 2010
<u>AB</u>	+		<u>1.5MG</u>	<u>N022421</u>	<u>003</u>	Feb 19, 2010
<u>AB</u>	+		<u>2.25MG</u>	<u>N022421</u>	<u>006</u>	Jun 17, 2011
<u>AB</u>	+		<u>3MG</u>	<u>N022421</u>	<u>004</u>	Feb 19, 2010
<u>AB</u>	+		<u>3.75MG</u>	<u>N022421</u>	<u>007</u>	Jun 17, 2011
<u>AB</u>	+		<u>4.5MG</u>	<u>N022421</u>	<u>005</u>	Feb 19, 2010

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>	ACTAVIS ELIZABETH	<u>0.375MG</u>	<u>A201963</u>	<u>001</u>	Apr 21, 2016
<u>AB</u>		<u>0.75MG</u>	<u>A201963</u>	<u>002</u>	Apr 21, 2016
<u>AB</u>		<u>1.5MG</u>	<u>A201963</u>	<u>003</u>	Apr 21, 2016
<u>AB</u>		<u>2.25MG</u>	<u>A203615</u>	<u>001</u>	Oct 14, 2016
<u>AB</u>		<u>3MG</u>	<u>A201963</u>	<u>004</u>	Apr 21, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-351 (of 436)

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

PRAMIPEXOLE DIHYDROCHLORIDE

<u>AB</u>		<u>3.75MG</u>	<u>A203615 002</u>	Jan 03, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A201963 005</u>	Apr 21, 2016
<u>AB</u>	ANCHEN PHARMS	<u>0.375MG</u>	<u>A202206 001</u>	Feb 06, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202206 002</u>	Feb 06, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202206 003</u>	Feb 06, 2014
<u>AB</u>		<u>2.25MG</u>	<u>A202206 004</u>	Feb 06, 2014
<u>AB</u>		<u>3MG</u>	<u>A202206 005</u>	Feb 06, 2014
<u>AB</u>		<u>3.75MG</u>	<u>A202206 006</u>	Feb 06, 2014
<u>AB</u>		<u>4.5MG</u>	<u>A202206 007</u>	Feb 06, 2014
<u>AB</u>	DR REDDYS LABS LTD	<u>0.375MG</u>	<u>A203354 001</u>	Aug 07, 2015
<u>AB</u>		<u>0.75MG</u>	<u>A203354 002</u>	Aug 07, 2015
<u>AB</u>		<u>1.5MG</u>	<u>A203354 003</u>	Aug 07, 2015
<u>AB</u>		<u>3MG</u>	<u>A203354 004</u>	Aug 07, 2015
<u>AB</u>		<u>4.5MG</u>	<u>A203354 005</u>	Aug 07, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>0.375MG</u>	<u>A206156 001</u>	Jun 24, 2016
<u>AB</u>		<u>0.75MG</u>	<u>A206156 002</u>	Jun 24, 2016
<u>AB</u>		<u>1.5MG</u>	<u>A206156 003</u>	Jun 24, 2016
<u>AB</u>		<u>2.25MG</u>	<u>A206156 004</u>	Jun 24, 2016
<u>AB</u>		<u>3MG</u>	<u>A206156 005</u>	Jun 24, 2016
<u>AB</u>		<u>3.75MG</u>	<u>A206156 007</u>	Jan 23, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A206156 006</u>	Jun 24, 2016
<u>AB</u>	SANDOZ INC	<u>0.375MG</u>	<u>A202353 001</u>	Dec 04, 2014
<u>AB</u>		<u>0.75MG</u>	<u>A202353 002</u>	Dec 04, 2014
<u>AB</u>		<u>1.5MG</u>	<u>A202353 003</u>	Dec 04, 2014
<u>AB</u>		<u>3MG</u>	<u>A202353 004</u>	Dec 04, 2014
<u>AB</u>		<u>4.5MG</u>	<u>A202353 005</u>	Dec 04, 2014
<u>AB</u>	ZYDUS PHARMS USA INC	<u>0.375MG</u>	<u>A202891 001</u>	Dec 12, 2017
<u>AB</u>		<u>0.75MG</u>	<u>A202891 002</u>	Dec 12, 2017
<u>AB</u>		<u>1.5MG</u>	<u>A202891 003</u>	Dec 12, 2017
<u>AB</u>		<u>2.25MG</u>	<u>A202891 004</u>	Dec 12, 2017
<u>AB</u>		<u>3MG</u>	<u>A202891 005</u>	Dec 12, 2017
<u>AB</u>		<u>3.75MG</u>	<u>A202891 006</u>	Dec 12, 2017
<u>AB</u>		<u>4.5MG</u>	<u>A202891 007</u>	Dec 12, 2017

PRAMILINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN

+ ASTRAZENECA AB

EQ 1.5MG BASE/1.5ML (EQ 1MG BASE/ML)

N021332 002 Sep 25, 2007

+

EQ 2.7MG BASE/2.7ML (EQ 1MG BASE/ML)

N021332 003 Sep 25, 2007

PRASTERONE

INSERT; VAGINAL

INTRAROSA

+! AMAG PHARMS INC

6.5MG

N208470 001 Nov 16, 2016

PRASUGREL HYDROCHLORIDE

TABLET; ORAL

EFFIENT

<u>AB</u>	+ ELI LILLY AND CO	<u>EQ 5MG BASE</u>	<u>N022307 001</u>	Jul 10, 2009
<u>AB</u>	+!	<u>EQ 10MG BASE</u>	<u>N022307 002</u>	Jul 10, 2009
<u>AB</u>	<u>PRASUGREL</u>			
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A205888 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205888 002</u>	Oct 16, 2017
<u>AB</u>	LIBERTY PHARMA INC	<u>EQ 5MG BASE</u>	<u>A205790 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205790 002</u>	Oct 16, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A205927 001</u>	Jul 12, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205927 002</u>	Jul 12, 2017
<u>AB</u>	PANACEA BIOTEC LTD	<u>EQ 5MG BASE</u>	<u>A205897 001</u>	Oct 16, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205897 002</u>	Oct 16, 2017

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>20MG</u>	<u>N019898 003</u>	Oct 31, 1991
<u>AB</u>	+	<u>40MG</u>	<u>N019898 004</u>	Mar 22, 1993
<u>AB</u>	+	<u>80MG</u>	<u>N019898 008</u>	Dec 18, 2001
<u>AB</u>	<u>PRAVASTATIN SODIUM</u>			
<u>AB</u>	ACCORD HLTHCARE	<u>10MG</u>	<u>A207068 001</u>	Nov 17, 2016
<u>AB</u>		<u>20MG</u>	<u>A207068 002</u>	Nov 17, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-352 (of 436)

PRAVASTATIN SODIUM

TABLET;ORAL

PRAVASTATIN SODIUM

<u>AB</u>		<u>40MG</u>	<u>A207068</u> <u>003</u>	Nov 17, 2016
<u>AB</u>		<u>80MG</u>	<u>A207068</u> <u>004</u>	Nov 17, 2016
<u>AB</u>	APOTEX INC	<u>10MG</u>	<u>A076341</u> <u>001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076341</u> <u>002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076341</u> <u>003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076341</u> <u>004</u>	Dec 28, 2007
<u>AB</u>	AUROBINDO PHARMA LTD	<u>10MG</u>	<u>A203367</u> <u>001</u>	Feb 02, 2017
<u>AB</u>		<u>20MG</u>	<u>A203367</u> <u>002</u>	Feb 02, 2017
<u>AB</u>		<u>40MG</u>	<u>A203367</u> <u>003</u>	Feb 02, 2017
<u>AB</u>		<u>80MG</u>	<u>A203367</u> <u>004</u>	Feb 02, 2017
<u>AB</u>	CIPLA LTD	<u>10MG</u>	<u>A077904</u> <u>001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A077904</u> <u>002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A077904</u> <u>003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A077904</u> <u>004</u>	Mar 22, 2016
<u>AB</u>	DR REDDYS LABS INC	<u>10MG</u>	<u>A076714</u> <u>001</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076714</u> <u>002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076714</u> <u>003</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076714</u> <u>004</u>	Dec 28, 2007
<u>AB</u>	GLENMARK GENERICS	<u>10MG</u>	<u>A077987</u> <u>001</u>	May 11, 2007
<u>AB</u>		<u>20MG</u>	<u>A077987</u> <u>002</u>	May 11, 2007
<u>AB</u>		<u>40MG</u>	<u>A077987</u> <u>003</u>	May 11, 2007
<u>AB</u>		<u>80MG</u>	<u>A077987</u> <u>004</u>	Dec 28, 2007
<u>AB</u>	LUPIN PHARMS	<u>10MG</u>	<u>A077917</u> <u>001</u>	Jan 08, 2008
<u>AB</u>		<u>20MG</u>	<u>A077917</u> <u>002</u>	Jan 08, 2008
<u>AB</u>		<u>40MG</u>	<u>A077917</u> <u>003</u>	Jan 08, 2008
<u>AB</u>		<u>80MG</u>	<u>A077917</u> <u>004</u>	Jan 08, 2008
<u>AB</u>	MYLAN PHARMS INC	<u>10MG</u>	<u>A079187</u> <u>001</u>	May 27, 2010
<u>AB</u>		<u>20MG</u>	<u>A079187</u> <u>002</u>	May 27, 2010
<u>AB</u>		<u>40MG</u>	<u>A079187</u> <u>003</u>	May 27, 2010
<u>AB</u>		<u>80MG</u>	<u>A079187</u> <u>004</u>	May 27, 2010
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>A076397</u> <u>003</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076397</u> <u>002</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076397</u> <u>001</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A077491</u> <u>001</u>	Feb 11, 2008
<u>AB</u>	TEVA	<u>10MG</u>	<u>A076056</u> <u>001</u>	Apr 24, 2006
<u>AB</u>		<u>20MG</u>	<u>A076056</u> <u>002</u>	Apr 24, 2006
<u>AB</u>		<u>40MG</u>	<u>A076056</u> <u>003</u>	Apr 24, 2006
<u>AB</u>	TEVA PHARMS	<u>80MG</u>	<u>A077793</u> <u>001</u>	Jan 15, 2008
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>A076939</u> <u>004</u>	Oct 23, 2006
<u>AB</u>		<u>20MG</u>	<u>A076939</u> <u>003</u>	Oct 23, 2006
<u>AB</u>		<u>40MG</u>	<u>A076939</u> <u>002</u>	Oct 23, 2006
<u>AB</u>		<u>80MG</u>	<u>A076939</u> <u>001</u>	Dec 28, 2007
<u>AB</u>	ZYDUS PHARMS USA	<u>10MG</u>	<u>A077751</u> <u>001</u>	Apr 30, 2008
<u>AB</u>		<u>20MG</u>	<u>A077751</u> <u>002</u>	Apr 30, 2008
<u>AB</u>		<u>40MG</u>	<u>A077751</u> <u>003</u>	Apr 30, 2008
<u>AB</u>		<u>80MG</u>	<u>A077751</u> <u>004</u>	Apr 30, 2008

PRAZIQUANTEL

TABLET;ORAL

BILTRICIDE

<u>AB</u>	+!	BAYER HLTHCARE	<u>600MG</u>	<u>N018714</u> <u>001</u>	Dec 29, 1982
<u>AB</u>		<u>PRAZIQUANTEL</u>		<u>A208820</u> <u>001</u>	Nov 27, 2017

PRAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

MINIPRESS

<u>AB</u>	+	PFIZER	<u>EQ 1MG BASE</u>	<u>N017442</u> <u>002</u>
<u>AB</u>	+		<u>EQ 2MG BASE</u>	<u>N017442</u> <u>003</u>
<u>AB</u>	+		<u>EQ 5MG BASE</u>	<u>N017442</u> <u>001</u>

PRAZOSIN HYDROCHLORIDE

<u>AB</u>		MYLAN	<u>EQ 1MG BASE</u>	<u>A072575</u> <u>003</u>	May 16, 1989
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A072575</u> <u>002</u>	May 16, 1989
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A072575</u> <u>001</u>	May 16, 1989
<u>AB</u>		TEVA PHARMS	<u>EQ 1MG BASE</u>	<u>A071745</u> <u>002</u>	Sep 12, 1988
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>A071745</u> <u>003</u>	Sep 12, 1988
<u>AB</u>			<u>EQ 5MG BASE</u>	<u>A071745</u> <u>001</u>	Sep 12, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-353 (of 436)

PREDNICARBATE

CREAM;TOPICAL

DERMATOP E EMOLLIENT

AB +! VALEANT BERMUDA 0.1% N020279 001 Oct 29, 1993

PREDNICARBATE

AB FOUGERA PHARMS 0.1% A077287 001 Sep 19, 2006  
 OINTMENT;TOPICAL

DERMATOP

AB +! VALEANT PHARMS NORTH 0.1% N019568 001 Sep 23, 1991

PREDNICARBATE

AB FOUGERA PHARMS 0.1% A077236 001 Mar 09, 2007

PREDNISOLONE

SYRUP;ORAL

PREDNISOLONE

AA ! HI TECH PHARMA CO 15MG/5ML A040401 001 Feb 27, 2003  
AA PHARM ASSOC 15MG/5ML A040399 001 Mar 05, 2003  
AA VINTAGE 15MG/5ML A040775 001 Sep 21, 2007  
AA VISTAPHARM 15MG/5ML A040323 001 May 13, 1999  
AA WOCKHARDT BIO AG 15MG/5ML A040313 001 Sep 10, 2003

PRELONE

AA TEVA 15MG/5ML A089081 001 Feb 04, 1986  
 TABLET;ORAL  
 PREDNISOLONE  
 ! WATSON LABS 5MG A080354 001

PREDNISOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

OMNIPRED

AB NOVARTIS PHARMS CORP 1% N017469 001

PRED FORTE

AB +! ALLERGAN 1% N017011 001  
 PRED MILD  
 +! ALLERGAN 0.12% N017100 001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

BLEPHAMIDE S.O.P.  
 ! ALLERGAN 0.2%;10% A087748 001 Dec 03, 1986

SUSPENSION;OPHTHALMIC

BLEPHAMIDE  
 +! ALLERGAN 0.2%;10% N012813 002

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION;ORAL

PEDIAPRED

AA +! SETON PHARM EQ 5MG BASE/5ML N019157 001 May 28, 1986

PREDNISOLONE SODIUM PHOSPHATE

AA CHARTWELL RX EQ 5MG BASE/5ML A075988 001 May 25, 2004  
AA EDENBRIDGE PHARMS EQ 10MG BASE/5ML A203559 001 Dec 20, 2016  
AA EQ 20MG BASE/5ML A203559 002 Dec 20, 2016  
AA ! HI TECH PHARMA EQ 5MG BASE/5ML A075183 001 Mar 26, 2003  
AA ! PHARM ASSOC EQ 10MG BASE/5ML A078465 001 Mar 07, 2008  
AA EQ 15MG BASE/5ML A076913 001 Apr 25, 2005  
AA ! EQ 20MG BASE/5ML A078988 001 Jun 09, 2008  
AA ! VINTAGE EQ 15MG BASE/5ML A079010 001 May 26, 2009  
AA ! WOCKHARDT EQ 15MG BASE/5ML A076895 001 Oct 04, 2004  
AA ! WOCKHARDT BIO AG EQ 5MG BASE/5ML A075099 001 Jun 28, 2002  
 ! MISSION PHARMA EQ 25MG BASE/5ML A091396 001 Sep 13, 2010

SOLUTION/DROPS;OPHTHALMIC

PREDNISOLONE SODIUM PHOSPHATE  
 ! BAUSCH AND LOMB EQ 0.9% PHOSPHATE A040070 001 Jul 29, 1994

TABLET, ORALLY DISINTEGRATING;ORAL

ORAPRED ODT

AB + CONCORDIA PHARMS INC EQ 10MG BASE N021959 001 Jun 01, 2006

AB + EQ 15MG BASE

AB +! EQ 30MG BASE N021959 002 Jun 01, 2006

PREDNISOLONE SODIUM PHOSPHATE

AB MYLAN PHARMS INC EQ 10MG BASE A202179 001 Apr 10, 2013  
AB EQ 15MG BASE A202179 002 Apr 10, 2013  
AB EQ 30MG BASE A202179 003 Apr 10, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-354 (of 436)

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

<u>AT</u>	!	BAUSCH AND LOMB	<u>EQ 0.23% PHOSPHATE,10%</u>
<u>AT</u>		SANDOZ INC	<u>EQ 0.23% PHOSPHATE,10%</u>

A074449 001 Dec 29, 1995  
A073630 001 May 27, 1993

PREDNISONE

SOLUTION;ORAL

PREDNISONE

!	WEST-WARD PHARMS	5MG/5ML
INT		

A088703 001 Nov 08, 1984  
A088810 001 Feb 20, 1985

TABLET;ORAL

PREDNISONE

<u>AB</u>	HIKMA PHARMS	<u>2.5MG</u>
<u>AB</u>		<u>50MG</u>
<u>AB</u>	JUBILANT CADISTA	<u>1MG</u>
<u>AB</u>		<u>5MG</u>
<u>AB</u>		<u>10MG</u>
<u>AB</u>		<u>20MG</u>
<u>AB</u>	MUTUAL PHARM	<u>5MG</u>
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>
<u>AB</u>		<u>10MG</u>
<u>AB</u>		<u>20MG</u>
<u>AB</u>	SUN PHARM INDUSTRIES	<u>10MG</u>
<u>AB</u>		<u>20MG</u>
<u>AB</u>	VINTAGE PHARMS	<u>1MG</u>
<u>AB</u>		<u>2.5MG</u>
<u>AB</u>		<u>5MG</u>
<u>AB</u>		<u>10MG</u>
<u>AB</u>		<u>20MG</u>
<u>AB</u>	WATSON LABS	<u>5MG</u>
<u>AB</u>		<u>10MG</u>
<u>AB</u>		<u>20MG</u>
<u>AB</u>	!	WEST-WARD PHARMS
	INT	<u>1MG</u>
<u>AB</u>	!	
		<u>2.5MG</u>
<u>AB</u>	!	
		<u>5MG</u>
<u>AB</u>	!	
		<u>10MG</u>
<u>AB</u>	!	
		<u>20MG</u>
<u>AB</u>	!	
		<u>50MG</u>

A040538 001 Jan 08, 2004  
A088465 001 Jun 01, 1984  
A040611 001 Jun 06, 2005  
A040362 002 Aug 29, 2001  
A040362 001 Aug 29, 2001  
A040362 003 Jun 29, 2005  
A089245 001 Dec 04, 1985  
A080292 001  
A088832 001 Dec 04, 1985  
A083677 001  
A089246 001 Dec 04, 1985  
A089247 001 Dec 04, 1985  
A040584 001 Dec 21, 2004  
A040581 001 Dec 21, 2004  
A040256 001 Jul 12, 2002  
A040256 002 Jul 12, 2002  
A040392 001 Feb 12, 2003  
A080356 001  
A085162 001  
A085161 001  
A087800 001 Apr 22, 1982  
A087801 001 Apr 22, 1982  
A080352 001  
A084122 001  
A087342 001  
A084283 001

TABLET, DELAYED RELEASE;ORAL

PREDNISONE

<u>AB</u>	ACTAVIS LABS FL INC	<u>1MG</u>
<u>AB</u>		<u>2MG</u>
<u>AB</u>		<u>5MG</u>

A204867 001 Apr 25, 2017  
A204867 002 Apr 25, 2017  
A204867 003 Apr 25, 2017

RAYOS

<u>AB</u>	+	HORIZON PHARMA	<u>1MG</u>
<u>AB</u>	+		<u>2MG</u>
<u>AB</u>	+		<u>5MG</u>

N202020 001 Jul 26, 2012  
N202020 002 Jul 26, 2012  
N202020 003 Jul 26, 2012

PREGABALIN

CAPSULE;ORAL

LYRICA

+	PF PRISM CV	25MG
+		50MG
+		75MG
+		100MG
+		150MG
+		200MG
+		225MG
+		300MG

N021446 001 Dec 30, 2004  
N021446 002 Dec 30, 2004  
N021446 003 Dec 30, 2004  
N021446 004 Dec 30, 2004  
N021446 005 Dec 30, 2004  
N021446 006 Dec 30, 2004  
N021446 007 Dec 30, 2004  
N021446 008 Dec 30, 2004

SOLUTION;ORAL

LYRICA

+	PF PRISM CV	20MG/ML
---	-------------	---------

N022488 001 Jan 04, 2010

TABLET, EXTENDED RELEASE;ORAL

LYRICA CR

+	Pfizer Inc	82.5MG
+		165MG
+		330MG

N209501 001 Oct 11, 2017  
N209501 002 Oct 11, 2017  
N209501 003 Oct 11, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-355 (of 436)

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION  
 PRILOCAINE HYDROCHLORIDE  
 ! SEPTODONT INC 4%

A079235 001 Sep 29, 2010

PRIMAQUINE PHOSPHATE

TABLET; ORAL

PRIMAQUINE

AB +! SANOFI AVENTIS US EQ 15MG BASE

N008316 001

PRIMAQUINE PHOSPHATE

AB ALVOGEN INC EQ 15MG BASE

A203924 001 Feb 03, 2014

AB BAYSHORE PHARMS LLC EQ 15MG BASE

A204476 001 Feb 25, 2014

AB INGENUS PHARMS NJ EQ 15MG BASE

A206043 001 Jun 23, 2016

PRIMIDONE

TABLET; ORAL

MYSOLINE

AB +! VALEANT 50MG

N009170 003

AB + 250MG

N009170 002

PRIMIDONE

AB AMNEAL PHARM 50MG

A040866 001 Apr 23, 2008

AB 250MG

A040866 002 Apr 23, 2008

AB ANDA REPOSITORY 50MG

A040626 001 Sep 29, 2005

AB 250MG

A040626 002 Sep 29, 2005

AB HIKMA INTL PHARMS 250MG

A040667 002 Jul 27, 2006

AB LANNETT 50MG

A084903 002 May 24, 2001

AB 250MG

A084903 001

AB VINTAGE PHARMS 50MG

A040586 001 Feb 24, 2005

AB 250MG

A040586 002 Feb 24, 2005

AB WATSON LABS 250MG

A083551 001

PROBENECID

TABLET; ORAL

PROBALAN

AB LANNETT 500MG

A080966 001

PROBENECID

AB ! MYLAN 500MG

A084211 002

AB WATSON LABS TEVA 500MG

A084442 004 Mar 29, 1983

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

AP ! HOSPIRA 100MG/ML

A089069 001 Feb 12, 1986

AP INTL MEDICATION 100MG/ML

A088636 001 Jul 31, 1984

AP NEXUS PHARMS 100MG/ML

A206332 001 Oct 13, 2017

AP ! HOSPIRA 500MG/ML

A206332 002 Oct 13, 2017

A089070 001 Feb 12, 1986

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL

MATULANE

+! LEADIANTE BIOSCI INC EQ 50MG BASE

N016785 001

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPRO

AB PADDICK LLC 25MG

A040246 001 Jun 28, 2000

PROCHLORPERAZINE

AB ! G AND W LABS 25MG

A040058 001 Nov 24, 1993

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

AP ! EMCURE PHARMS LTD EQ 5MG BASE/ML

A204147 001 Oct 15, 2013

AP WEST-WARD PHARMS INT EQ 5MG BASE/ML

A089903 001 Aug 29, 1989

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

AB MYLAN EQ 5MG BASE

A040185 002 Oct 28, 1996

AB EQ 10MG BASE

A040185 001 Oct 28, 1996

AB SANDOZ EQ 5MG BASE

A040101 001 Jul 19, 1996

AB ! TEVA PHARMS EQ 10MG BASE

A040101 002 Jul 19, 1996

AB TEVA PHARMS EQ 5MG BASE

A040120 001 Jul 11, 1996

AB TEVA PHARMS EQ 10MG BASE

A040120 002 Jul 11, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-356 (of 436)

PROCHLORPERAZINE MALEATE

TABLET;ORAL

PROCOMP

<u>AB</u>	JUBILANT CADISTA	<u>EQ 5MG BASE</u>
<u>AB</u>		<u>EQ 10MG BASE</u>

<u>A040268</u>	<u>001</u>	Feb 27, 1998
<u>A040268</u>	<u>002</u>	Feb 27, 1998

PROGESTERONE

CAPSULE;ORAL

PROGESTERONE

<u>AB</u>	AMNEAL PHARMS NY	<u>100MG</u>
<u>AB</u>		<u>200MG</u>
<u>AB</u>	BIONPHARMA INC	<u>100MG</u>
<u>AB</u>		<u>200MG</u>
<u>AB</u>	DR REDDYS LABS INC	<u>100MG</u>
<u>AB</u>		<u>200MG</u>
<u>AB</u>	SANDOZ INC	<u>100MG</u>
<u>AB</u>		<u>200MG</u>
<u>AB</u>	SOFGEN PHARMS	<u>100MG</u>
<u>AB</u>		<u>200MG</u>
<u>AB</u>	TEVA PHARMS	<u>100MG</u>
<u>AB</u>		<u>200MG</u>

<u>A207724</u>	<u>001</u>	Sep 07, 2017
<u>A207724</u>	<u>002</u>	Sep 07, 2017
<u>A200900</u>	<u>001</u>	Aug 16, 2013
<u>A200900</u>	<u>002</u>	Aug 16, 2013
<u>A208801</u>	<u>001</u>	Feb 28, 2017
<u>A208801</u>	<u>002</u>	Feb 28, 2017
<u>A205229</u>	<u>001</u>	Oct 20, 2017
<u>A205229</u>	<u>002</u>	Oct 20, 2017
<u>A200456</u>	<u>001</u>	Sep 28, 2012
<u>A200456</u>	<u>002</u>	Sep 28, 2012
<u>A202121</u>	<u>001</u>	Feb 29, 2012
<u>A202121</u>	<u>002</u>	Feb 29, 2012

PROMETRIUM

<u>AB</u>	+	VIRTUS PHARMS	<u>100MG</u>
<u>AB</u>	+!		<u>200MG</u>

<u>N019781</u>	<u>001</u>	May 14, 1998
<u>N019781</u>	<u>002</u>	Oct 15, 1999

GEL;VAGINAL

CRINONE

+!	ALLERGAN SALES LLC	4%
+!		8%

N020701	001	Jul 31, 1997
N020701	002	Jul 31, 1997

INJECTABLE;INJECTION

PROGESTERONE

<u>AO</u>	+!	ACTAVIS LABS UT INC	<u>50MG/ML</u>
<u>AO</u>		FRESENIUS KABI USA	<u>50MG/ML</u>
<u>AO</u>		HIKMA FARMACEUTICA	<u>50MG/ML</u>
<u>AO</u>		LUITPOLD	<u>50MG/ML</u>

<u>N017362</u>	<u>002</u>	
<u>A075906</u>	<u>001</u>	Apr 25, 2001
<u>A091033</u>	<u>001</u>	Oct 28, 2010
<u>A090845</u>	<u>001</u>	Jun 22, 2009

INSERT;VAGINAL

ENDOMETRIN

+!	FERRING	100MG
----	---------	-------

N022057	001	Jun 21, 2007
---------	-----	--------------

PROMETHAZINE HYDROCHLORIDE

INJECTABLE;INJECTION

PROMETHAZINE HYDROCHLORIDE

<u>AP</u>	!	WEST-WARD PHARMS INT	<u>25MG/ML</u>
<u>AP</u>	!		<u>50MG/ML</u>
<u>AP</u>		X-GEN PHARMS	<u>25MG/ML</u>
<u>AP</u>			<u>50MG/ML</u>

<u>A083312</u>	<u>001</u>	
<u>A083312</u>	<u>002</u>	
<u>A040737</u>	<u>001</u>	Apr 24, 2008
<u>A040737</u>	<u>002</u>	Apr 24, 2008

SUPPOSITORY;RECTAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	G AND W LABS	<u>12.5MG</u>
<u>AB</u>	!	<u>25MG</u>
<u>AB</u>	PERRIGO NEW YORK	<u>12.5MG</u>
<u>AB</u>		<u>25MG</u>
<u>AB</u>	TARO	<u>12.5MG</u>
<u>AB</u>		<u>25MG</u>
<u>AB</u>	WATSON LABS INC	<u>12.5MG</u>
<u>AB</u>		<u>25MG</u>

<u>A040428</u>	<u>002</u>	Mar 31, 2003
<u>A040428</u>	<u>001</u>	Feb 05, 2002
<u>A040500</u>	<u>001</u>	Jun 30, 2003
<u>A040500</u>	<u>002</u>	Jun 30, 2003
<u>A040603</u>	<u>001</u>	Oct 26, 2006
<u>A040603</u>	<u>002</u>	Oct 26, 2006
<u>A040479</u>	<u>001</u>	Jun 24, 2003
<u>A040479</u>	<u>002</u>	Jun 24, 2003

PROMETHEGAN

!	G AND W LABS	50MG
---	--------------	------

A087165	001	Aug 14, 1987
---------	-----	--------------

SYRUP;ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AA</u>	AMNEAL PHARMS	<u>6.25MG/5ML</u>
<u>AA</u>	HI TECH PHARMA	<u>6.25MG/5ML</u>
<u>AA</u>	NOSTRUM LABS INC	<u>6.25MG/5ML</u>
<u>AA</u>	TARO	<u>6.25MG/5ML</u>
<u>AA</u>	TRIS PHARMA INC	<u>6.25MG/5ML</u>
<u>AA</u>	VINTAGE	<u>6.25MG/5ML</u>

<u>A040882</u>	<u>001</u>	Dec 30, 2009
<u>A040026</u>	<u>001</u>	Sep 25, 1998
<u>A040891</u>	<u>001</u>	Mar 13, 2009
<u>A040718</u>	<u>001</u>	Apr 04, 2007
<u>A091675</u>	<u>001</u>	Jun 28, 2012
<u>A040643</u>	<u>001</u>	Apr 26, 2006

PROMETHAZINE PLAIN

<u>AA</u>	!	WOCKHARDT BIO AG	<u>6.25MG/5ML</u>
-----------	---	------------------	-------------------

<u>A087953</u>	<u>001</u>	Nov 15, 1982
----------------	------------	--------------

TABLET;ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	AMNEAL PHARMS NY	<u>12.5MG</u>
<u>AB</u>		<u>25MG</u>
<u>AB</u>		<u>50MG</u>
<u>AB</u>	HERITAGE PHARMA	<u>12.5MG</u>

<u>A091179</u>	<u>001</u>	Dec 13, 2010
<u>A091179</u>	<u>002</u>	Dec 13, 2010
<u>A091179</u>	<u>003</u>	Dec 13, 2010
<u>A040673</u>	<u>001</u>	Mar 05, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-357 (of 436)

PROMETHAZINE HYDROCHLORIDE

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>		<u>25MG</u>	<u>A040673 002</u>	Mar 05, 2008
<u>AB</u>		<u>50MG</u>	<u>A040673 003</u>	Mar 05, 2008
<u>AB</u>	KVK TECH	<u>12.5MG</u>	<u>A040712 002</u>	May 04, 2007
<u>AB</u>		<u>25MG</u>	<u>A040712 001</u>	Jul 31, 2006
<u>AB</u>		<u>50MG</u>	<u>A040712 003</u>	Jul 31, 2006
<u>AB</u>	PRINSTON INC	<u>12.5MG</u>	<u>A040622 001</u>	Jul 18, 2006
<u>AB</u>		<u>25MG</u>	<u>A040622 002</u>	Jul 18, 2006
<u>AB</u>		<u>50MG</u>	<u>A040622 003</u>	Jul 18, 2006
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A084176 003</u>	
<u>AB</u>		<u>50MG</u>	<u>A084176 001</u>	
<u>AB</u>	STRIDES PHARMA	<u>12.5MG</u>	<u>A209177 001</u>	Jun 30, 2017
<u>AB</u>		<u>25MG</u>	<u>A209177 002</u>	Jun 30, 2017
<u>AB</u>		<u>50MG</u>	<u>A209177 003</u>	Jun 30, 2017
<u>AB</u>	SUN PHARM INDS INC	<u>12.5MG</u>	<u>A040863 001</u>	Dec 30, 2008
<u>AB</u>		<u>25MG</u>	<u>A040863 002</u>	Dec 30, 2008
<u>AB</u>		<u>50MG</u>	<u>A040863 003</u>	Dec 30, 2008
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>A083426 001</u>	
<u>AB</u>		<u>50MG</u>	<u>A083711 001</u>	
<u>AB</u>	ZYDUS PHARMS USA	<u>12.5MG</u>	<u>A040596 001</u>	Nov 18, 2005
<u>AB</u>		<u>25MG</u>	<u>A040596 002</u>	Nov 18, 2005
<u>AB</u>		<u>50MG</u>	<u>A040596 003</u>	Nov 18, 2005
<u>AB</u>	IMPAX LABS	12.5MG	A040791 002	Feb 12, 2008
		25MG	A040791 003	Feb 12, 2008

PROPAFENONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	GLENMARK PHARMS LTD	<u>225MG</u>	<u>A205268 001</u>	Sep 08, 2017
<u>AB</u>		<u>325MG</u>	<u>A205268 002</u>	Sep 08, 2017
<u>AB</u>		<u>425MG</u>	<u>A205268 003</u>	Sep 08, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>225MG</u>	<u>A203803 001</u>	Apr 29, 2016
<u>AB</u>		<u>325MG</u>	<u>A203803 002</u>	Apr 29, 2016
<u>AB</u>		<u>425MG</u>	<u>A203803 003</u>	Apr 29, 2016
<u>AB</u>	PAR PHARM	<u>225MG</u>	<u>A078540 001</u>	Oct 18, 2010
<u>AB</u>		<u>325MG</u>	<u>A078540 002</u>	Oct 18, 2010
<u>AB</u>		<u>425MG</u>	<u>A078540 003</u>	Oct 18, 2010
<u>AB</u>	WATSON LABS INC	<u>225MG</u>	<u>A202688 001</u>	Aug 24, 2015
<u>AB</u>		<u>325MG</u>	<u>A202688 002</u>	Aug 24, 2015
<u>AB</u>		<u>425MG</u>	<u>A202688 003</u>	Aug 24, 2015
<b>RYTHMOL SR</b>				
<u>AB</u>	+ GLAXOSMITHKLINE LLC	<u>225MG</u>	<u>N021416 001</u>	Sep 04, 2003
<u>AB</u>	+	<u>325MG</u>	<u>N021416 002</u>	Sep 04, 2003
<u>AB</u>	+!	<u>425MG</u>	<u>N021416 003</u>	Sep 04, 2003

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	ANI PHARMS INC	<u>150MG</u>	<u>A076550 001</u>	Apr 23, 2004
<u>AB</u>		<u>225MG</u>	<u>A076550 002</u>	Apr 23, 2004
<u>AB</u>		<u>300MG</u>	<u>A076550 003</u>	Apr 23, 2004
<u>AB</u>	AUROBINDO PHARMA LTD	<u>150MG</u>	<u>A202445 001</u>	May 11, 2016
<u>AB</u>		<u>225MG</u>	<u>A202445 002</u>	May 11, 2016
<u>AB</u>		<u>300MG</u>	<u>A202445 003</u>	May 11, 2016
<u>AB</u>	SUN PHARM INDUSTRIES	<u>150MG</u>	<u>A075998 001</u>	Nov 29, 2001
<u>AB</u>		<u>225MG</u>	<u>A075998 002</u>	Nov 29, 2001
<u>AB</u>		<u>300MG</u>	<u>A075998 003</u>	Nov 29, 2001
<u>AB</u>	VINTAGE PHARMS	<u>150MG</u>	<u>A075938 001</u>	Oct 17, 2002
<u>AB</u>		<u>225MG</u>	<u>A075938 002</u>	Oct 17, 2002
<u>AB</u>		<u>300MG</u>	<u>A075938 003</u>	Oct 17, 2002
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>A075203 001</u>	Oct 24, 2000
<u>AB</u>		<u>225MG</u>	<u>A075203 002</u>	Oct 24, 2000
<b>RYTHMOL</b>				
<u>AB</u>	+ GLAXOSMITHKLINE LLC	<u>150MG</u>	<u>N019151 001</u>	Nov 27, 1989
<u>AB</u>	+	<u>225MG</u>	<u>N019151 003</u>	Nov 20, 1992
<u>AB</u>	+!	<u>300MG</u>	<u>N019151 002</u>	Nov 27, 1989

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-358 (of 436)

PROPANTHELINE BROMIDE

TABLET;ORAL

PROPANTHELINE BROMIDE  
 ! WEST-WARD PHARMS 15MG

A080927 002

PROPARACAIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

ALCAINE

**AT** ! NOVARTIS PHARMS 0.5%  
 CORP

**A080027 001**

PROPARACAIN HYDROCHLORIDE

**AT** AKORN INC 0.5%  
**AT** BAUSCH AND LOMB 0.5%

**A040277 001** Mar 16, 2000  
**A040074 001** Sep 29, 1995

PROPOFOL

INJECTABLE;INJECTION

DIPRIVAN

**AB** +! FRESENIUS KABI USA 10MG/ML

**N019627 002** Jun 11, 1996

PROPOFOL

**AB** HOSPIRA 10MG/ML  
**AB** SAGENT PHARMS 10MG/ML  
**AB** WATSON LABS INC 10MG/ML

**A077908 001** Mar 17, 2006  
**A075102 001** Jan 04, 1999  
**A205307 001** Dec 22, 2015

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

Inderal LA

**AB** + ANI PHARMS INC 60MG  
**AB** + 80MG  
**AB** + 120MG  
**AB** +! 160MG

**N018553 004** Mar 18, 1987  
**N018553 002** Apr 19, 1983  
**N018553 003** Apr 19, 1983  
**N018553 001** Apr 19, 1983

PROPRANOLOL HYDROCHLORIDE

**AB** ACTAVIS ELIZABETH 60MG  
**AB** 80MG  
**AB** 120MG  
**AB** 160MG

**A078494 001** Aug 10, 2007  
**A078494 002** Aug 10, 2007  
**A078494 003** Aug 10, 2007  
**A078494 004** Aug 10, 2007

**AB** MYLAN 60MG  
**AB** 80MG  
**AB** 120MG  
**AB** 160MG

**A078022 001** Feb 15, 2007  
**A078022 002** Feb 15, 2007  
**A078022 003** Feb 15, 2007  
**A078022 004** Feb 15, 2007

**AB** NORTEC DEV ASSOC 60MG  
**AB** 80MG  
**AB** 120MG  
**AB** 160MG

**A078065 001** Jan 26, 2007  
**A078065 002** Jan 26, 2007  
**A078065 003** Jan 26, 2007  
**A078065 004** Jan 26, 2007

**AB** RP SCHERER 60MG  
**AB** 80MG  
**AB** 120MG  
**AB** 160MG

**A078703 001** Jul 15, 2011  
**A078703 002** Jul 15, 2011  
**A078703 003** Jul 15, 2011  
**A078703 004** Jul 15, 2011

**AB** ZYDUS PHARMS USA 60MG  
 INC  
**AB** 80MG  
**AB** 120MG  
**AB** 160MG

**A090321 001** Mar 25, 2011  
**A090321 002** Mar 25, 2011  
**A090321 003** Mar 25, 2011  
**A090321 004** Mar 25, 2011

INNOPRAN XL

BX ANI PHARMS INC 80MG  
 BX 120MG

N021438 001 Mar 12, 2003  
 N021438 002 Mar 12, 2003

INJECTABLE;INJECTION

PROPRANOLOL HYDROCHLORIDE

**AP** ATHENEX INC 1MG/ML  
**AP** +! BAXTER HLTHCARE CORP 1MG/ML

**A075792 001** Aug 29, 2000  
**N016419 001**

**AP** FRESENIUS KABI USA 1MG/ML  
**AP** HIKMA FARMACEUTICA 1MG/ML

**A075826 001** Aug 31, 2001  
**A077760 001** Jan 31, 2008

SOLUTION;ORAL

HEMANGEOL

+! PIERRE FABRE DERMA 4.28MG/ML  
 PROPRANOLOL HYDROCHLORIDE  
 ! WEST-WARD PHARMS 20MG/5ML  
 INT  
 ! 40MG/5ML

N205410 001 Mar 14, 2014  
 A070979 001 May 15, 1987  
 A070690 001 May 15, 1987

TABLET;ORAL

PROPRANOLOL HYDROCHLORIDE

**AB** IMPAX LABS INC 10MG  
**AB** 20MG  
**AB** 40MG

**A071972 001** Apr 06, 1988  
**A071973 001** Apr 06, 1988  
**A071974 001** Apr 06, 1988

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-359 (of 436)

PROPRANOLOL HYDROCHLORIDE

TABLET;ORAL

PROPRANOLOL HYDROCHLORIDE

<u>AB</u>		<u>60MG</u>	<u>A071975</u> <u>001</u>	Apr 06, 1988
<u>AB</u>	!	<u>80MG</u>	<u>A071976</u> <u>001</u>	Apr 06, 1988
<u>AB</u>	IPCA LABS LTD	<u>10MG</u>	<u>A078955</u> <u>001</u>	Jun 02, 2008
<u>AB</u>		<u>20MG</u>	<u>A078955</u> <u>002</u>	Jun 02, 2008
<u>AB</u>		<u>40MG</u>	<u>A078955</u> <u>003</u>	Jun 02, 2008
<u>AB</u>		<u>60MG</u>	<u>A078955</u> <u>004</u>	Jun 02, 2008
<u>AB</u>		<u>80MG</u>	<u>A078955</u> <u>005</u>	Jun 02, 2008
<u>AB</u>	MYLAN	<u>10MG</u>	<u>A070213</u> <u>002</u>	Nov 19, 1985
<u>AB</u>		<u>20MG</u>	<u>A070213</u> <u>003</u>	Nov 19, 1985
<u>AB</u>		<u>40MG</u>	<u>A070213</u> <u>001</u>	Nov 19, 1985
<u>AB</u>		<u>60MG</u>	<u>A070213</u> <u>005</u>	Apr 08, 2011
<u>AB</u>		<u>80MG</u>	<u>A070213</u> <u>004</u>	Nov 19, 1985
<u>AB</u>	NORTHSTAR HLTHCARE	<u>10MG</u>	<u>A078213</u> <u>001</u>	Jan 10, 2008
<u>AB</u>		<u>20MG</u>	<u>A078213</u> <u>002</u>	Jan 10, 2008
<u>AB</u>		<u>40MG</u>	<u>A078213</u> <u>003</u>	Jan 10, 2008
<u>AB</u>		<u>60MG</u>	<u>A078213</u> <u>004</u>	Jan 10, 2008
<u>AB</u>		<u>80MG</u>	<u>A078213</u> <u>005</u>	Jan 10, 2008
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>A070221</u> <u>002</u>	Aug 01, 1986
<u>AB</u>		<u>20MG</u>	<u>A070221</u> <u>003</u>	Aug 01, 1986
<u>AB</u>		<u>40MG</u>	<u>A070219</u> <u>001</u>	Aug 01, 1986
<u>AB</u>		<u>60MG</u>	<u>A070221</u> <u>004</u>	Aug 01, 1986
<u>AB</u>		<u>80MG</u>	<u>A070221</u> <u>005</u>	Sep 24, 1986
<u>AB</u>		<u>10MG</u>	<u>A070221</u> <u>001</u>	Apr 14, 1986
<u>AB</u>		<u>20MG</u>	<u>A070175</u> <u>001</u>	May 13, 1986
<u>AB</u>		<u>40MG</u>	<u>A070176</u> <u>001</u>	May 13, 1986
<u>AB</u>		<u>80MG</u>	<u>A070177</u> <u>001</u>	May 13, 1986
<u>AB</u>			<u>A070178</u> <u>001</u>	May 13, 1986

PROPYLTHIOURACIL

TABLET;ORAL

PROPYLTHIOURACIL

BD	ACTAVIS ELIZABETH	50MG	A080172 001
BD	+! DAVA PHARMS INC	50MG	N006188 001

PROTAMINE SULFATE

INJECTABLE;INJECTION

PROTAMINE SULFATE

!	FRESENIUS KABI USA	10MG/ML	A089454 001
---	--------------------	---------	-------------

Apr 07, 1987

PROTRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

PROTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	EPIC PHARMA LLC	<u>5MG</u>	<u>A202220</u> <u>001</u>	Nov 19, 2012
<u>AB</u>		<u>10MG</u>	<u>A202220</u> <u>002</u>	Nov 19, 2012
<u>AB</u>	SIGMAPHARM LABS LLC	<u>5MG</u>	<u>A090462</u> <u>001</u>	May 03, 2010
<u>AB</u>		<u>10MG</u>	<u>A090462</u> <u>002</u>	May 03, 2010
<u>AB</u>	WEST-WARD PHARMS INT	<u>5MG</u>	<u>A078913</u> <u>001</u>	Sep 16, 2008
<u>AB</u>		<u>10MG</u>	<u>A078913</u> <u>002</u>	Sep 16, 2008
<u>VIVACTIL</u>				
<u>AB</u>	ODYSSEY PHARMS	<u>5MG</u>	<u>A073644</u> <u>001</u>	Aug 24, 1995
<u>AB</u>	!	<u>10MG</u>	<u>A073645</u> <u>001</u>	Aug 24, 1995

PYRAZINAMIDE

TABLET;ORAL

PYRAZINAMIDE

<u>AB</u>	AKORN	<u>500MG</u>	<u>A081319</u> <u>001</u>	Jun 30, 1992
<u>AB</u>	! DAVA PHARMS INC	<u>500MG</u>	<u>A080157</u> <u>001</u>	

PYRIDOSTIGMINE BROMIDE

INJECTABLE;INJECTION

MESTINON

<u>AP</u>	+! VALEANT PHARM INTL	<u>5MG/ML</u>	<u>N009830</u> <u>001</u>
-----------	-----------------------	---------------	---------------------------

REGONOL

<u>AP</u>	SANDOZ INC	<u>5MG/ML</u>	<u>N017398</u> <u>001</u>
-----------	------------	---------------	---------------------------

SYRUP;ORAL

MESTINON

+!	VALEANT PHARMS	60MG/5ML	N015193 001
----	----------------	----------	-------------

TABLET;ORAL

MESTINON

<u>AB</u>	+! VALEANT PHARMS LLC	<u>60MG</u>	<u>N009829</u> <u>002</u>
-----------	-----------------------	-------------	---------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-360 (of 436)

PYRIDOSTIGMINE BROMIDE

TABLET;ORAL

**PYRIDOSTIGMINE BROMIDE**

<b>AB</b>	IMPAK LABS	<u>60MG</u>	<b>A040502 001</b>	Apr 24, 2003
<b>AB</b>	ZYDUS PHARMS USA INC	<u>60MG</u>	<b>A205650 001</b>	Jun 22, 2015

TABLET, EXTENDED RELEASE;ORAL

**MESTINON**

<b>AB</b>	+!	VALEANT PHARMS LLC	<u>180MG</u>	<b>N011665 001</b>
-----------	----	--------------------	--------------	--------------------

**PYRIDOSTIGMINE BROMIDE**

<b>AB</b>	ALVOGEN MALTA	<u>180MG</u>	<b>A204737 001</b>	Jun 26, 2015
<b>AB</b>	IMPAK LABS INC	<u>180MG</u>	<b>A203184 001</b>	Sep 15, 2015
<b>AB</b>	KINEDEXE UK	<u>180MG</u>	<b>A205464 001</b>	Aug 15, 2017

PYRIDOXINE HYDROCHLORIDE

INJECTABLE;INJECTION

**PYRIDOXINE HYDROCHLORIDE**

<b>AP</b>	! FRESENIUS KABI USA	<u>100MG/ML</u>	<b>A080618 001</b>	
<b>AP</b>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<b>A204879 001</b>	Jul 14, 2016

PYRIMETHAMINE

TABLET;ORAL

DARAPRIM

+!	VYERA PHARMS LLC	<u>25MG</u>	N008578 001
----	------------------	-------------	-------------

QUAZEPAM

TABLET;ORAL

DORAL

+!	CUTIS HEALTH LLC	<u>15MG</u>	N018708 001
----	------------------	-------------	-------------

Dec 27, 1985

QUETIAPINE FUMARATE

TABLET;ORAL

**QUETIAPINE FUMARATE**

<b>AB</b>	ACCORD HLTHCARE	<u>EQ 25MG BASE</u>	<b>A202152 001</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 50MG BASE</u>	<b>A202152 002</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 100MG BASE</u>	<b>A202152 003</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 200MG BASE</u>	<b>A202152 004</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A202152 005</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 400MG BASE</u>	<b>A202152 006</b>	Mar 27, 2012
<b>AB</b>	ALEMBIC PHARMS LTD	<u>EQ 25MG BASE</u>	<b>A203390 001</b>	Oct 28, 2014
<b>AB</b>		<u>EQ 50MG BASE</u>	<b>A203390 002</b>	Oct 28, 2014
<b>AB</b>		<u>EQ 100MG BASE</u>	<b>A203390 003</b>	Oct 28, 2014
<b>AB</b>		<u>EQ 200MG BASE</u>	<b>A203390 004</b>	Oct 28, 2014
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A203390 005</b>	Oct 28, 2014
<b>AB</b>		<u>EQ 400MG BASE</u>	<b>A203390 006</b>	Oct 28, 2014
<b>AB</b>	ALKEM LABS LTD	<u>EQ 25MG BASE</u>	<b>A201504 001</b>	Feb 12, 2013
<b>AB</b>		<u>EQ 50MG BASE</u>	<b>A201504 002</b>	Feb 12, 2013
<b>AB</b>		<u>EQ 100MG BASE</u>	<b>A201504 003</b>	Feb 12, 2013
<b>AB</b>		<u>EQ 150MG BASE</u>	<b>A201504 004</b>	Feb 12, 2013
<b>AB</b>		<u>EQ 200MG BASE</u>	<b>A201504 005</b>	Feb 12, 2013
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A201504 006</b>	Feb 12, 2013
<b>AB</b>		<u>EQ 400MG BASE</u>	<b>A201504 007</b>	Feb 12, 2013
<b>AB</b>	APOTEX INC	<u>EQ 25MG BASE</u>	<b>A090960 001</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 50MG BASE</u>	<b>A090960 002</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 100MG BASE</u>	<b>A090960 003</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 200MG BASE</u>	<b>A090960 004</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A090960 005</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 400MG BASE</u>	<b>A090960 006</b>	Mar 27, 2012
<b>AB</b>	AUROBINDO PHARMA LTD	<u>EQ 25MG BASE</u>	<b>A091388 001</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 50MG BASE</u>	<b>A091388 002</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 100MG BASE</u>	<b>A091388 003</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 150MG BASE</u>	<b>A091388 004</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 200MG BASE</u>	<b>A091388 005</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A091388 006</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 400MG BASE</u>	<b>A091388 007</b>	Mar 27, 2012
<b>AB</b>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<b>A077380 001</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 50MG BASE</u>	<b>A077380 002</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 100MG BASE</u>	<b>A077380 003</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 150MG BASE</u>	<b>A077380 004</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 200MG BASE</u>	<b>A077380 005</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A077380 006</b>	Mar 27, 2012
<b>AB</b>		<u>EQ 400MG BASE</u>	<b>A077380 007</b>	Mar 27, 2012
<b>AB</b>	JUBILANT GENERICS	<u>EQ 25MG BASE</u>	<b>A203150 001</b>	Nov 26, 2013
<b>AB</b>	LUPIN LTD	<u>EQ 25MG BASE</u>	<b>A201109 001</b>	Mar 27, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-361 (of 436)

**QUETIAPINE FUMARATE**

TABLET;ORAL

**QUETIAPINE FUMARATE**

<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A201109 002</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A201109 003</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A201109 004</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A201109 005</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A201109 006</u></b>	Mar 27, 2012
<b><u>AB</u></b>	MACLEODS PHARMS LTD	<b><u>EQ 25MG BASE</u></b>	<b><u>A203359 001</u></b>	May 17, 2016
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A203359 002</u></b>	May 17, 2016
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A203359 003</u></b>	May 17, 2016
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A203359 004</u></b>	May 17, 2016
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A203359 005</u></b>	May 17, 2016
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A203359 006</u></b>	May 17, 2016
<b><u>AB</u></b>	SANDOZ	<b><u>EQ 25MG BASE</u></b>	<b><u>A078679 001</u></b>	Dec 14, 2012
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A078679 002</u></b>	Dec 14, 2012
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A078679 003</u></b>	Dec 14, 2012
<b><u>AB</u></b>		<b><u>EQ 150MG BASE</u></b>	<b><u>A078679 004</u></b>	Dec 14, 2012
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A078679 005</u></b>	Dec 14, 2012
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A078679 006</u></b>	Dec 14, 2012
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A078679 007</u></b>	Dec 14, 2012
<b><u>AB</u></b>	SUN PHARMA GLOBAL	<b><u>EQ 25MG BASE</u></b>	<b><u>A201190 001</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A201190 002</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A201190 003</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A201190 004</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A201190 005</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A201190 006</u></b>	Mar 27, 2012
<b><u>AB</u></b>	TEVA PHARMS	<b><u>EQ 25MG BASE</u></b>	<b><u>A077745 001</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A077745 002</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A077745 003</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 150MG BASE</u></b>	<b><u>A077745 004</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A077745 005</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A077745 006</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A077745 007</u></b>	Mar 27, 2012
<b><u>AB</u></b>	TORRENT PHARMS LTD	<b><u>EQ 25MG BASE</u></b>	<b><u>A200363 001</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A200363 002</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A200363 003</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A200363 004</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A200363 005</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A200363 006</u></b>	Mar 27, 2012
<b><u>AB</u></b>	UNICHEM LABS LTD	<b><u>EQ 25MG BASE</u></b>	<b><u>A202674 001</u></b>	Mar 08, 2016
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A202674 002</u></b>	Mar 08, 2016
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A202674 003</u></b>	Mar 08, 2016
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A202674 004</u></b>	Mar 08, 2016
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A202674 005</u></b>	Mar 08, 2016
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A202674 006</u></b>	Mar 08, 2016
<b><u>AB</u></b>	WEST-WARD PHARMS INT	<b><u>EQ 25MG BASE</u></b>	<b><u>A202674 007</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 50MG BASE</u></b>	<b><u>A090749 001</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 100MG BASE</u></b>	<b><u>A090749 002</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A090749 003</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A090749 004</u></b>	Mar 27, 2012
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A090749 005</u></b>	Mar 27, 2012

**SEROQUEL**

<b><u>AB</u></b>	+ ! ASTRAZENECA PHARMS	<b><u>EQ 25MG BASE</u></b>	<b><u>N020639 001</u></b>	Sep 26, 1997
<b><u>AB</u></b>	+	<b><u>EQ 50MG BASE</u></b>	<b><u>N020639 007</u></b>	Oct 04, 2005
<b><u>AB</u></b>	+	<b><u>EQ 100MG BASE</u></b>	<b><u>N020639 002</u></b>	Sep 26, 1997
<b><u>AB</u></b>	+	<b><u>EQ 200MG BASE</u></b>	<b><u>N020639 003</u></b>	Sep 26, 1997
<b><u>AB</u></b>	! +	<b><u>EQ 300MG BASE</u></b>	<b><u>N020639 005</u></b>	Jul 26, 2000
<b><u>AB</u></b>	+	<b><u>EQ 400MG BASE</u></b>	<b><u>N020639 006</u></b>	Oct 04, 2005

TABLET, EXTENDED RELEASE;ORAL

**QUETIAPINE FUMARATE**

<b><u>AB</u></b>	ACCORD HLTHCARE	<b><u>EQ 50MG BASE</u></b>	<b><u>A206252 001</u></b>	Nov 29, 2017
<b><u>AB</u></b>		<b><u>EQ 150MG BASE</u></b>	<b><u>A090681 001</u></b>	May 09, 2017
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A090681 002</u></b>	May 09, 2017
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A090681 003</u></b>	May 09, 2017
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A090681 004</u></b>	Nov 01, 2016
<b><u>AB</u></b>	ANCHEM PHARMS	<b><u>EQ 150MG BASE</u></b>	<b><u>A090757 001</u></b>	Dec 01, 2017
<b><u>AB</u></b>		<b><u>EQ 200MG BASE</u></b>	<b><u>A090757 002</u></b>	Dec 01, 2017
<b><u>AB</u></b>		<b><u>EQ 300MG BASE</u></b>	<b><u>A090757 003</u></b>	Dec 01, 2017
<b><u>AB</u></b>		<b><u>EQ 400MG BASE</u></b>	<b><u>A090757 004</u></b>	Dec 01, 2017
<b><u>AB</u></b>	AUROBINDO PHARMA LTD	<b><u>EQ 50MG BASE</u></b>	<b><u>A207655 001</u></b>	Nov 29, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-362 (of 436)

QUETIAPINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

QUETIAPINE FUMARATE

<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A207655 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A207655 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A207655 004</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A207655 005</u>	Nov 29, 2017
<u>AB</u>	INTELLIPHARMACEUTIC S	<u>EQ 50MG BASE</u>	<u>A202939 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A202939 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A202939 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A202939 004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A202939 005</u>	May 09, 2017
<u>AB</u>	LUPIN LTD	<u>EQ 50MG BASE</u>	<u>A204203 001</u>	May 17, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A204203 002</u>	May 17, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204203 003</u>	May 17, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204203 004</u>	May 17, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204203 005</u>	May 17, 2017
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 150MG BASE</u>	<u>A204253 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A204253 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A204253 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A204253 004</u>	Nov 29, 2017
<u>AB</u>	NOVAST LABS LTD	<u>EQ 50MG BASE</u>	<u>A208947 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A208947 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A208947 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A208947 004</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A208947 005</u>	Nov 29, 2017
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>A090482 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A090482 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A090482 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A090482 004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A090482 005</u>	May 09, 2017
<u>AB</u>	PHARMADAX INC	<u>EQ 50MG BASE</u>	<u>A206260 001</u>	May 09, 2017
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A206260 002</u>	May 09, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A206260 003</u>	May 09, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A206260 004</u>	May 09, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A206260 005</u>	May 09, 2017
<u>AB</u>	SCIEGEN PHARMS INC	<u>EQ 150MG BASE</u>	<u>A209635 001</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 200MG BASE</u>	<u>A209635 002</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A209635 003</u>	Nov 29, 2017
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>A209635 004</u>	Nov 29, 2017

SEROQUEL XR

<u>AB</u>	+	ASTRAZENECA	<u>EQ 50MG BASE</u>	<u>N022047 001</u>	May 17, 2007
<u>AB</u>	+		<u>EQ 150MG BASE</u>	<u>N022047 005</u>	Aug 11, 2008
<u>AB</u>	+!		<u>EQ 200MG BASE</u>	<u>N022047 002</u>	May 17, 2007
<u>AB</u>	+		<u>EQ 300MG BASE</u>	<u>N022047 003</u>	May 17, 2007
<u>AB</u>	+		<u>EQ 400MG BASE</u>	<u>N022047 004</u>	May 17, 2007

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

ACCUPRIL

<u>AB</u>	+	PFIZER PHARMS	<u>EQ 5MG BASE</u>	<u>N019885 001</u>	Nov 19, 1991
<u>AB</u>	+		<u>EQ 10MG BASE</u>	<u>N019885 002</u>	Nov 19, 1991
<u>AB</u>	+		<u>EQ 20MG BASE</u>	<u>N019885 003</u>	Nov 19, 1991
<u>AB</u>	+!		<u>EQ 40MG BASE</u>	<u>N019885 004</u>	Nov 19, 1991

QUINAPRIL HYDROCHLORIDE

<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A202725 001</u>	Apr 29, 2013
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A202725 002</u>	Apr 29, 2013
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A202725 003</u>	Apr 29, 2013
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A202725 004</u>	Apr 29, 2013
<u>AB</u>		INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A078457 001</u>	Aug 24, 2007
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A078457 002</u>	Aug 24, 2007
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A078457 003</u>	Aug 24, 2007
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A078457 004</u>	Aug 24, 2007
<u>AB</u>		LUPIN	<u>EQ 5MG BASE</u>	<u>A077690 001</u>	Jun 20, 2006
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A077690 002</u>	Jun 20, 2006
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A077690 003</u>	Jun 20, 2006
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A077690 004</u>	Jun 20, 2006
<u>AB</u>		MYLAN	<u>EQ 5MG BASE</u>	<u>A076694 001</u>	Dec 23, 2004
<u>AB</u>			<u>EQ 10MG BASE</u>	<u>A076694 002</u>	Dec 23, 2004
<u>AB</u>			<u>EQ 20MG BASE</u>	<u>A076694 003</u>	Dec 23, 2004
<u>AB</u>			<u>EQ 40MG BASE</u>	<u>A076694 004</u>	Dec 23, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-363 (of 436)

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

QUINAPRIL HYDROCHLORIDE

<u>AB</u>	PRINSTON INC	<u>EQ 5MG BASE</u>	<u>A205823 001</u>	Sep 15, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A205823 002</u>	Sep 15, 2016
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A205823 003</u>	Sep 15, 2016
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A205823 004</u>	Sep 15, 2016
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 5MG BASE</u>	<u>A076607 001</u>	Dec 15, 2004
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A076607 002</u>	Dec 15, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A076607 003</u>	Dec 15, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A076607 004</u>	Dec 15, 2004
<u>AB</u>	TEVA	<u>EQ 5MG BASE</u>	<u>A075504 001</u>	Aug 24, 2007
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075504 002</u>	Aug 24, 2007
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>A075504 003</u>	Aug 24, 2007
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>A075504 004</u>	Aug 24, 2007

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE GLUCONATE

BX	!	SUN PHARM INDUSTRIES	324MG	A089338 001	Feb 11, 1987
----	---	----------------------	-------	-------------	--------------

QUINIDINE SULFATE

TABLET;ORAL

QUINIDINE SULFATE

<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A088072 002</u>	
<u>AB</u>		<u>300MG</u>	<u>A088072 001</u>	Sep 26, 1983
<u>AB</u>	SUN PHARM INDUSTRIES	<u>200MG</u>	<u>A081030 001</u>	Apr 14, 1989
<u>AB</u>		<u>300MG</u>	<u>A081031 001</u>	Apr 14, 1989
<u>AB</u>	! WATSON LABS	<u>200MG</u>	<u>A083288 001</u>	
<u>AB</u>	!	<u>300MG</u>	<u>A085583 001</u>	

TABLET, EXTENDED RELEASE;ORAL

QUINIDINE SULFATE

!	G AND W LABS INC	300MG
---	------------------	-------

A040045 001 Jun 30, 1994

QUININE SULFATE

CAPSULE;ORAL

QUALAQUIN

<u>AB</u>	+! SUN PHARM INDUSTRIES	<u>324MG</u>	<u>N021799 001</u>	Aug 12, 2005
-----------	-------------------------	--------------	--------------------	--------------

QUININE SULFATE

<u>AB</u>	AMNEAL PHARMS	<u>324MG</u>	<u>A203729 001</u>	Jul 15, 2015
<u>AB</u>	LUPIN LTD	<u>324MG</u>	<u>A203112 001</u>	Apr 24, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>324MG</u>	<u>A202581 001</u>	Dec 14, 2012
<u>AB</u>	RICONPHARMA LLC	<u>324MG</u>	<u>A204372 001</u>	Jul 22, 2015
<u>AB</u>	TEVA PHARMS	<u>324MG</u>	<u>A091661 001</u>	Sep 28, 2012

RABEPRAZOLE SODIUM

CAPSULE, DELAYED RELEASE;ORAL

ACIPHEX SPRINKLE

+	AVADEL PHARMS	5MG
! !		10MG

N204736 001 Mar 26, 2013

N204736 002 Mar 26, 2013

TABLET, DELAYED RELEASE;ORAL

ACIPHEX

<u>AB</u>	+! EISAI INC	<u>20MG</u>	<u>N020973 002</u>	Aug 19, 1999
-----------	--------------	-------------	--------------------	--------------

RABEPRAZOLE SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>20MG</u>	<u>A204179 001</u>	Jul 31, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A205761 001</u>	Feb 17, 2017
<u>AB</u>	BRECKENRIDGE PHARM	<u>20MG</u>	<u>A204237 001</u>	Nov 18, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>20MG</u>	<u>A076824 001</u>	Nov 08, 2013
<u>AB</u>	KREMERS URBAN PHARMS	<u>20MG</u>	<u>A090678 001</u>	Nov 08, 2013
<u>AB</u>	LUPIN LTD	<u>20MG</u>	<u>A078964 001</u>	Nov 08, 2013
<u>AB</u>	MYLAN PHARMS INC	<u>20MG</u>	<u>A076885 001</u>	Nov 08, 2013
<u>AB</u>	TEVA PHARMS USA	<u>20MG</u>	<u>A076822 001</u>	Nov 08, 2013
<u>AB</u>	TORRENT PHARMS LTD	<u>20MG</u>	<u>A202376 001</u>	Nov 08, 2013

RADIUM RA-223 DICHLORIDE

SOLUTION;INTRAVENOUS

XOFIGO

! !	BAYER HLTHCARE	162mCi/6ML (27mCi/ML)
-----	----------------	-----------------------

N203971 001 May 15, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-364 (of 436)

RALOXIFENE HYDROCHLORIDE

TABLET;ORAL

EVISTA

<u>AB</u>	+!	LILLY	<u>60MG</u>	<u>N020815 001</u>	Dec 09, 1997
<u>RALOXIFENE HYDROCHLORIDE</u>					
<u>AB</u>		AMNEAL PHARMS	<u>60MG</u>	<u>A208206 001</u>	Apr 08, 2016
<u>AB</u>		AUROBINDO PHARMA LTD	<u>60MG</u>	<u>A204310 001</u>	Aug 28, 2015
<u>AB</u>		GLENMARK PHARMS LTD	<u>60MG</u>	<u>A204491 001</u>	Mar 22, 2016
<u>AB</u>		INVAGEN PHARMS	<u>60MG</u>	<u>A090842 001</u>	Sep 24, 2014
<u>AB</u>		SCIEGEN PHARMS INC	<u>60MG</u>	<u>A206384 001</u>	Oct 12, 2016
<u>AB</u>		TEVA PHARMS USA	<u>60MG</u>	<u>A078193 001</u>	Mar 04, 2014
<u>AB</u>		WATSON LABS INC	<u>60MG</u>	<u>A200825 001</u>	Jan 21, 2015

RALTEGRAVIR POTASSIUM

POWDER;ORAL

ISENTRESS

+! MERCK SHARP DOHME EQ 100MG BASE/PACKET N205786 001 Dec 20, 2013

TABLET;ORAL

ISENTRESS

+! MERCK SHARP DOHME EQ 400MG BASE N022145 001 Oct 12, 2007

ISENTRESS HD

+ MERCK SHARP DOHME EQ 600MG BASE N022145 002 May 26, 2017

TABLET, CHEWABLE;ORAL

ISENTRESS

+ MERCK SHARP DOHME EQ 25MG BASE N203045 001 Dec 21, 2011

+! MERCK SHARP DOHME EQ 100MG BASE N203045 002 Dec 21, 2011

RAMELTEON

TABLET;ORAL

RAMELTEON

<u>AB</u>		ACTAVIS LABS FL INC	<u>8MG</u>	<u>A091610 001</u>	Aug 19, 2015
<u>AB</u>		DR REDDYS LABS INTL	<u>8MG</u>	<u>A091693 001</u>	Jul 26, 2013

ROZEREM

<u>AB</u>	+!	TAKEDA PHARMS USA	<u>8MG</u>	<u>N021782 001</u>	Jul 22, 2005
-----------	----	-------------------	------------	--------------------	--------------

RAMIPRIL

CAPSULE;ORAL

ALTACE

<u>AB</u>	+	KING PHARMS LLC	<u>1.25MG</u>	<u>N019901 001</u>	Jan 28, 1991
<u>AB</u>	+		<u>2.5MG</u>	<u>N019901 002</u>	Jan 28, 1991
<u>AB</u>	+		<u>5MG</u>	<u>N019901 003</u>	Jan 28, 1991
<u>AB</u>	+!		<u>10MG</u>	<u>N019901 004</u>	Jan 28, 1991

RAMIPRIL

<u>AB</u>		ACCORD HLTHCARE	<u>1.25MG</u>	<u>A202392 001</u>	Apr 15, 2014
<u>AB</u>			<u>2.5MG</u>	<u>A202392 002</u>	Apr 15, 2014
<u>AB</u>			<u>5MG</u>	<u>A202392 003</u>	Apr 15, 2014
<u>AB</u>			<u>10MG</u>	<u>A202392 004</u>	Apr 15, 2014
<u>AB</u>		APOTEX	<u>1.25MG</u>	<u>A079116 001</u>	Jun 20, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A079116 002</u>	Jun 20, 2008
<u>AB</u>			<u>5MG</u>	<u>A079116 003</u>	Jun 20, 2008
<u>AB</u>			<u>10MG</u>	<u>A079116 004</u>	Jun 20, 2008
<u>AB</u>		AUROBINDO PHARMA LTD	<u>1.25MG</u>	<u>A091604 001</u>	Jun 08, 2011
<u>AB</u>			<u>2.5MG</u>	<u>A091604 002</u>	Jun 08, 2011
<u>AB</u>			<u>5MG</u>	<u>A091604 003</u>	Jun 08, 2011
<u>AB</u>			<u>10MG</u>	<u>A091604 004</u>	Jun 08, 2011
<u>AB</u>		DR REDDYS LABS LTD	<u>1.25MG</u>	<u>A078191 001</u>	Jun 18, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A078191 002</u>	Jun 18, 2008
<u>AB</u>			<u>5MG</u>	<u>A078191 003</u>	Jun 18, 2008
<u>AB</u>			<u>10MG</u>	<u>A078191 004</u>	Jun 18, 2008
<u>AB</u>		INVAGEN PHARMS	<u>1.25MG</u>	<u>A078745 001</u>	Jun 18, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A078745 002</u>	Jun 18, 2008
<u>AB</u>			<u>5MG</u>	<u>A078745 003</u>	Jun 18, 2008
<u>AB</u>			<u>10MG</u>	<u>A078745 004</u>	Jun 18, 2008
<u>AB</u>		LUPIN	<u>1.25MG</u>	<u>A077626 001</u>	Jun 09, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A077626 002</u>	Jun 09, 2008
<u>AB</u>			<u>5MG</u>	<u>A077626 003</u>	Jun 09, 2008
<u>AB</u>			<u>10MG</u>	<u>A077626 004</u>	Jun 09, 2008
<u>AB</u>		TEVA PHARMS	<u>1.25MG</u>	<u>A077470 001</u>	Jun 18, 2008
<u>AB</u>			<u>2.5MG</u>	<u>A077470 002</u>	Jun 18, 2008
<u>AB</u>			<u>5MG</u>	<u>A077470 003</u>	Jun 18, 2008
<u>AB</u>			<u>10MG</u>	<u>A077470 004</u>	Jun 18, 2008
<u>AB</u>		WATSON LABS	<u>1.25MG</u>	<u>A076549 001</u>	Oct 24, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-365 (of 436)

RAMIPRIL

CAPSULE;ORAL

RAMIPRIL

<u>AB</u>		<u>2.5MG</u>	<u>A076549 002</u>	Oct 24, 2005
<u>AB</u>		<u>5MG</u>	<u>A076549 003</u>	Oct 24, 2005
<u>AB</u>		<u>10MG</u>	<u>A076549 004</u>	Oct 24, 2005
<u>AB</u>	WEST-WARD PHARMS INT	<u>1.25MG</u>	<u>A077900 001</u>	Jun 18, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A077900 002</u>	Jun 18, 2008
<u>AB</u>		<u>5MG</u>	<u>A077900 003</u>	Jun 18, 2008
<u>AB</u>		<u>10MG</u>	<u>A077900 004</u>	Jun 18, 2008
<u>AB</u>	ZYDUS PHARMS USA	<u>1.25MG</u>	<u>A078832 001</u>	Sep 02, 2008
<u>AB</u>		<u>2.5MG</u>	<u>A078832 002</u>	Sep 02, 2008
<u>AB</u>		<u>5MG</u>	<u>A078832 003</u>	Sep 02, 2008
<u>AB</u>		<u>10MG</u>	<u>A078832 004</u>	Sep 02, 2008

TABLET;ORAL

RAMIPRIL

APOTEX INC	1.25MG	A091069 001	Dec 02, 2015
	2.5MG	A091069 002	Dec 02, 2015
	5MG	A091069 003	Dec 02, 2015
!	10MG	A091069 004	Dec 02, 2015

RANITIDINE HYDROCHLORIDE

CAPSULE;ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>A075742 001</u>	Nov 29, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075742 002</u>	Nov 29, 2000
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>A074655 001</u>	Oct 22, 1997
<u>AB</u>	!	<u>EQ 300MG BASE</u>	<u>A074655 002</u>	Oct 22, 1997

INJECTABLE;INJECTION

RANITIDINE HYDROCHLORIDE

<u>AP</u>	MYLAN LABS LTD	<u>EQ 25MG BASE/ML</u>	<u>A079076 001</u>	Jun 09, 2016
<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 25MG BASE/ML</u>	<u>A074777 001</u>	Mar 02, 2005
<u>AP</u>		<u>EQ 25MG BASE/ML</u>	<u>A077458 001</u>	Feb 16, 2006
<u>AP</u>	ZYDUS PHARMS USA INC	<u>EQ 25MG BASE/ML</u>	<u>A091534 001</u>	Feb 22, 2013
<b>ZANTAC</b>				

AP +! TELIGENT

SYRUP;ORAL

EQ 25MG BASE/ML

N019090 001 Oct 19, 1984

RANITIDINE HYDROCHLORIDE

<u>AA</u>	ACTAVIS MID ATLANTIC	<u>EQ 15MG BASE/ML</u>	<u>A076124 001</u>	Feb 21, 2007
<u>AA</u>	AMNEAL PHARMS	<u>EQ 15MG BASE/ML</u>	<u>A078312 001</u>	Sep 02, 2008
<u>AA</u>	AUROBINDO PHARMA LTD	<u>EQ 15MG BASE/ML</u>	<u>A090623 001</u>	Jul 28, 2010
<u>AA</u>	BIO PHARM INC	<u>EQ 15MG BASE/ML</u>	<u>A090102 001</u>	May 26, 2009
<u>AA</u>	BRECKENRIDGE PHARM	<u>EQ 15MG BASE/ML</u>	<u>A078684 001</u>	Aug 27, 2009
<u>AA</u>	HI TECH PHARMA	<u>EQ 15MG BASE/ML</u>	<u>A091078 001</u>	Mar 22, 2011
<u>AA</u>	NOSTRUM LABS INC	<u>EQ 15MG BASE/ML</u>	<u>A091091 001</u>	Sep 20, 2011
<u>AA</u>	PHARM ASSOC	<u>EQ 15MG BASE/ML</u>	<u>A077405 001</u>	Sep 21, 2007
<u>AA</u>	SILARX	<u>EQ 15MG BASE/ML</u>	<u>A091288 001</u>	Dec 09, 2010
<u>AA</u>	TARO	<u>EQ 15MG BASE/ML</u>	<u>A077476 001</u>	Jun 13, 2011
<u>AA</u>	TOLMAR	<u>EQ 15MG BASE/ML</u>	<u>A090054 001</u>	Nov 15, 2010
<u>AA</u>	VINTAGE PHARMS	<u>EQ 15MG BASE/ML</u>	<u>A078890 001</u>	Jul 01, 2010
<b>ZANTAC</b>				

AA +! GLAXO GRP LTD

EQ 15MG BASE/ML

N019675 001 Dec 30, 1988

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	ACIC PHARMS	<u>EQ 150MG BASE</u>	<u>A203694 001</u>	Nov 30, 2017
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A203694 002</u>	Nov 30, 2017
<u>AB</u>	AMNEAL PHARMS NY	<u>EQ 150MG BASE</u>	<u>A077824 001</u>	Oct 13, 2006
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A077824 002</u>	Oct 13, 2006
<u>AB</u>	APOTEX	<u>EQ 150MG BASE</u>	<u>A074680 001</u>	Sep 12, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A074680 002</u>	Sep 12, 1997
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 150MG BASE</u>	<u>A076705 001</u>	Jul 27, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A076705 002</u>	Jul 27, 2005
<u>AB</u>	GLENMARK PHARMS INC	<u>EQ 150MG BASE</u>	<u>A078542 001</u>	Nov 19, 2008
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A078542 002</u>	Nov 19, 2008
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 150MG BASE</u>	<u>A075165 001</u>	Sep 30, 1998
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075165 002</u>	Sep 30, 1998
<u>AB</u>	PAR PHARM	<u>EQ 150MG BASE</u>	<u>A075180 001</u>	Jan 28, 1999
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>A075180 002</u>	Jan 28, 1999

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-366 (of 436)

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

**RANITIDINE HYDROCHLORIDE**

<b>AB</b>	SANDOZ	<u>EQ 150MG BASE</u>	<b>A074467 001</b>	Aug 29, 1997
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A074467 002</b>	Aug 29, 1997
<b>AB</b>	STRIDES PHARMA	<u>EQ 150MG BASE</u>	<b>A205512 001</b>	Aug 22, 2016
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A205512 002</b>	Aug 22, 2016
<b>AB</b>	TEVA	<u>EQ 150MG BASE</u>	<b>A074488 001</b>	Jul 31, 1997
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A074488 002</b>	Jul 31, 1997
<b>AB</b>	WOCKHARDT LTD	<u>EQ 150MG BASE</u>	<b>A075208 001</b>	Dec 17, 1998
<b>AB</b>		<u>EQ 300MG BASE</u>	<b>A075208 002</b>	Dec 17, 1998
	<b>ZANTAC 150</b>			
<b>AB</b>	+ GLAXO GRP LTD	<u>EQ 150MG BASE</u>	<b>N018703 001</b>	Jun 09, 1983
	<b>ZANTAC 300</b>			
<b>AB</b>	+! GLAXO GRP LTD	<u>EQ 300MG BASE</u>	<b>N018703 002</b>	Dec 09, 1985

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

+ GILEAD	500MG	N021526 002	Jan 27, 2006
+!	1GM	N021526 001	Feb 12, 2007

RASAGILINE MESYLATE

TABLET; ORAL

**AZILECT**

<b>AB</b>	+ TEVA	<u>EQ 0.5MG BASE</u>	<b>N021641 001</b>	May 16, 2006
<b>AB</b>	+!	<u>EQ 1MG BASE</u>	<b>N021641 002</b>	May 16, 2006

**RASAGILINE MESYLATE**

<b>AB</b>	ALKEM LABS LTD	<u>EQ 0.5MG BASE</u>	<b>A201889 001</b>	Oct 30, 2017
<b>AB</b>		<u>EQ 1MG BASE</u>	<b>A201889 002</b>	Oct 30, 2017
<b>AB</b>	APOTEX INC	<u>EQ 0.5MG BASE</u>	<b>A201950 001</b>	Sep 12, 2013
<b>AB</b>		<u>EQ 1MG BASE</u>	<b>A201950 002</b>	Sep 12, 2013
<b>AB</b>	MYLAN PHARMS INC	<u>EQ 0.5MG BASE</u>	<b>A201971 001</b>	May 15, 2017
<b>AB</b>		<u>EQ 1MG BASE</u>	<b>A201971 002</b>	May 15, 2017
<b>AB</b>	ORCHID HLTHCARE	<u>EQ 0.5MG BASE</u>	<b>A201970 001</b>	Mar 15, 2016
<b>AB</b>		<u>EQ 1MG BASE</u>	<b>A201970 002</b>	Mar 15, 2016
<b>AB</b>	WATSON LABS INC	<u>EQ 0.5MG BASE</u>	<b>A201823 001</b>	Jul 01, 2013
<b>AB</b>		<u>EQ 1MG BASE</u>	<b>A201823 002</b>	Jul 01, 2013

REGADERONOSON

SOLUTION; INTRAVENOUS

LEXISCAN

+! ASTELLAS	0.4MG/5ML (0.08MG/ML)	N022161 001	Apr 10, 2008
-------------	-----------------------	-------------	--------------

REGORAFENIB

TABLET; ORAL

STIVARGA

+! BAYER HLTHCARE	40MG	N203085 001	Sep 27, 2012
-------------------	------	-------------	--------------

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ULTIVA

+ MYLAN INSTITUTIONAL	<u>EQ 1MG BASE/VIAL</u>	N020630 001	Jul 12, 1996
+	<u>EQ 2MG BASE/VIAL</u>	N020630 002	Jul 12, 1996
+!	<u>EQ 5MG BASE/VIAL</u>	N020630 003	Jul 12, 1996

REPAGLINIDE

TABLET; ORAL

**PRANDIN**

<b>AB</b>	+ GEMINI LABS LLC	<u>0.5MG</u>	<b>N020741 001</b>	Dec 22, 1997
<b>AB</b>	+	<u>1MG</u>	<b>N020741 002</b>	Dec 22, 1997
<b>AB</b>	+!	<u>2MG</u>	<b>N020741 003</b>	Dec 22, 1997

**REPAGLINIDE**

<b>AB</b>	ACTAVIS TOTOWA	<u>0.5MG</u>	<b>A090008 001</b>	Jan 22, 2014
<b>AB</b>		<u>1MG</u>	<b>A090008 002</b>	Jan 22, 2014
<b>AB</b>		<u>2MG</u>	<b>A090008 003</b>	Jan 22, 2014
<b>AB</b>	AUROBINDO PHARMA LTD	<u>0.5MG</u>	<b>A203820 001</b>	Jan 22, 2014
<b>AB</b>		<u>1MG</u>	<b>A203820 002</b>	Jan 22, 2014
<b>AB</b>		<u>2MG</u>	<b>A203820 003</b>	Jan 22, 2014
<b>AB</b>	BOSCOGEN	<u>0.5MG</u>	<b>A091517 001</b>	Apr 24, 2015
<b>AB</b>		<u>1MG</u>	<b>A091517 002</b>	Apr 24, 2015
<b>AB</b>		<u>2MG</u>	<b>A091517 003</b>	Apr 24, 2015
<b>AB</b>	MYLAN PHARMS INC	<u>0.5MG</u>	<b>A090252 001</b>	Aug 23, 2013
<b>AB</b>		<u>1MG</u>	<b>A090252 002</b>	Jan 22, 2014
<b>AB</b>		<u>2MG</u>	<b>A090252 003</b>	Jan 22, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-367 (of 436)

REPAGLINIDE

TABLET;ORAL

REPAGLINIDE

<u>AB</u>	PADDOCK LLC	<u>0.5MG</u>	<u>A201189 001</u>	Jul 17, 2013
<u>AB</u>		<u>1MG</u>	<u>A201189 002</u>	Jan 22, 2014
<u>AB</u>		<u>2MG</u>	<u>A201189 003</u>	Jan 22, 2014
<u>AB</u>	SANDOZ INC	<u>0.5MG</u>	<u>A078555 001</u>	Nov 22, 2013
<u>AB</u>		<u>1MG</u>	<u>A078555 002</u>	Jan 22, 2014
<u>AB</u>		<u>2MG</u>	<u>A078555 003</u>	Jan 22, 2014
<u>AB</u>	SUN PHARM INDNS INC	<u>1MG</u>	<u>A077571 002</u>	Jul 11, 2013
<u>AB</u>		<u>2MG</u>	<u>A077571 003</u>	Jul 11, 2013

RETAPAMULIN

OINTMENT;TOPICAL

ALTABAX

+!	AQUA PHARMS LLC	1%
		N022055 001 Apr 12, 2007

RIBAVIRIN

CAPSULE;ORAL

REBETOL

<u>AB</u>	+! MERCK SHARP DOHME	<u>200MG</u>	<u>N020903 002</u>	Jul 25, 2001
<u>AB</u>		<u>RIBOSPHERE</u>		
<u>AB</u>	KADMON PHARMS LLC	<u>200MG</u>	<u>A076203 001</u>	Apr 06, 2004
<u>AB</u>		<u>RIBAVARIN</u>		
<u>AB</u>	AUROBINDO PHARMA	<u>200MG</u>	<u>A079117 001</u>	Sep 17, 2009
<u>AB</u>		<u>RIBAVIRIN</u>		
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A076192 001</u>	Apr 06, 2004
<u>AB</u>	TEVA	<u>200MG</u>	<u>A076277 001</u>	Oct 04, 2004
<u>AB</u>	ZYDUS PHARMS USA	<u>200MG</u>	<u>A077224 001</u>	Oct 28, 2005

FOR SOLUTION;INHALATION

RIBAVIRIN

<u>AN</u>	NAVINTA LLC	<u>6GM/VIAL</u>	<u>A207366 001</u>	Oct 06, 2016
<u>AN</u>		<u>VIRAZOLE</u>		

<u>AN</u>	+! VALEANT PHARM INTL	<u>6GM/VIAL</u>	<u>N018859 001</u>	Dec 31, 1985

SOLUTION;ORAL

REBETOL

+!	SCHERING	40MG/ML
		N021546 001 Jul 29, 2003

TABLET;ORAL

COPEGUS

<u>AB</u>	ROCHE	<u>200MG</u>	<u>N021511 001</u>	Dec 03, 2002
		<u>RIBAVIRIN</u>		
<u>AB</u>	AUROBINDO PHARMA	<u>200MG</u>	<u>A079111 001</u>	Sep 17, 2009
<u>AB</u>	KADMON PHARMS LLC	<u>200MG</u>	<u>A077456 001</u>	Dec 05, 2005
<u>AB</u>		<u>400MG</u>	<u>A077456 002</u>	Dec 05, 2005
<u>AB</u>	!	<u>600MG</u>	<u>A077456 003</u>	Dec 05, 2005
<u>AB</u>	SANDOZ	<u>200MG</u>	<u>A077743 001</u>	Oct 03, 2006
<u>AB</u>	SANDOZ INC	<u>200MG</u>	<u>A202546 001</u>	Aug 12, 2014
<u>AB</u>		<u>400MG</u>	<u>A202546 002</u>	Aug 12, 2014
<u>AB</u>		<u>500MG</u>	<u>A202546 003</u>	Aug 12, 2014
<u>AB</u>		<u>600MG</u>	<u>A202546 004</u>	Aug 12, 2014
<u>AB</u>	TEVA	<u>200MG</u>	<u>A077053 001</u>	Dec 05, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>200MG</u>	<u>A077094 001</u>	Dec 05, 2005
<u>AB</u>		<u>400MG</u>	<u>A077094 002</u>	Mar 16, 2007
<u>AB</u>		<u>500MG</u>	<u>A077094 004</u>	Apr 18, 2008
<u>AB</u>		<u>600MG</u>	<u>A077094 003</u>	Mar 16, 2007

RIBOCICLIB SUCCINATE

TABLET;ORAL

KISQALI

+!	NOVARTIS PHARMS CORP	EQ 200MG BASE
		N209092 001 Mar 13, 2017

RIBOFLAVIN 5'-PHOSPHATE SODIUM

SOLUTION/DROPS;OPHTHALMIC

PHOTREXA

+!	AVEDRO INC	0.146%	N203324 001	Apr 15, 2016
		PHOTREXA VISCOSU IN DEXTRAN 20%		
+!	AVEDRO INC	0.146%	N203324 002	Apr 15, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-368 (of 436)

RIFABUTIN

CAPSULE;ORAL

MYCOBUTIN

AB +! PHARMACIA AND UPJOHN 150MG N050689 001 Dec 23, 1992

RIFABUTIN

AB LUPIN LTD 150MG A090033 001 Feb 24, 2014

RIFAMPIN

CAPSULE;ORAL

RIFADIN

AB SANOFI AVENTIS US 150MG A062303 001  
AB +! 300MG N050420 001

RIFAMPIN

AB AKORN 150MG A065028 001 Mar 14, 2001  
AB 300MG A065028 002 Mar 14, 2001  
AB LANNETT 150MG A065390 001 Mar 28, 2008  
AB 300MG A065390 002 Mar 28, 2008  
AB LUPIN PHARMS 150MG A090034 001 Aug 21, 2013  
AB 300MG A090034 002 Aug 21, 2013  
AB SANDOZ 150MG A064150 002 Jan 02, 1998  
AB 300MG A064150 001 May 28, 1997

RIMACTANE

AB OXFORD PHARMS 300MG N050429 001  
 INJECTABLE;INJECTION

RIFADIN

AP +! SANOFI AVENTIS US 600MG/VIAL N050627 001 May 25, 1989  
RIFAMPIN

AP AKORN 600MG/VIAL A065502 001 Sep 21, 2010  
AP EMCURE PHARMS LTD 600MG/VIAL A204101 001 Aug 18, 2014  
AP FRESENIUS KABI USA 600MG/VIAL A091181 001 Aug 21, 2014  
AP HIKMA PHARMS 600MG/VIAL A205039 001 Mar 03, 2016  
AP MYLAN LABS LTD 600MG/VIAL A065421 001 May 22, 2008  
AP WATSON PHARMS TEVA 600MG/VIAL A206736 001 Jan 19, 2016  
AP WEST-WARD PHARMS INT 600MG/VIAL A064217 001 Oct 29, 1999

RIFAPENTINE

TABLET;ORAL

PRIFTIN

+! SANOFI AVENTIS US 150MG N021024 001 Jun 22, 1998

RIFAXIMIN

TABLET;ORAL

XIFAXAN

+! SALIX PHARMS 200MG N021361 001 May 25, 2004  
 +! 550MG N022554 001 Mar 24, 2010

RILPIVIRINE HYDROCHLORIDE

TABLET;ORAL

EDURANT

+! JANSSEN PRODS EQ 25MG BASE N202022 001 May 20, 2011

RILUZOLE

TABLET;ORAL

RILUTEK

AB +! COVIS PHARMA BV 50MG N020599 001 Dec 12, 1995  
RILUZOLE  
AB ALKEM LABS LTD 50MG A204048 001 Mar 30, 2016  
AB APOTEX CORP 50MG A091300 001 Jun 18, 2013  
AB GLENMARK PHARMS LTD 50MG A091394 001 Jun 18, 2013  
AB IMPAX LABS 50MG A076173 001 Jan 29, 2003  
AB MYLAN PHARMS INC 50MG A203042 001 Jul 01, 2013  
AB SUN PHARM INDs LTD 50MG A091417 001 Jun 18, 2013

RIMANTADINE HYDROCHLORIDE

TABLET;ORAL

FLUMADINE

AB +! SUN PHARM INDs INC 100MG N019649 001 Sep 17, 1993

RIMANTADINE HYDROCHLORIDE

AB IMPAX LABS 100MG A076132 001 Aug 30, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-369 (of 436)

RIOCIGUAT

TABLET;ORAL

ADEMPAS

+ BAYER HLTHCARE	0.5MG	N204819 001	Oct 08, 2013
+	1MG	N204819 002	Oct 08, 2013
+	1.5MG	N204819 003	Oct 08, 2013
+	2MG	N204819 004	Oct 08, 2013
+!	2.5MG	N204819 005	Oct 08, 2013

RISEDRONATE SODIUM

TABLET;ORAL

ACTONEL

<u>AB</u> + APIL	<u>5MG</u>	<u>N020835 002</u>	Apr 14, 2000
<u>AB</u> +	<u>30MG</u>	<u>N020835 001</u>	Mar 27, 1998
<u>AB</u> +!	<u>35MG</u>	<u>N020835 003</u>	May 25, 2002
<u>AB</u> +!	<u>150MG</u>	<u>N020835 005</u>	Apr 22, 2008

RISEDRONATE SODIUM

<u>AB</u> APOTEX INC	<u>35MG</u>	<u>A090877 001</u>	Nov 30, 2015
<u>AB</u>	<u>75MG</u>	<u>A090877 002</u>	Jun 10, 2014
<u>AB</u>	<u>150MG</u>	<u>A090877 003</u>	Jun 10, 2014
<u>AB</u> AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A200296 001</u>	Nov 30, 2015
<u>AB</u>	<u>30MG</u>	<u>A200296 002</u>	Nov 30, 2015
<u>AB</u>	<u>35MG</u>	<u>A200296 003</u>	Nov 30, 2015
<u>AB</u>	<u>150MG</u>	<u>A206768 001</u>	Oct 21, 2016
<u>AB</u> MACLEODS PHARMS LTD	<u>5MG</u>	<u>A203533 001</u>	Dec 09, 2015
<u>AB</u>	<u>30MG</u>	<u>A203533 002</u>	Dec 09, 2015
<u>AB</u>	<u>35MG</u>	<u>A203533 003</u>	Nov 29, 2016
<u>AB</u> MYLAN PHARMS INC	<u>5MG</u>	<u>A200477 001</u>	Nov 30, 2015
<u>AB</u>	<u>30MG</u>	<u>A200477 002</u>	Nov 30, 2015
<u>AB</u>	<u>35MG</u>	<u>A200477 003</u>	Nov 30, 2015
<u>AB</u>	<u>75MG</u>	<u>A200477 004</u>	Jun 10, 2014
<u>AB</u>	<u>150MG</u>	<u>A200477 005</u>	Jun 10, 2014
<u>AB</u> SUN PHARMA GLOBAL	<u>5MG</u>	<u>A090886 001</u>	Nov 30, 2015
<u>AB</u>	<u>30MG</u>	<u>A090886 002</u>	Nov 30, 2015
<u>AB</u>	<u>35MG</u>	<u>A090886 003</u>	Nov 30, 2015
<u>AB</u>	<u>75MG</u>	<u>A090886 004</u>	Jun 10, 2014
<u>AB</u>	<u>150MG</u>	<u>A090886 005</u>	Jun 10, 2014
<u>AB</u> TEVA PHARMS USA	<u>5MG</u>	<u>A077132 001</u>	Oct 05, 2007
<u>AB</u>	<u>30MG</u>	<u>A077132 002</u>	Oct 05, 2007
<u>AB</u>	<u>35MG</u>	<u>A077132 003</u>	Oct 05, 2007
<u>AB</u>	<u>150MG</u>	<u>A079215 001</u>	Jun 13, 2014

TABLET, DELAYED RELEASE;ORAL

ATELVIA

<u>AB</u> +! APIL	<u>35MG</u>	<u>N022560 001</u>	Oct 08, 2010
<u>AB</u> TEVA PHARMS USA	<u>35MG</u>	<u>A203217 001</u>	May 18, 2015

RISPERIDONE

INJECTABLE;INTRAMUSCULAR

RISPERDAL CONSTA

+ JANSSEN PHARMS	12.5MG/VIAL	N021346 004	Apr 12, 2007
+!	25MG/VIAL	N021346 001	Oct 29, 2003
+	37.5MG/VIAL	N021346 002	Oct 29, 2003
+	50MG/VIAL	N021346 003	Oct 29, 2003

SOLUTION;ORAL

RISPERDAL

<u>AA</u> +! JANSSEN PHARMS	<u>1MG/ML</u>	<u>N020588 001</u>	Jun 10, 1996
<u>AA</u> RISPERIDONE	<u>1MG/ML</u>	<u>A091384 001</u>	May 25, 2011
<u>AA</u> AMNEAL PHARMS	<u>1MG/ML</u>	<u>A076440 001</u>	Jan 30, 2009
<u>AA</u> ANI PHARMS INC	<u>1MG/ML</u>	<u>A077719 001</u>	Jul 29, 2009
<u>AA</u> APOTEX INC	<u>1MG/ML</u>	<u>A078909 001</u>	Jul 29, 2009
<u>AA</u> BIO PHARM INC	<u>1MG/ML</u>	<u>A078452 001</u>	Sep 04, 2009
<u>AA</u> LIFESTAR PHARMA	<u>1MG/ML</u>	<u>A076797 001</u>	Jun 28, 2010
<u>AA</u> PRECISION DOSE	<u>1MG/ML</u>	<u>A090347 001</u>	Feb 07, 2011
<u>AA</u> TARO	<u>1MG/ML</u>	<u>A079059 001</u>	Dec 12, 2012
<u>AA</u> TRIS PHARMA INC	<u>1MG/ML</u>	<u>A079158 001</u>	Dec 03, 2010
<u>AA</u> VINTAGE	<u>1MG/ML</u>	<u>A076904 001</u>	Jul 29, 2009
<u>AA</u> WEST-WARD PHARMS INT	<u>1MG/ML</u>		

TABLET;ORAL

RISPERDAL

<u>AB</u> + JANSSEN PHARMS	<u>0.25MG</u>	<u>N020272 008</u>	May 10, 1999
<u>AB</u> +	<u>0.5MG</u>	<u>N020272 007</u>	Jan 27, 1999

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-370 (of 436)

RISPERIDONE

TABLET;ORAL

RISPERDAL

<u>AB</u>	+!	<u>1MG</u>	<u>N020272</u> <u>001</u>	Dec 29, 1993
<u>AB</u>	+	<u>2MG</u>	<u>N020272</u> <u>002</u>	Dec 29, 1993
<u>AB</u>	+	<u>3MG</u>	<u>N020272</u> <u>003</u>	Dec 29, 1993
<u>AB</u>	+	<u>4MG</u>	<u>N020272</u> <u>004</u>	Dec 29, 1993
<b>RISPERIDONE</b>				
<u>AB</u>	AJANTA PHARMA LTD	<u>0 .25MG</u>	<u>A201003</u> <u>001</u>	Aug 24, 2011
<u>AB</u>		<u>0 .5MG</u>	<u>A201003</u> <u>002</u>	Aug 24, 2011
<u>AB</u>		<u>1MG</u>	<u>A201003</u> <u>003</u>	Aug 24, 2011
<u>AB</u>		<u>2MG</u>	<u>A201003</u> <u>004</u>	Aug 24, 2011
<u>AB</u>		<u>3MG</u>	<u>A201003</u> <u>005</u>	Aug 24, 2011
<u>AB</u>		<u>4MG</u>	<u>A201003</u> <u>006</u>	Aug 24, 2011
<u>AB</u>	APOTEX INC	<u>0 .25MG</u>	<u>A077953</u> <u>001</u>	Sep 15, 2008
<u>AB</u>		<u>0 .5MG</u>	<u>A077953</u> <u>002</u>	Sep 15, 2008
<u>AB</u>		<u>1MG</u>	<u>A077953</u> <u>003</u>	Sep 15, 2008
<u>AB</u>		<u>2MG</u>	<u>A077953</u> <u>004</u>	Sep 15, 2008
<u>AB</u>		<u>3MG</u>	<u>A077953</u> <u>005</u>	Sep 15, 2008
<u>AB</u>		<u>4MG</u>	<u>A077953</u> <u>006</u>	Sep 15, 2008
<u>AB</u>	AUROBINDO PHARMA	<u>0 .25MG</u>	<u>A078269</u> <u>001</u>	Oct 08, 2008
<u>AB</u>		<u>0 .5MG</u>	<u>A078269</u> <u>002</u>	Oct 08, 2008
<u>AB</u>		<u>1MG</u>	<u>A078269</u> <u>003</u>	Oct 08, 2008
<u>AB</u>		<u>2MG</u>	<u>A078269</u> <u>004</u>	Oct 08, 2008
<u>AB</u>		<u>3MG</u>	<u>A078269</u> <u>005</u>	Oct 08, 2008
<u>AB</u>		<u>4MG</u>	<u>A078269</u> <u>006</u>	Oct 08, 2008
<u>AB</u>	CIPLA	<u>0 .25MG</u>	<u>A077543</u> <u>001</u>	May 18, 2011
<u>AB</u>		<u>0 .5MG</u>	<u>A077543</u> <u>002</u>	May 18, 2011
<u>AB</u>		<u>1MG</u>	<u>A077543</u> <u>003</u>	May 18, 2011
<u>AB</u>		<u>2MG</u>	<u>A077543</u> <u>004</u>	May 18, 2011
<u>AB</u>		<u>3MG</u>	<u>A077543</u> <u>005</u>	May 18, 2011
<u>AB</u>		<u>4MG</u>	<u>A077543</u> <u>006</u>	May 18, 2011
<u>AB</u>	DR REDDYS LABS LTD	<u>0 .25MG</u>	<u>A076879</u> <u>001</u>	Oct 24, 2008
<u>AB</u>		<u>0 .5MG</u>	<u>A076879</u> <u>002</u>	Oct 24, 2008
<u>AB</u>		<u>1MG</u>	<u>A076879</u> <u>003</u>	Oct 24, 2008
<u>AB</u>		<u>2MG</u>	<u>A076879</u> <u>004</u>	Oct 24, 2008
<u>AB</u>		<u>3MG</u>	<u>A076879</u> <u>005</u>	Oct 24, 2008
<u>AB</u>		<u>4MG</u>	<u>A076879</u> <u>006</u>	Oct 24, 2008
<u>AB</u>	MYLAN	<u>0 .25MG</u>	<u>A076288</u> <u>001</u>	Sep 15, 2008
<u>AB</u>		<u>0 .5MG</u>	<u>A076288</u> <u>002</u>	Sep 15, 2008
<u>AB</u>		<u>1MG</u>	<u>A076288</u> <u>003</u>	Sep 15, 2008
<u>AB</u>		<u>2MG</u>	<u>A076288</u> <u>004</u>	Sep 15, 2008
<u>AB</u>		<u>3MG</u>	<u>A076288</u> <u>005</u>	Sep 15, 2008
<u>AB</u>		<u>4MG</u>	<u>A076288</u> <u>006</u>	Sep 15, 2008
<u>AB</u>	OXFORD PHARMS	<u>0 .25MG</u>	<u>A078071</u> <u>001</u>	Jun 17, 2009
<u>AB</u>		<u>0 .5MG</u>	<u>A078071</u> <u>002</u>	Jun 17, 2009
<u>AB</u>		<u>1MG</u>	<u>A078071</u> <u>003</u>	Jun 17, 2009
<u>AB</u>		<u>2MG</u>	<u>A078071</u> <u>004</u>	Jun 17, 2009
<u>AB</u>		<u>3MG</u>	<u>A078071</u> <u>005</u>	Jun 17, 2009
<u>AB</u>		<u>4MG</u>	<u>A078071</u> <u>006</u>	Jun 17, 2009
<u>AB</u>	PLIVA HRVATSKA DOO	<u>0 .25MG</u>	<u>A077769</u> <u>001</u>	Oct 16, 2008
<u>AB</u>		<u>0 .5MG</u>	<u>A077769</u> <u>002</u>	Oct 16, 2008
<u>AB</u>		<u>1MG</u>	<u>A077769</u> <u>003</u>	Oct 16, 2008
<u>AB</u>		<u>2MG</u>	<u>A077769</u> <u>004</u>	Oct 16, 2008
<u>AB</u>		<u>3MG</u>	<u>A077769</u> <u>005</u>	Oct 16, 2008
<u>AB</u>		<u>4MG</u>	<u>A077769</u> <u>006</u>	Oct 16, 2008
<u>AB</u>	PRINSTON INC	<u>0 .25MG</u>	<u>A077493</u> <u>001</u>	Nov 29, 2011
<u>AB</u>		<u>0 .25MG</u>	<u>A078707</u> <u>001</u>	Dec 29, 2008
<u>AB</u>		<u>0 .5MG</u>	<u>A077493</u> <u>002</u>	Nov 29, 2011
<u>AB</u>		<u>0 .5MG</u>	<u>A078707</u> <u>002</u>	Dec 29, 2008
<u>AB</u>		<u>1MG</u>	<u>A077493</u> <u>003</u>	Nov 29, 2011
<u>AB</u>		<u>1MG</u>	<u>A078707</u> <u>003</u>	Dec 29, 2008
<u>AB</u>		<u>2MG</u>	<u>A077493</u> <u>004</u>	Nov 29, 2011
<u>AB</u>		<u>2MG</u>	<u>A078707</u> <u>004</u>	Dec 29, 2008
<u>AB</u>		<u>3MG</u>	<u>A077493</u> <u>005</u>	Nov 29, 2011
<u>AB</u>		<u>3MG</u>	<u>A078707</u> <u>005</u>	Dec 29, 2008
<u>AB</u>		<u>4MG</u>	<u>A077493</u> <u>006</u>	Nov 29, 2011
<u>AB</u>		<u>4MG</u>	<u>A078707</u> <u>006</u>	Dec 29, 2008
<u>AB</u>	SANDOZ	<u>0 .25MG</u>	<u>A078528</u> <u>001</u>	Oct 16, 2009
<u>AB</u>		<u>0 .5MG</u>	<u>A078528</u> <u>002</u>	Oct 16, 2009
<u>AB</u>		<u>1MG</u>	<u>A078528</u> <u>003</u>	Oct 16, 2009
<u>AB</u>		<u>2MG</u>	<u>A078528</u> <u>004</u>	Oct 16, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-371 (of 436)

**RISPERIDONE**

TABLET; ORAL

**RISPERIDONE**

<b>AB</b>		<b>3MG</b>	<b>A078528 005</b>	Oct 16, 2009
<b>AB</b>		<b>4MG</b>	<b>A078528 006</b>	Oct 16, 2009
<b>AB</b>	SUN PHARM INDNS INC	<b>0 .25MG</b>	<b>A078036 001</b>	Mar 10, 2014
<b>AB</b>		<b>0 .5MG</b>	<b>A078036 002</b>	Mar 10, 2014
<b>AB</b>		<b>1MG</b>	<b>A078036 003</b>	Mar 10, 2014
<b>AB</b>		<b>2MG</b>	<b>A078036 004</b>	Mar 10, 2014
<b>AB</b>		<b>3MG</b>	<b>A078036 005</b>	Mar 10, 2014
<b>AB</b>		<b>4MG</b>	<b>A078036 006</b>	Mar 10, 2014
<b>AB</b>	TEVA	<b>0 .25MG</b>	<b>A076228 001</b>	Jun 30, 2008
<b>AB</b>		<b>0 .5MG</b>	<b>A076228 002</b>	Jun 30, 2008
<b>AB</b>		<b>1MG</b>	<b>A076228 003</b>	Jun 30, 2008
<b>AB</b>		<b>2MG</b>	<b>A076228 004</b>	Jun 30, 2008
<b>AB</b>		<b>3MG</b>	<b>A076228 005</b>	Jun 30, 2008
<b>AB</b>		<b>4MG</b>	<b>A076228 006</b>	Jun 30, 2008
<b>AB</b>	TORRENT PHARMS	<b>0 .25MG</b>	<b>A079088 001</b>	Oct 30, 2008
<b>AB</b>		<b>0 .5MG</b>	<b>A079088 002</b>	Oct 30, 2008
<b>AB</b>		<b>1MG</b>	<b>A079088 003</b>	Oct 30, 2008
<b>AB</b>		<b>2MG</b>	<b>A079088 004</b>	Oct 30, 2008
<b>AB</b>		<b>3MG</b>	<b>A079088 005</b>	Oct 30, 2008
<b>AB</b>		<b>4MG</b>	<b>A079088 006</b>	Oct 30, 2008
<b>AB</b>	WOCKHARDT	<b>0 .25MG</b>	<b>A078871 001</b>	Oct 09, 2008
<b>AB</b>		<b>0 .5MG</b>	<b>A078871 002</b>	Oct 09, 2008
<b>AB</b>		<b>1MG</b>	<b>A078871 003</b>	Oct 09, 2008
<b>AB</b>		<b>2MG</b>	<b>A078871 004</b>	Oct 09, 2008
<b>AB</b>		<b>3MG</b>	<b>A078871 005</b>	Oct 09, 2008
<b>AB</b>		<b>4MG</b>	<b>A078871 006</b>	Oct 09, 2008
<b>AB</b>	ZYDUS PHARMS USA INC	<b>0 .25MG</b>	<b>A078040 001</b>	Oct 16, 2008
<b>AB</b>		<b>0 .5MG</b>	<b>A078040 002</b>	Oct 16, 2008
<b>AB</b>		<b>1MG</b>	<b>A078040 003</b>	Oct 16, 2008
<b>AB</b>		<b>2MG</b>	<b>A078040 004</b>	Oct 16, 2008
<b>AB</b>		<b>3MG</b>	<b>A078040 005</b>	Oct 16, 2008
<b>AB</b>		<b>4MG</b>	<b>A078040 006</b>	Oct 16, 2008

TABLET, ORALLY DISINTEGRATING; ORAL

**RISPERDAL**

<b>AB</b>	+	JANSSEN PHARMS	<b>0 .5MG</b>	<b>N021444 001</b>	Apr 02, 2003
<b>AB</b>	+!		<b>1MG</b>	<b>N021444 002</b>	Apr 02, 2003
<b>AB</b>	+		<b>2MG</b>	<b>N021444 003</b>	Apr 02, 2003
<b>AB</b>	+		<b>3MG</b>	<b>N021444 004</b>	Dec 23, 2004
<b>AB</b>	+		<b>4MG</b>	<b>N021444 005</b>	Dec 23, 2004

**RISPERIDONE**

<b>AB</b>	ACTAVIS LABS FL INC	<b>0 .5MG</b>	<b>A076996 001</b>	Apr 19, 2011
<b>AB</b>		<b>1MG</b>	<b>A076996 002</b>	Apr 19, 2011
<b>AB</b>		<b>2MG</b>	<b>A076996 003</b>	Apr 19, 2011
<b>AB</b>		<b>3MG</b>	<b>A076996 004</b>	Apr 19, 2011
<b>AB</b>		<b>4MG</b>	<b>A076996 005</b>	Apr 19, 2011
<b>AB</b>	DR REDDYS LABS LTD	<b>0 .5MG</b>	<b>A077328 001</b>	Feb 24, 2009
<b>AB</b>		<b>1MG</b>	<b>A077328 002</b>	Oct 05, 2009
<b>AB</b>		<b>2MG</b>	<b>A077328 003</b>	Feb 24, 2009
<b>AB</b>		<b>3MG</b>	<b>A077328 004</b>	Nov 30, 2009
<b>AB</b>		<b>4MG</b>	<b>A077328 005</b>	Nov 30, 2009
<b>AB</b>	JUBILANT GENERICS	<b>0 .5MG</b>	<b>A090839 001</b>	Nov 04, 2011
<b>AB</b>		<b>1MG</b>	<b>A090839 002</b>	Nov 04, 2011
<b>AB</b>		<b>2MG</b>	<b>A090839 003</b>	Nov 04, 2011
<b>AB</b>		<b>3MG</b>	<b>A090839 004</b>	Nov 04, 2011
<b>AB</b>		<b>4MG</b>	<b>A090839 005</b>	Nov 04, 2011
<b>AB</b>	PAR PHARM	<b>0 .5MG</b>	<b>A077494 002</b>	Apr 30, 2009
<b>AB</b>		<b>1MG</b>	<b>A077494 003</b>	Oct 26, 2009
<b>AB</b>		<b>2MG</b>	<b>A077494 004</b>	Apr 30, 2009
<b>AB</b>		<b>3MG</b>	<b>A077494 005</b>	Apr 30, 2009
<b>AB</b>		<b>4MG</b>	<b>A077494 006</b>	Apr 30, 2009
<b>AB</b>	SANDOZ	<b>0 .5MG</b>	<b>A078116 001</b>	Dec 22, 2009
<b>AB</b>		<b>1MG</b>	<b>A078116 002</b>	Dec 22, 2009
<b>AB</b>		<b>2MG</b>	<b>A078116 003</b>	Dec 22, 2009
<b>AB</b>		<b>3MG</b>	<b>A078116 004</b>	Dec 22, 2009
<b>AB</b>		<b>4MG</b>	<b>A078116 005</b>	Dec 22, 2009
<b>AB</b>	SUN PHARM INDNS LTD	<b>0 .5MG</b>	<b>A077542 001</b>	Aug 06, 2010
<b>AB</b>		<b>0 .5MG</b>	<b>A078464 001</b>	Apr 08, 2013
<b>AB</b>		<b>1MG</b>	<b>A077542 002</b>	Aug 06, 2010
<b>AB</b>		<b>1MG</b>	<b>A078464 002</b>	Apr 08, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-372 (of 436)

**RISPERIDONE**

TABLET, ORALLY DISINTEGRATING;ORAL

**RISPERIDONE**

<b>AB</b>		<b>2MG</b>	<b>A077542 003</b>	Aug 06, 2010
<b>AB</b>		<b>2MG</b>	<b>A078464 003</b>	Apr 08, 2013
<b>AB</b>		<b>3MG</b>	<b>A078464 004</b>	Apr 08, 2013
<b>AB</b>		<b>3MG</b>	<b>A078474 001</b>	Aug 06, 2010
<b>AB</b>		<b>4MG</b>	<b>A078464 005</b>	Apr 08, 2013
<b>AB</b>		<b>4MG</b>	<b>A078474 002</b>	Aug 06, 2010
<b>AB</b>	TEVA	<b>0 . 5MG</b>	<b>A076908 001</b>	Mar 12, 2012
<b>AB</b>		<b>1MG</b>	<b>A076908 002</b>	Mar 12, 2012
<b>AB</b>		<b>2MG</b>	<b>A076908 003</b>	Mar 12, 2012
<b>AB</b>	ZYDUS PHARMS USA	<b>0 . 5MG</b>	<b>A078516 001</b>	May 01, 2009
<b>AB</b>		<b>2MG</b>	<b>A078516 003</b>	May 01, 2009
	PAR PHARM	0 . 25MG	A077494 001	Apr 30, 2009

**RITONAVIR**

CAPSULE;ORAL

NORVIR

+! ABBVIE

100MG

N020945 001 Jun 29, 1999

POWDER;ORAL

NORVIR

+! ABBVIE INC

100MG/PACKET

N209512 001 Jun 07, 2017

SOLUTION;ORAL

NORVIR

+! ABBVIE

80MG/ML

N020659 001 Mar 01, 1996

TABLET;ORAL

**NORVIR**

**AB** +! ABBVIE

**100MG**

**N022417 001** Feb 10, 2010

**RITONAVIR**

**AB** WEST-WARD PHARMS INT

**100MG**

**A202573 001** Jan 15, 2015

**RIVAROXABAN**

TABLET;ORAL

XARELTO

+

JANSSEN PHARMS

10MG

N022406 001 Jul 01, 2011

+

15MG

N022406 002 Nov 04, 2011

!+

20MG

N022406 003 Nov 04, 2011

**RIVASTIGMINE**

FILM, EXTENDED RELEASE;TRANSDERMAL

**EXELON**

**AB** + NOVARTIS

**4 . 6MG/24HR**

**N022083 001** Jul 06, 2007

**AB** +!

**9 . 5MG/24HR**

**N022083 002** Jul 06, 2007

**AB** +

**13 . 3MG/24HR**

**N022083 005** Aug 31, 2012

**RIVASTIGMINE**

**AB** ALVOGEN MALTA

**4 . 6MG/24HR**

**A204403 001** Sep 03, 2015

**AB**

**9 . 5MG/24HR**

**A204403 002** Sep 03, 2015

**AB**

**13 . 3MG/24HR**

**A204403 003** Aug 31, 2015

**RIVASTIGMINE TARTRATE**

CAPSULE;ORAL

**EXELON**

**AB** +! NOVARTIS

**EQ 1 . 5MG BASE**

**N020823 003** Apr 21, 2000

**AB** +

**EQ 3MG BASE**

**N020823 004** Apr 21, 2000

**AB** +

**EQ 4 . 5MG BASE**

**N020823 005** Apr 21, 2000

**AB** +!

**EQ 6MG BASE**

**N020823 006** Apr 21, 2000

**RIVASTIGMINE TARTRATE**

**AB** AJANTA PHARMA LTD

**EQ 1 . 5MG BASE**

**A207797 001** Sep 28, 2017

**AB**

**EQ 3MG BASE**

**A207797 002** Sep 28, 2017

**AB**

**EQ 4 . 5MG BASE**

**A207797 003** Sep 28, 2017

**AB**

**EQ 6MG BASE**

**A207797 004** Sep 28, 2017

**AB** ALEMBIC PHARMS LTD

**EQ 1 . 5MG BASE**

**A091689 001** Jun 12, 2012

**AB**

**EQ 3MG BASE**

**A091689 002** Jun 12, 2012

**AB**

**EQ 4 . 5MG BASE**

**A091689 003** Jun 12, 2012

**AB**

**EQ 6MG BASE**

**A091689 004** Jun 12, 2012

**AB** APOTEX INC

**EQ 1 . 5MG BASE**

**A091072 001** May 16, 2013

**AB**

**EQ 3MG BASE**

**A091072 002** May 16, 2013

**AB**

**EQ 4 . 5MG BASE**

**A091072 003** May 16, 2013

**AB**

**EQ 6MG BASE**

**A091072 004** May 16, 2013

**AB** AUROBINDO PHARMA LTD

**EQ 1 . 5MG BASE**

**A204572 001** Mar 25, 2016

**AB**

**EQ 3MG BASE**

**A204572 002** Mar 25, 2016

**AB**

**EQ 4 . 5MG BASE**

**A204572 003** Mar 25, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-373 (of 436)

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

**RIVASTIGMINE TARTRATE**

<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A204572 004</u>	Mar 25, 2016
<u>AB</u>	CADILA PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A203844 001</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A203844 002</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A203844 003</u>	Feb 13, 2017
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A203844 004</u>	Feb 13, 2017
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 1.5MG BASE</u>	<u>A077130 001</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077130 002</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077130 003</u>	Oct 31, 2007
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077130 004</u>	Oct 31, 2007
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 1.5MG BASE</u>	<u>A203148 001</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A203148 002</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A203148 003</u>	Aug 22, 2014
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A203148 004</u>	Aug 22, 2014
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 1.5MG BASE</u>	<u>A090879 001</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090879 002</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A090879 003</u>	Jun 10, 2015
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A090879 004</u>	Jun 10, 2015
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 1.5MG BASE</u>	<u>A077131 001</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077131 002</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077131 003</u>	Oct 22, 2007
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077131 004</u>	Oct 22, 2007
<u>AB</u>	WATSON LABS	<u>EQ 1.5MG BASE</u>	<u>A077129 001</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077129 002</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 4.5MG BASE</u>	<u>A077129 003</u>	Jan 08, 2008
<u>AB</u>		<u>EQ 6MG BASE</u>	<u>A077129 004</u>	Jan 08, 2008

RIZATRIPTAN BENZOATE

TABLET; ORAL

**MAXALT**

<u>AB</u>	+	MERCK	<u>EQ 5MG BASE</u>	<u>N020864 001</u>	Jun 29, 1998
<u>AB</u>	+!		<u>EQ 10MG BASE</u>	<u>N020864 002</u>	Jun 29, 1998

**RIZATRIPTAN BENZOATE**

<u>AB</u>	ALKEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A203269 001</u>	Feb 18, 2016
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203269 002</u>	Feb 18, 2016
<u>AB</u>	APOTEX INC	<u>EQ 5MG BASE</u>	<u>A202244 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202244 002</u>	Dec 31, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A202490 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202490 002</u>	Dec 31, 2012
<u>AB</u>	CIPLA LTD	<u>EQ 5MG BASE</u>	<u>A077526 001</u>	Mar 26, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077526 002</u>	Mar 26, 2013
<u>AB</u>	ECI PHARMS LLC	<u>EQ 5MG BASE</u>	<u>A202047 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202047 002</u>	Dec 31, 2012
<u>AB</u>	EMCURE PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A204090 001</u>	Nov 26, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204090 002</u>	Nov 26, 2013
<u>AB</u>	GLENMARK GENERICS	<u>EQ 5MG BASE</u>	<u>A201967 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201967 002</u>	Dec 31, 2012
<u>AB</u>	INVAGEN PHARMS	<u>EQ 5MG BASE</u>	<u>A204339 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204339 002</u>	Jul 01, 2013
<u>AB</u>	JUBILANT GENERICS	<u>EQ 5MG BASE</u>	<u>A203252 001</u>	Dec 31, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203252 002</u>	Dec 31, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A203147 001</u>	Feb 11, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203147 002</u>	Feb 11, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A201993 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201993 002</u>	Dec 31, 2012
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A200482 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A200482 002</u>	Dec 31, 2012
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>A079230 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A079230 002</u>	Dec 31, 2012
<u>AB</u>	TEVA PHARMS	<u>EQ 5MG BASE</u>	<u>A077263 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A077263 002</u>	Dec 31, 2012
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A207836 001</u>	Mar 07, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207836 002</u>	Mar 07, 2017

TABLET, ORALLY DISINTEGRATING; ORAL

**MAXALT-MLT**

<u>AB</u>	+	MERCK	<u>EQ 5MG BASE</u>	<u>N020865 001</u>	Jun 29, 1998
<u>AB</u>	+!		<u>EQ 10MG BASE</u>	<u>N020865 002</u>	Jun 29, 1998

**RIZATRIPTAN BENZOATE**

<u>AB</u>	APOTEX INC	<u>EQ 5MG BASE</u>	<u>A202477 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A202477 002</u>	Jul 01, 2013

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-374 (of 436)

RIZATRIPTAN BENZOATE

TABLET, ORALLY DISINTEGRATING;ORAL

**RIZATRIPTAN BENZOATE**

<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A203062 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203062 002</u>	Jul 01, 2013
<u>AB</u>	GLENMARK PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A201914 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A201914 002</u>	Jul 01, 2013
<u>AB</u>	JUBILANT GENERICS	<u>EQ 5MG BASE</u>	<u>A203334 001</u>	Oct 16, 2015
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203334 002</u>	Oct 16, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 5MG BASE</u>	<u>A203146 001</u>	Sep 19, 2014
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203146 002</u>	Sep 19, 2014
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 5MG BASE</u>	<u>A078173 001</u>	Dec 31, 2012
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078173 002</u>	Dec 31, 2012
<u>AB</u>	NATCO PHARMA LTD	<u>EQ 5MG BASE</u>	<u>A203478 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A203478 002</u>	Jul 01, 2013
<u>AB</u>	PANACEA BIOTEC LTD	<u>EQ 5MG BASE</u>	<u>A204722 001</u>	Jan 11, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A204722 002</u>	Jan 11, 2017
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>A078739 001</u>	Jul 01, 2013
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A078739 002</u>	Jul 01, 2013
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 5MG BASE</u>	<u>A207835 001</u>	Mar 07, 2017
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A207835 002</u>	Mar 07, 2017

ROCURONIUM BROMIDE

INJECTABLE;INJECTION

**ROCURONIUM BROMIDE**

<u>AP</u>	AUROBINDO PHARMA LTD	<u>50MG/5ML (10MG/ML)</u>	<u>A206206 001</u>	Apr 12, 2017
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A206206 002</u>	Apr 12, 2017
<u>AP</u>	FRESENIUS KABI USA	<u>50MG/5ML (10MG/ML)</u>	<u>A078651 001</u>	Dec 29, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078651 002</u>	Dec 29, 2008
<u>AP</u>	HOSPIRA	<u>50MG/5ML (10MG/ML)</u>	<u>A078519 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078519 002</u>	Nov 26, 2008
<u>AP</u>	MYLAN INSTITUTIONAL	<u>50MG/5ML (10MG/ML)</u>	<u>A079199 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A079199 002</u>	Nov 26, 2008
<u>AP</u>	SAGENT PHARMS	<u>50MG/5ML (10MG/ML)</u>	<u>A091458 001</u>	Jul 28, 2010
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A091458 002</u>	Jul 28, 2010
<u>AP</u>	! SANDOZ INC	<u>50MG/5ML (10MG/ML)</u>	<u>A079195 001</u>	Dec 05, 2008
<u>AP</u>	!	<u>100MG/10ML (10MG/ML)</u>	<u>A079195 002</u>	Dec 05, 2008
<u>AP</u>	TAMARANG	<u>50MG/5ML (10MG/ML)</u>	<u>A091115 001</u>	Aug 27, 2012
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A091115 002</u>	Aug 27, 2012
<u>AP</u>	TEVA PHARMS	<u>50MG/5ML (10MG/ML)</u>	<u>A078717 001</u>	Nov 26, 2008
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A078717 002</u>	Nov 26, 2008
<u>AP</u>	WEST WARD PHARM CORP	<u>50MG/5ML (10MG/ML)</u>	<u>A204679 001</u>	Feb 28, 2017
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A204679 002</u>	Feb 28, 2017

ROFLUMILAST

TABLET;ORAL

DALIRESP

+! ASTRAZENECA PHARMS 500MCG

N022522 001 Feb 28, 2011

ROLAPITANT HYDROCHLORIDE

EMULSION;IV (INFUSION)

VARUBI

+! TESARO INC

EQ 166.5MG BASE/92.5ML (EQ 1.8MG BASE/ML)

N208399 001 Oct 25, 2017

TABLET;ORAL

VARUBI

+! TESARO INC

EQ 90MG BASE

N206500 001 Sep 01, 2015

ROMIDEPSIN

POWDER;IV (INFUSION)

ISTODAX

+! CELGENE

10MG/VIAL

N022393 001 Nov 05, 2009

ROPINIROLE HYDROCHLORIDE

TABLET;ORAL

**REQUIP**

<u>AB</u>	+! GLAXOSMITHKLINE LLC	<u>EQ 0.25MG BASE</u>	<u>N020658 001</u>	Sep 19, 1997
<u>AB</u>	+	<u>EQ 0.5MG BASE</u>	<u>N020658 002</u>	Sep 19, 1997
<u>AB</u>	+	<u>EQ 1MG BASE</u>	<u>N020658 003</u>	Sep 19, 1997
<u>AB</u>	+	<u>EQ 2MG BASE</u>	<u>N020658 004</u>	Sep 19, 1997
<u>AB</u>	+	<u>EQ 3MG BASE</u>	<u>N020658 006</u>	Jan 27, 1999
<u>AB</u>	+	<u>EQ 4MG BASE</u>	<u>N020658 007</u>	Jan 27, 1999
<u>AB</u>	+	<u>EQ 5MG BASE</u>	<u>N020658 005</u>	Sep 19, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-375 (of 436)

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 0 .25MG BASE</u>	<u>A204022 001</u>	Feb 28, 2017
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A204022 002</u>	Feb 28, 2017
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A204022 003</u>	Feb 28, 2017
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A204022 004</u>	Feb 28, 2017
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A204022 005</u>	Feb 28, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A204022 006</u>	Feb 28, 2017
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A204022 007</u>	Feb 28, 2017
<u>AB</u>	ALEMBIC LTD	<u>EQ 0 .25MG BASE</u>	<u>A090429 001</u>	Mar 24, 2010
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A090429 002</u>	Mar 24, 2010
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A090429 003</u>	Mar 24, 2010
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A090429 004</u>	Mar 24, 2010
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090429 005</u>	Mar 24, 2010
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A090429 006</u>	Mar 24, 2010
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090429 007</u>	Mar 24, 2010
<u>AB</u>	APOTEX	<u>EQ 0 .25MG BASE</u>	<u>A079165 001</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A079165 002</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079165 003</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079165 004</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079165 005</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079165 006</u>	Feb 07, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079165 007</u>	Feb 07, 2012
<u>AB</u>	G AND W LABS INC	<u>EQ 0 .25MG BASE</u>	<u>A077460 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A077460 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A077460 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A077460 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077460 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A077460 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077460 007</u>	May 19, 2008
<u>AB</u>	GLENMARK GENERICS	<u>EQ 0 .25MG BASE</u>	<u>A090135 001</u>	Feb 25, 2010
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A090135 002</u>	Feb 25, 2010
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A090135 003</u>	Feb 25, 2010
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A090135 004</u>	Feb 25, 2010
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090135 005</u>	Feb 25, 2010
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A090135 006</u>	Feb 25, 2010
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090135 007</u>	Feb 25, 2010
<u>AB</u>	MYLAN	<u>EQ 0 .25MG BASE</u>	<u>A078881 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A078881 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A078881 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A078881 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A078881 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A078881 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078881 007</u>	May 19, 2008
<u>AB</u>	ORCHID HLTHCARE	<u>EQ 0 .25MG BASE</u>	<u>A079229 001</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A079229 002</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079229 003</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079229 004</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079229 005</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079229 006</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079229 007</u>	Nov 28, 2012
<u>AB</u>	PRINSTON INC	<u>EQ 0 .25MG BASE</u>	<u>A078110 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A078110 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A078110 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A078110 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A078110 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A078110 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A078110 007</u>	Jul 11, 2008
<u>AB</u>	WEST-WARD PHARMS INT	<u>EQ 0 .25MG BASE</u>	<u>A077852 001</u>	May 05, 2008
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A077852 002</u>	May 05, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A077852 003</u>	May 05, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A077852 004</u>	May 05, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A077852 005</u>	May 05, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A077852 006</u>	May 05, 2008
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A077852 007</u>	May 19, 2008
<u>AB</u>	WOCKHARDT	<u>EQ 0 .25MG BASE</u>	<u>A079050 001</u>	May 29, 2008
<u>AB</u>		<u>EQ 0 .5MG BASE</u>	<u>A079050 002</u>	May 29, 2008
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A079050 003</u>	May 29, 2008
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A079050 004</u>	May 29, 2008
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A079050 005</u>	May 29, 2008
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A079050 006</u>	May 29, 2008

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-376 (of 436)

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A079050 007</u>	May 29, 2008
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 0.25MG BASE</u>	<u>A090411 001</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 0.5MG BASE</u>	<u>A090411 002</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>A090411 003</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A090411 004</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 3MG BASE</u>	<u>A090411 005</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A090411 006</u>	Jun 01, 2009
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A090411 007</u>	Jun 01, 2009

TABLET, EXTENDED RELEASE; ORAL

REQUIP XL

<u>AB</u>	+!	GLAXOSMITHKLINE LLC	<u>EQ 2MG BASE</u>	<u>N022008 001</u>	Jun 13, 2008
<u>AB</u>	+		<u>EQ 4MG BASE</u>	<u>N022008 003</u>	Jun 13, 2008
<u>AB</u>	+		<u>EQ 6MG BASE</u>	<u>N022008 006</u>	Apr 10, 2009
<u>AB</u>	+		<u>EQ 8MG BASE</u>	<u>N022008 004</u>	Jun 13, 2008
<u>AB</u>	+		<u>EQ 12MG BASE</u>	<u>N022008 005</u>	Oct 31, 2008

ROPINIROLE HYDROCHLORIDE

<u>AB</u>		ACTAVIS ELIZABETH	<u>EQ 2MG BASE</u>	<u>A090869 001</u>	May 17, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A090869 002</u>	May 17, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A090869 003</u>	May 17, 2012
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A090869 004</u>	May 17, 2012
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A090869 005</u>	May 17, 2012
<u>AB</u>		ALEMBIC PHARMS LTD	<u>EQ 2MG BASE</u>	<u>A202786 001</u>	Apr 22, 2013
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A202786 002</u>	Apr 22, 2013
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A202786 003</u>	Apr 22, 2013
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A202786 004</u>	Apr 22, 2013
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A202786 005</u>	Apr 22, 2013
<u>AB</u>		DR REDDYS LABS LTD	<u>EQ 2MG BASE</u>	<u>A201576 001</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A201576 002</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A201576 003</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A201576 004</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A201576 005</u>	Jun 06, 2012
<u>AB</u>		MYLAN PHARMS INC	<u>EQ 2MG BASE</u>	<u>A200462 001</u>	Oct 15, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A200462 003</u>	Oct 15, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A200462 004</u>	Oct 15, 2012
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A200462 005</u>	Oct 15, 2012
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A200462 006</u>	Oct 15, 2012
<u>AB</u>		SANDOZ INC	<u>EQ 2MG BASE</u>	<u>A201047 001</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A201047 003</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A201047 004</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A201047 005</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A201047 006</u>	Jun 06, 2012
<u>AB</u>		WATSON LABS INC	<u>EQ 2MG BASE</u>	<u>A200431 001</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A200431 002</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A200431 003</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A200431 004</u>	Jun 06, 2012
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A200431 005</u>	Jun 06, 2012
<u>AB</u>		WOCKHARDT LTD	<u>EQ 2MG BASE</u>	<u>A091395 001</u>	Aug 27, 2012
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A091395 002</u>	Aug 27, 2012
<u>AB</u>			<u>EQ 6MG BASE</u>	<u>A091395 003</u>	Aug 27, 2012
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A091395 004</u>	Aug 27, 2012
<u>AB</u>			<u>EQ 12MG BASE</u>	<u>A091395 005</u>	Aug 27, 2012

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

NAROPIN

<u>AP</u>	+	FRESENIUS KABI USA	<u>20MG/10ML (2MG/ML)</u>	<u>N020533 001</u>	May 01, 1998
<u>AP</u>	+		<u>40MG/20ML (2MG/ML)</u>	<u>N020533 002</u>	Sep 24, 1996
<u>AP</u>	+		<u>100MG/20ML (5MG/ML)</u>	<u>N020533 003</u>	May 01, 1998
<u>AP</u>	+		<u>100MG/10ML (10MG/ML)</u>	<u>N020533 005</u>	Sep 24, 1996
<u>AP</u>	+		<u>150MG/20ML (7.5MG/ML)</u>	<u>N020533 004</u>	Sep 24, 1996
<u>AP</u>	+		<u>150MG/30ML (5MG/ML)</u>	<u>N020533 008</u>	Sep 24, 1996
<u>AP</u>	+		<u>200MG/100ML (2MG/ML)</u>	<u>N020533 006</u>	Sep 24, 1996
<u>AP</u>	+		<u>200MG/20ML (10MG/ML)</u>	<u>N020533 011</u>	Sep 24, 1996

ROPIVACAINE HYDROCHLORIDE

<u>AP</u>		AKORN INC	<u>150MG/30ML (5MG/ML)</u>	<u>A203955 001</u>	Apr 11, 2016
<u>AP</u>		AUROBINDO PHARMA LTD	<u>40MG/20ML (2MG/ML)</u>	<u>A205612 001</u>	Jul 13, 2016
<u>AP</u>			<u>100MG/20ML (5MG/ML)</u>	<u>A205612 003</u>	Jul 13, 2016
<u>AP</u>			<u>100MG/10ML (10MG/ML)</u>	<u>A205612 006</u>	Jul 13, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-377 (of 436)

ROPIVACAINE HYDROCHLORIDE

SOLUTION; INJECTION

ROPIVACAINE HYDROCHLORIDE

<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A205612 004</u>	Jul 13, 2016
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A205612 005</u>	Jul 13, 2016
<u>AP</u>		<u>200MG/100ML (2MG/ML)</u>	<u>A205612 002</u>	Jul 13, 2016
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A205612 007</u>	Jul 13, 2016
<u>AP</u>	HOSPIRA	<u>20MG/10ML (2MG/ML)</u>	<u>A090194 001</u>	Sep 23, 2014
<u>AP</u>		<u>40MG/20ML (2MG/ML)</u>	<u>A090194 005</u>	Sep 23, 2014
<u>AP</u>		<u>100MG/10ML (10MG/ML)</u>	<u>A090194 004</u>	Sep 23, 2014
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A090194 002</u>	Sep 23, 2014
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A090194 003</u>	Sep 23, 2014
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A090194 006</u>	Sep 23, 2014
<u>AP</u>	NAVINTA LLC	<u>150MG/30ML (5MG/ML)</u>	<u>A078601 002</u>	Jul 17, 2014
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A078601 003</u>	Jul 17, 2014
<u>AP</u>	SAGENT STRIDES	<u>40MG/20ML (2MG/ML)</u>	<u>A090318 001</u>	Sep 23, 2014
<u>AP</u>		<u>150MG/30ML (5MG/ML)</u>	<u>A090318 002</u>	Sep 23, 2014
<u>AP</u>		<u>150MG/20ML (7.5MG/ML)</u>	<u>A090318 003</u>	Sep 23, 2014
<u>AP</u>		<u>200MG/20ML (10MG/ML)</u>	<u>A090318 004</u>	Sep 23, 2014
<u>NAROPIN</u>				
+ FRESENIUS KABI USA		<u>400MG/200ML (2MG/ML)</u>	<u>N020533 007</u>	Sep 24, 1996
+		<u>500MG/100ML (5MG/ML)</u>	<u>N020533 009</u>	Jan 04, 2011
+		<u>1GM/200ML (5MG/ML)</u>	<u>N020533 010</u>	Jan 04, 2011

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

<u>AB</u>	+	SB PHARMCO	<u>EQ 2MG BASE</u>	<u>N021071 002</u>	May 25, 1999
<u>AB</u>	+		<u>EQ 4MG BASE</u>	<u>N021071 003</u>	May 25, 1999
<u>AB</u>	+	!	<u>EQ 8MG BASE</u>	<u>N021071 004</u>	May 25, 1999
<u>ROSIGLITAZONE MALEATE</u>					
<u>AB</u>		TEVA	<u>EQ 2MG BASE</u>	<u>A076747 001</u>	Jan 25, 2013
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>A076747 002</u>	Jan 25, 2013
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>A076747 003</u>	Jan 25, 2013

ROSUVASTATIN CALCIUM

TABLET; ORAL

CRESTOR

<u>AB</u>	+	IPR	<u>5MG</u>	<u>N021366 002</u>	Aug 12, 2003
<u>AB</u>	+		<u>10MG</u>	<u>N021366 003</u>	Aug 12, 2003
<u>AB</u>	+		<u>20MG</u>	<u>N021366 004</u>	Aug 12, 2003
<u>AB</u>	+	!	<u>40MG</u>	<u>N021366 005</u>	Aug 12, 2003

ROSVUSTATIN CALCIUM

<u>AB</u>		ACCORD HLTHCARE	<u>5MG</u>	<u>A206434 001</u>	Oct 31, 2016
<u>AB</u>			<u>10MG</u>	<u>A206434 002</u>	Oct 31, 2016
<u>AB</u>			<u>20MG</u>	<u>A206434 003</u>	Oct 31, 2016
<u>AB</u>			<u>40MG</u>	<u>A206434 004</u>	Oct 31, 2016
<u>AB</u>		ALKEM LABS LTD	<u>5MG</u>	<u>A206465 001</u>	Mar 21, 2017
<u>AB</u>			<u>10MG</u>	<u>A206465 002</u>	Mar 21, 2017
<u>AB</u>			<u>20MG</u>	<u>A206465 003</u>	Mar 21, 2017
<u>AB</u>			<u>40MG</u>	<u>A206465 004</u>	Mar 21, 2017
<u>AB</u>		APOTEX INC	<u>5MG</u>	<u>A079145 001</u>	Jul 19, 2016
<u>AB</u>			<u>10MG</u>	<u>A079145 002</u>	Jul 19, 2016
<u>AB</u>			<u>20MG</u>	<u>A079145 003</u>	Jul 19, 2016
<u>AB</u>			<u>40MG</u>	<u>A079145 004</u>	Jul 19, 2016
<u>AB</u>		AUROBINDO PHARMA LTD	<u>5MG</u>	<u>A079170 001</u>	Jul 19, 2016
<u>AB</u>			<u>10MG</u>	<u>A079170 002</u>	Jul 19, 2016
<u>AB</u>			<u>20MG</u>	<u>A079170 003</u>	Jul 19, 2016
<u>AB</u>			<u>40MG</u>	<u>A079170 004</u>	Jul 19, 2016
<u>AB</u>		BIOCON LTD	<u>5MG</u>	<u>A207752 001</u>	Oct 31, 2016
<u>AB</u>			<u>10MG</u>	<u>A207752 002</u>	Oct 31, 2016
<u>AB</u>			<u>20MG</u>	<u>A207752 003</u>	Oct 31, 2016
<u>AB</u>			<u>40MG</u>	<u>A207752 004</u>	Oct 31, 2016
<u>AB</u>		CADILA PHARMS LTD	<u>5MG</u>	<u>A207453 001</u>	Nov 23, 2016
<u>AB</u>			<u>10MG</u>	<u>A207453 002</u>	Nov 23, 2016
<u>AB</u>			<u>20MG</u>	<u>A207453 003</u>	Nov 23, 2016
<u>AB</u>			<u>40MG</u>	<u>A207453 004</u>	Nov 23, 2016
<u>AB</u>		CHANGZHOU PHARM	<u>5MG</u>	<u>A207408 001</u>	Oct 31, 2016
<u>AB</u>			<u>10MG</u>	<u>A207408 002</u>	Oct 31, 2016
<u>AB</u>			<u>20MG</u>	<u>A207408 003</u>	Oct 31, 2016
<u>AB</u>			<u>40MG</u>	<u>A207408 004</u>	Oct 31, 2016
<u>AB</u>		GLENMARK PHARMS	<u>5MG</u>	<u>A079172 001</u>	Jul 19, 2016
<u>AB</u>			<u>10MG</u>	<u>A079172 002</u>	Jul 19, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-378 (of 436)

ROSUVASTATIN CALCIUM

TABLET;ORAL

**ROSUVASTATIN CALCIUM**

<u>AB</u>		<u>20MG</u>	<u>A079172 003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079172 004</u>	Jul 19, 2016
<u>AB</u>	HETERO LABS LTD V	<u>5MG</u>	<u>A207616 001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A207616 002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A207616 003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A207616 004</u>	Oct 31, 2016
<u>AB</u>	JUBILANT GENERICS	<u>5MG</u>	<u>A207062 001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A207062 002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A207062 003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A207062 004</u>	Oct 31, 2016
<u>AB</u>	LUPIN LTD	<u>5MG</u>	<u>A205587 001</u>	Jul 31, 2017
<u>AB</u>		<u>10MG</u>	<u>A205587 002</u>	Jul 31, 2017
<u>AB</u>		<u>20MG</u>	<u>A205587 003</u>	Jul 31, 2017
<u>AB</u>		<u>40MG</u>	<u>A205587 004</u>	Jul 31, 2017
<u>AB</u>	MSN LABS PVT LTD	<u>5MG</u>	<u>A208898 001</u>	Nov 22, 2017
<u>AB</u>		<u>10MG</u>	<u>A208898 002</u>	Nov 22, 2017
<u>AB</u>		<u>20MG</u>	<u>A208898 003</u>	Nov 22, 2017
<u>AB</u>		<u>40MG</u>	<u>A208898 004</u>	Nov 22, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A079161 001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079161 002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079161 003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079161 004</u>	Jul 19, 2016
<u>AB</u>	PAR PHARM INC	<u>5MG</u>	<u>A079168 001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079168 002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079168 003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079168 004</u>	Jul 19, 2016
<u>AB</u>	SANDOZ INC	<u>5MG</u>	<u>A079171 001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079171 002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079171 003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079171 004</u>	Jul 19, 2016
<u>AB</u>	SUN PHARMA GLOBAL	<u>5MG</u>	<u>A079169 001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079169 002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079169 003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079169 004</u>	Jul 19, 2016
<u>AB</u>	TEVA PHARMS USA	<u>5MG</u>	<u>A079166 001</u>	Jul 19, 2016
<u>AB</u>		<u>10MG</u>	<u>A079166 002</u>	Jul 19, 2016
<u>AB</u>		<u>20MG</u>	<u>A079166 003</u>	Jul 19, 2016
<u>AB</u>		<u>40MG</u>	<u>A079166 004</u>	Jul 19, 2016
<u>AB</u>	TORRENT PHARMS LTD	<u>5MG</u>	<u>A201619 001</u>	Oct 31, 2016
<u>AB</u>		<u>10MG</u>	<u>A201619 002</u>	Oct 31, 2016
<u>AB</u>		<u>20MG</u>	<u>A201619 003</u>	Oct 31, 2016
<u>AB</u>		<u>40MG</u>	<u>A201619 004</u>	Oct 31, 2016
<u>AB</u>	WATSON LABS INC	<u>5MG</u>	<u>A079167 001</u>	Apr 29, 2016
<u>AB</u>		<u>10MG</u>	<u>A079167 002</u>	Apr 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A079167 003</u>	Apr 29, 2016
<u>AB</u>		<u>40MG</u>	<u>A079167 004</u>	Apr 29, 2016

ROTIGOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NEUPRO

+	UCB INC	1MG/24HR	N021829 004	Apr 02, 2012
+		2MG/24HR	N021829 001	May 09, 2007
+		3MG/24HR	N021829 005	Apr 02, 2012
+		4MG/24HR	N021829 002	May 09, 2007
+		6MG/24HR	N021829 003	May 09, 2007
+		8MG/24HR	N021829 006	Apr 02, 2012

RUBIDIUM CHLORIDE RB-82

INJECTABLE;INJECTION

CARDIOGEN-82

BRACCO

N/A

N019414 001 Dec 29, 1989

SOLUTION;INTRAVENOUS

RUBY-FILL

JUBILANT DRAXIMAGE N/A

N202153 001 Sep 30, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-379 (of 436)

RUCAPARIB CAMSYLATE

TABLET;ORAL

RUBRACA

+ CLOVIS ONCOLOGY INC	EQ 200MG BASE
+ CLOVIS ONCOLOGY INC	EQ 250MG BASE
+! CLOVIS ONCOLOGY INC	EQ 300MG BASE

N209115 001	Dec 19, 2016
N209115 003	May 01, 2017
N209115 002	Dec 19, 2016

RUFINAMIDE

SUSPENSION;ORAL

BANZEL

+! EISAI INC	40MG/ML
--------------	---------

N201367 001	Mar 03, 2011
-------------	--------------

TABLET;ORAL

**BANZEL**

<b>AB</b> + EISAI INC	<b>200MG</b>	<b>N021911 002</b> Nov 14, 2008
<b>AB</b> +!	<b>400MG</b>	<b>N021911 003</b> Nov 14, 2008

RUFINAMIDE

<b>AB</b> GLENMARK PHARMS LTD	<b>200MG</b>	<b>A205075 001</b> May 16, 2016
-------------------------------	--------------	---------------------------------

<b>AB</b>	<b>400MG</b>	<b>A205075 002</b> May 16, 2016
-----------	--------------	---------------------------------

<b>AB</b> MYLAN PHARMS INC	<b>200MG</b>	<b>A205095 001</b> May 16, 2016
----------------------------	--------------	---------------------------------

<b>AB</b>	<b>400MG</b>	<b>A205095 002</b> May 16, 2016
-----------	--------------	---------------------------------

<b>AB</b> WEST-WARD PHARMS INT	<b>200MG</b>	<b>A204988 001</b> May 16, 2016
--------------------------------	--------------	---------------------------------

<b>AB</b>	<b>400MG</b>	<b>A204988 002</b> May 16, 2016
-----------	--------------	---------------------------------

RUXOLITINIB PHOSPHATE

TABLET;ORAL

JAKAFI

+ INCYTE CORP	EQ 5MG BASE
+	EQ 10MG BASE
+	EQ 15MG BASE
+	EQ 20MG BASE
+!	EQ 25MG BASE

N202192 001	Nov 16, 2011
N202192 002	Nov 16, 2011
N202192 003	Nov 16, 2011
N202192 004	Nov 16, 2011
N202192 005	Nov 16, 2011

SACROSIDASE

SOLUTION;ORAL

SUCRAID

+! QOL MEDCL	8,500 IU/ML
--------------	-------------

N020772 001	Apr 09, 1998
-------------	--------------

SACUBITRIL; VALSARTAN

TABLET;ORAL

ENTRESTO

+ NOVARTIS PHARMS CORP	24MG;26MG
+	49MG;51MG
+!	97MG;103MG

N207620 001	Jul 07, 2015
N207620 002	Jul 07, 2015
N207620 003	Jul 07, 2015

SAFINAMIDE MESYLATE

TABLET;ORAL

XADAGO

+ US WORLDMEDS LLC	50MG
+!	100MG

N207145 001	Mar 21, 2017
N207145 002	Mar 21, 2017

SALMETEROL XINAFOATE

POWDER;INHALATION

SEREVENT

+! GLAXOSMITHKLINE	EQ 0.05MG BASE/INH
--------------------	--------------------

N020692 001	Sep 19, 1997
-------------	--------------

SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

INJECTABLE;INJECTION

QUADRAMET

+! LANTHEUS MEDICAL	50mCi/ML
---------------------	----------

N020570 001	Mar 28, 1997
-------------	--------------

SAPROPTERIN DIHYDROCHLORIDE

POWDER;ORAL

KUVAN

+! BIOMARIN PHARM	100MG/PACKET
+	500MG/PACKET

N205065 001	Dec 19, 2013
N205065 002	Oct 27, 2015

TABLET;ORAL

KUVAN

+! BIOMARIN PHARM	100MG
-------------------	-------

N022181 001	Dec 13, 2007
-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-380 (of 436)

SAQUINAVIR MESYLATE

CAPSULE;ORAL			
INVIRASE			
+! HOFFMANN LA ROCHE	EQ 200MG BASE		N020628 001 Dec 06, 1995
TABLET;ORAL			
INVIRASE			
+! HOFFMANN-LA ROCHE	EQ 500MG BASE		N021785 001 Dec 17, 2004

SAXAGLIPTIN HYDROCHLORIDE

TABLET;ORAL			
ONGLYZA			
+ ASTRAZENECA AB	EQ 2.5MG BASE		N022350 001 Jul 31, 2009
+!	EQ 5MG BASE		N022350 002 Jul 31, 2009

SCOPOLAMINE

FILM, EXTENDED RELEASE;TRANSDERMAL

<u>SCOPOLAMINE</u>			
<u>AB</u> PERRIGO PHARMS CO	<u>1MG/72HR</u>		<u>A078830 001</u> Jan 30, 2015
<u>TRANSDERM SCOP</u>			
<u>AB</u> +! GLAXOSMITHKLINE CON	<u>1MG/72HR</u>		<u>N017874 001</u>

SECNIDAZOLE

GRANULE;ORAL			
SOLOSEC			
+! LUPIN	2GM/PACKET		N209363 001 Sep 15, 2017

SECOBARBITAL SODIUM

CAPSULE;ORAL			
SECONAL SODIUM			
+! VALEANT PHARMS NORTH	50MG		A086101 001 Oct 03, 1983
!	100MG		A086101 002 Oct 03, 1983

SECRETIN SYNTHETIC HUMAN

FOR SOLUTION;INTRAVENOUS			
CHIRHOSTIM			
+! CHIRHOCLIN	16MCG/VIAL		N021256 001 Apr 09, 2004
+	40MCG/VIAL		N021256 002 Jun 21, 2007

SELEGILINE

FILM, EXTENDED RELEASE;TRANSDERMAL			
EMSAM			
+! SOMERSET	6MG/24HR		N021336 001 Feb 27, 2006
+	9MG/24HR		N021336 002 Feb 27, 2006
+	12MG/24HR		N021336 003 Feb 27, 2006

SELEGILINE HYDROCHLORIDE

CAPSULE;ORAL			
<u>ELDEPRYL</u>			
<u>AB</u> +! SOMERSET	<u>5MG</u>		<u>N020647 001</u> May 15, 1996
<u>SELEGILINE HYDROCHLORIDE</u>			
<u>AB</u> APOTEX	<u>5MG</u>		<u>A075321 001</u> Dec 04, 1998
<u>AB</u> DAVA PHARMS INC	<u>5MG</u>		<u>A075352 001</u> Nov 30, 1998
TABLET;ORAL			
<u>SELEGILINE HYDROCHLORIDE</u>			
<u>AB</u> ! APOTEX INC	<u>5MG</u>		<u>A074871 001</u> Jun 06, 1997
<u>AB</u> BOSCOGEN	<u>5MG</u>		<u>A074912 001</u> Apr 30, 1998
<u>AB</u> MYLAN	<u>5MG</u>		<u>A074866 001</u> Nov 26, 1997
TABLET, ORALLY DISINTEGRATING;ORAL			
ZELAPAR			
+! VALEANT PHARM INTL	1.25MG		N021479 001 Jun 14, 2006

SELENIUM SULFIDE

LOTION/SHAMPOO;TOPICAL			
<u>SELENIUM SULFIDE</u>			
<u>AT</u> PERRIGO NEW YORK	<u>2.5%</u>		<u>A089996 001</u> Jan 10, 1991
<u>AT</u> WOCKHARDT BIO AG	<u>2.5%</u>		<u>A088228 001</u> Sep 01, 1983
<u>SELSUN</u>			
<u>AT</u> +! CHATTEM	<u>2.5%</u>		<u>N007936 001</u>

SELEXIPAG

TABLET;ORAL			
UPTRAVI			
+ ACTELION PHARMS LTD	0.2MG		N207947 001 Dec 21, 2015
+	0.4MG		N207947 002 Dec 21, 2015
+	0.6MG		N207947 003 Dec 21, 2015
+	0.8MG		N207947 004 Dec 21, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-381 (of 436)

SELEXIPAG

TABLET;ORAL  
 UPTRAVI

+	1MG	N207947 005	Dec 21, 2015
+	1.2MG	N207947 006	Dec 21, 2015
+	1.4MG	N207947 007	Dec 21, 2015
+!	1.6MG	N207947 008	Dec 21, 2015

SEMAGLUTIDE

SOLUTION;SUBCUTANEOUS  
 OZEMPIC

+!	NOVO NORDISK INC	2MG/1.5ML (1.34MG/ML)	N209637 001	Dec 05, 2017
----	------------------	-----------------------	-------------	--------------

SERTACONAZOLE NITRATE

CREAM;TOPICAL  
 ERTACZO

+!	VALEANT LUXEMBOURG	2%	N021385 001	Dec 10, 2003
----	--------------------	----	-------------	--------------

SERTRALINE HYDROCHLORIDE

CONCENTRATE;ORAL

SERTRALINE HYDROCHLORIDE

<u>AA</u>	AUROBINDO PHARMA	<u>EQ 20MG BASE/ML</u>	<u>A078861 001</u>	Oct 31, 2008
-----------	------------------	------------------------	--------------------	--------------

ZOLOFT

<u>AA</u>	+!	PFIZER	<u>EQ 20MG BASE/ML</u>
-----------	----	--------	------------------------

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

<u>AB</u>	ACCORD HLTHCARE	<u>EQ 25MG BASE</u>	<u>A202825 001</u>	Nov 07, 2014
-----------	-----------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202825 002</u>	Nov 07, 2014
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202825 003</u>	Nov 07, 2014
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	APOTEX INC	<u>EQ 25MG BASE</u>	<u>A076882 001</u>	Feb 06, 2007
-----------	------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076882 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076882 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A077206 001</u>	Feb 06, 2007
-----------	------------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077206 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077206 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	AUSTARPHARMA LLC	<u>EQ 25MG BASE</u>	<u>A078677 001</u>	Mar 04, 2009
-----------	------------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078677 002</u>	Mar 04, 2009
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078677 003</u>	Mar 04, 2009
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	INVAGEN PHARMS	<u>EQ 25MG BASE</u>	<u>A077397 001</u>	Feb 06, 2007
-----------	----------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077397 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077397 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	LUPIN	<u>EQ 25MG BASE</u>	<u>A077670 001</u>	Feb 06, 2007
-----------	-------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077670 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077670 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	MYLAN PHARMS INC	<u>EQ 25MG BASE</u>	<u>A078626 001</u>	Jan 31, 2008
-----------	------------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078626 002</u>	Jan 31, 2008
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078626 003</u>	Jan 31, 2008
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	OXFORD PHARMS	<u>EQ 25MG BASE</u>	<u>A078175 001</u>	Jul 21, 2010
-----------	---------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078175 002</u>	Jul 21, 2010
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078175 003</u>	Jul 21, 2010
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 25MG BASE</u>	<u>A077977 001</u>	Feb 06, 2007
-----------	--------------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077977 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077977 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076465 001</u>	Aug 11, 2006
-----------	------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076465 002</u>	Aug 11, 2006
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076465 003</u>	Aug 11, 2006
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	TORRENT PHARMS	<u>EQ 25MG BASE</u>	<u>A077765 001</u>	Feb 06, 2007
-----------	----------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077765 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077765 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	WOCKHARDT	<u>EQ 25MG BASE</u>	<u>A078403 001</u>	Jan 08, 2008
-----------	-----------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078403 002</u>	Jan 08, 2008
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078403 003</u>	Jan 08, 2008
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A077106 001</u>	Feb 06, 2007
-----------	------------------	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077106 002</u>	Feb 06, 2007
-----------	--	---------------------	--------------------	--------------

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077106 003</u>	Feb 06, 2007
-----------	--	----------------------	--------------------	--------------

<u>AB</u>	<u>ZOLOFT</u>	<u>EQ 25MG BASE</u>	<u>N019839 005</u>	Mar 06, 1996
-----------	---------------	---------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-382 (of 436)

SEVELAMER CARBONATE

FOR SUSPENSION;ORAL

RENELA

<u>AB</u>	+	GENZYME	<u>800MG/PACKET</u>	<u>N022318 001</u>	Aug 12, 2009
<u>AB</u>	+!		<u>2.4GM/PACKET</u>	<u>N022318 002</u>	Feb 18, 2009

SEVELAMER CARBONATE

<u>AB</u>	AUROBINDO PHARMA LTD	<u>800MG/PACKET</u>	<u>A207624 001</u>	Jun 13, 2017
<u>AB</u>		<u>2.4GM/PACKET</u>	<u>A207624 002</u>	Jun 13, 2017

TABLET;ORAL

RENELA

<u>AB</u>	+!	GENZYME	<u>800MG</u>	<u>N022127 001</u>	Oct 19, 2007
-----------	----	---------	--------------	--------------------	--------------

SEVELAMER CARBONATE

<u>AB</u>	AMNEAL PHARMS CO	<u>800MG</u>	<u>A207288 001</u>	Nov 28, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>800MG</u>	<u>A207179 001</u>	Jul 17, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>800MG</u>	<u>A206094 001</u>	Sep 29, 2017
<u>AB</u>	IMPAX LABS INC	<u>800MG</u>	<u>A090975 001</u>	Oct 23, 2017
<u>AB</u>	INVAGEN PHARMS	<u>800MG</u>	<u>A203860 001</u>	Oct 26, 2017

SEVELAMER HYDROCHLORIDE

TABLET;ORAL

RENAGEL

+	GENZYME	400MG	N021179 001	Jul 12, 2000
+!		800MG	N021179 002	Jul 12, 2000

SEVOFLURANE

LIQUID;INHALATION

SEVOFLURANE

<u>AN</u>	BAXTER HLTHCARE	<u>100%</u>	<u>A075895 001</u>	Jul 02, 2002
<u>AN</u>	HALOCARBON PRODS	<u>100%</u>	<u>A078650 001</u>	Nov 19, 2007
<u>AN</u>	SHANGHAI HENGRI	<u>100%</u>	<u>A203793 001</u>	Nov 03, 2015
	<u>SOJOURN</u>			
<u>AN</u>	PIRAMAL CRITICAL	<u>100%</u>	<u>A077867 001</u>	May 02, 2007
	<u>ULTANE</u>			
<u>AN</u>	+! ABBVIE	<u>100%</u>	<u>N020478 001</u>	Jun 07, 1995

SILDENAFIL CITRATE

FOR SUSPENSION;ORAL

REVATIO

+!	PFIZER	EQ 10MG BASE/ML	N203109 001	Aug 30, 2012
----	--------	-----------------	-------------	--------------

SOLUTION;INTRAVENOUS

REVATIO

<u>AP</u>	+! PFIZER	<u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u>	<u>N022473 001</u>	Nov 18, 2009
-----------	-----------	---	--------------------	--------------

SILDENAFIL CITRATE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 10MG BASE/12.5ML (EQ 0.8MG BASE/ML)</u>	<u>A203988 001</u>	Apr 01, 2015
-----------	----------------------	---	--------------------	--------------

TABLET;ORAL

REVATIO

<u>AB</u>	+! PFIZER	<u>EQ 20MG BASE</u>	<u>N021845 001</u>	Jun 03, 2005
-----------	-----------	---------------------	--------------------	--------------

SILDENAFIL CITRATE

<u>AB</u>	AMNEAL PHARMS	<u>EQ 20MG BASE</u>	<u>A202025 001</u>	Feb 28, 2013
<u>AB</u>	APOTEX CORP	<u>EQ 20MG BASE</u>	<u>A091379 001</u>	Nov 06, 2012
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 20MG BASE</u>	<u>A203963 001</u>	Nov 18, 2015
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 20MG BASE</u>	<u>A202598 001</u>	Nov 06, 2012
<u>AB</u>	HETERO LABS LTD V	<u>EQ 20MG BASE</u>	<u>A203623 001</u>	Nov 26, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A203814 001</u>	Dec 17, 2013
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 20MG BASE</u>	<u>A201150 001</u>	Nov 09, 2012
<u>AB</u>	RUBICON RES PVT LTD	<u>EQ 20MG BASE</u>	<u>A204883 001</u>	Jun 20, 2016
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A077342 001</u>	Mar 09, 2016
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077342 002</u>	Mar 09, 2016
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077342 003</u>	Mar 09, 2016
<u>AB</u>	TEVA PHARMS	<u>EQ 20MG BASE</u>	<u>A078380 001</u>	Jan 07, 2013
<u>AB</u>	TORRENT PHARMS LTD	<u>EQ 20MG BASE</u>	<u>A091479 001</u>	Nov 06, 2012
<u>AB</u>	WATSON LABS INC	<u>EQ 20MG BASE</u>	<u>A202503 001</u>	Nov 06, 2012
	<u>VIAGRA</u>			
<u>AB</u>	+ PFIZER INC	<u>EQ 25MG BASE</u>	<u>N020895 001</u>	Mar 27, 1998
<u>AB</u>	+	<u>EQ 50MG BASE</u>	<u>N020895 002</u>	Mar 27, 1998
<u>AB</u>	+!	<u>EQ 100MG BASE</u>	<u>N020895 003</u>	Mar 27, 1998

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-383 (of 436)

SILODOSIN

CAPSULE;ORAL

RAPAFLO

AB +! ALLERGAN SALES LLC 4MG

AB + 8MG

SILODOSIN

AB SANDOZ INC 4MG

AB 8MG

SILVER SULFADIAZINE

CREAM;TOPICAL

SILVADENE

AB +! KING PHARMS LLC 1%

SSD

AB DR REDDYS LA 1%

THERMAZENE

AB THEPHARMANETWORK LLC 1%

SSD AF

BX DR REDDYS LA 1%

N022206 001 Oct 08, 2008

N022206 002 Oct 08, 2008

A204726 001 Mar 31, 2017

A204726 002 Mar 31, 2017

SIMEPREVIR SODIUM

CAPSULE;ORAL

OLYSIO

+! JANSSEN PRODS

EQ 150MG BASE

N205123 001 Nov 22, 2013

SIMVASTATIN

SUSPENSION;ORAL

FLOLIPID

+ TCG FLUENT PHARMA 20MG/5ML

+! 40MG/5ML

N206679 001 Apr 21, 2016

N206679 002 Apr 21, 2016

TABLET;ORAL

SIMVASTATIN

AB ACCORD HLTHCARE 5MG

A078155 005 Apr 05, 2013

AB 10MG

A078155 002 Feb 26, 2008

AB 20MG

A078155 003 Feb 26, 2008

AB 40MG

A078155 004 Feb 26, 2008

AB 80MG

A078155 001 Feb 26, 2008

AB AUROBINDO PHARMA 5MG

A077691 001 Dec 20, 2006

AB 10MG

A077691 002 Dec 20, 2006

AB 20MG

A077691 003 Dec 20, 2006

AB 40MG

A077691 004 Dec 20, 2006

AB 80MG

A077691 005 Dec 20, 2006

AB BIOCON LIMITED 5MG

A078034 001 Dec 20, 2006

AB 10MG

A078034 002 Dec 20, 2006

AB 20MG

A078034 003 Dec 20, 2006

AB 40MG

A078034 004 Dec 20, 2006

AB 80MG

A078034 005 Dec 20, 2006

AB DR REDDYS LABS INC 5MG

A077752 005 Jan 23, 2008

AB 10MG

A077752 001 Dec 20, 2006

AB 20MG

A077752 002 Dec 20, 2006

AB 40MG

A077752 003 Dec 20, 2006

AB 80MG

A077752 004 Dec 20, 2006

AB HETERO LABS LTD III 5MG

A200895 001 Nov 25, 2014

AB 10MG

A200895 002 Nov 25, 2014

AB 20MG

A200895 003 Nov 25, 2014

AB 40MG

A200895 004 Nov 25, 2014

AB 80MG

A200895 005 Nov 25, 2014

AB HISUN PHARM HANGZHOU 10MG

A206557 001 Nov 13, 2017

AB 20MG

A206557 002 Nov 13, 2017

AB 40MG

A206557 003 Nov 13, 2017

AB 80MG

A206557 004 Nov 13, 2017

AB IVAX SUB TEVA PHARMS 5MG

A076052 001 Jun 23, 2006

AB 10MG

A076052 002 Jun 23, 2006

AB 20MG

A076052 003 Jun 23, 2006

AB 40MG

A076052 004 Jun 23, 2006

AB 80MG

A076052 005 Dec 20, 2006

AB LUPIN 5MG

A078103 005 Apr 14, 2009

AB 10MG

A078103 001 May 11, 2007

AB 20MG

A078103 002 May 11, 2007

AB 40MG

A078103 003 May 11, 2007

AB 80MG

A078103 004 May 11, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-384 (of 436)

SIMVASTATIN

TABLET;ORAL

**SIMVASTATIN**

<b>AB</b>	OXFORD PHARMS	<u>5MG</u>	<b>A078735 001</b>	Aug 30, 2010
<b>AB</b>		<u>10MG</u>	<b>A078735 002</b>	Aug 30, 2010
<b>AB</b>		<u>20MG</u>	<b>A078735 003</b>	Aug 30, 2010
<b>AB</b>		<u>40MG</u>	<b>A078735 004</b>	Aug 30, 2010
<b>AB</b>		<u>80MG</u>	<b>A078735 005</b>	Aug 30, 2010
<b>AB</b>	VIVA HLTHCARE	<u>5MG</u>	<b>A090383 001</b>	Sep 16, 2011
<b>AB</b>		<u>10MG</u>	<b>A090383 002</b>	Sep 16, 2011
<b>AB</b>		<u>20MG</u>	<b>A090383 003</b>	Sep 16, 2011
<b>AB</b>		<u>40MG</u>	<b>A090383 004</b>	Sep 16, 2011
<b>AB</b>		<u>80MG</u>	<b>A090383 005</b>	Sep 16, 2011
<b>AB</b>	WATSON LABS TEVA	<u>5MG</u>	<b>A076685 001</b>	Dec 20, 2006
<b>AB</b>		<u>10MG</u>	<b>A076685 002</b>	Dec 20, 2006
<b>AB</b>		<u>20MG</u>	<b>A076685 003</b>	Dec 20, 2006
<b>AB</b>		<u>40MG</u>	<b>A076685 004</b>	Dec 20, 2006
<b>AB</b>		<u>80MG</u>	<b>A076685 005</b>	Dec 20, 2006
<b>AB</b>	ZYDUS PHARMS USA	<u>5MG</u>	<b>A077837 001</b>	Dec 20, 2006
<b>AB</b>		<u>10MG</u>	<b>A077837 002</b>	Dec 20, 2006
<b>AB</b>		<u>20MG</u>	<b>A077837 003</b>	Dec 20, 2006
<b>AB</b>		<u>40MG</u>	<b>A077837 004</b>	Dec 20, 2006
<b>AB</b>		<u>80MG</u>	<b>A077837 005</b>	Dec 20, 2006
<b>ZOCOR</b>				
<b>AB</b>	+ MSD MERCK CO	<u>5MG</u>	<b>N019766 001</b>	Dec 23, 1991
<b>AB</b>	+	<u>10MG</u>	<b>N019766 002</b>	Dec 23, 1991
<b>AB</b>	+	<u>20MG</u>	<b>N019766 003</b>	Dec 23, 1991
<b>AB</b>	+	<u>40MG</u>	<b>N019766 004</b>	Dec 23, 1991
<b>AB</b>	++!	<u>80MG</u>	<b>N019766 005</b>	Jul 10, 1998

SINCALIDE

INJECTABLE;INJECTION

KINEVAC

+! BRACCO

0.005MG/VIAL

N017697 001

SINECATECHINS

OINTMENT;TOPICAL

VEREGEN

+! MEDIGENE AG

15%

N021902 001 Oct 31, 2006

SIROLIMUS

SOLUTION;ORAL

RAPAMUNE

+! PF PRISM CV

1MG/ML

N021083 001 Sep 15, 1999

TABLET;ORAL

**RAPAMUNE**

<b>AB</b>	+	PF PRISM CV	<u>0 .5MG</u>	<b>N021110 004</b>	Jan 25, 2010
<b>AB</b>	+		<u>1MG</u>	<b>N021110 001</b>	Aug 25, 2000
<b>AB</b>	++!		<u>2MG</u>	<b>N021110 002</b>	Aug 22, 2002
<b>SIROLIMUS</b>					
<b>AB</b>	DR REDDYS LABS LTD	<u>1MG</u>	<b>A201578 001</b>	Oct 27, 2014	
<b>AB</b>		<u>2MG</u>	<b>A201578 002</b>	Oct 27, 2014	
<b>AB</b>	ZYDUS PHARMS USA INC	<u>0 .5MG</u>	<b>A201676 003</b>	Jan 08, 2014	

SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JANUVIA

+ MERCK SHARP DOHME

EQ 25MG BASE

N021995 001 Oct 16, 2006

+

EQ 50MG BASE

N021995 002 Oct 16, 2006

+!

EQ 100MG BASE

N021995 003 Oct 16, 2006

SODIUM ACETATE

INJECTABLE;INJECTION

SODIUM ACETATE

FRESENIUS KABI USA

4MEQ/ML

A206687 001 Oct 30, 2017

+! HOSPIRA

2MEQ/ML

N018893 001 May 04, 1983

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION;IV (INFUSION)

**AMMONUL**

<b>AP</b>	++!	MEDICIS	<u>10%:10% (5GM/50ML:5GM/50ML)</u>	<b>N020645 001</b>	Feb 17, 2005
<b>SODIUM PHENYLACETATE AND SODIUM BENZOATE</b>					
<b>AP</b>	AILEX PHARMS LLC	<u>10%:10% (5GM/50ML:5GM/50ML)</u>	<b>A207096 001</b>	Feb 24, 2016	
<b>AP</b>	MAIA PHARMS INC	<u>10%:10% (5GM/50ML:5GM/50ML)</u>	<b>A208521 001</b>	May 08, 2017	
<b>AP</b>	NAVINTA LLC	<u>10%:10% (5GM/50ML:5GM/50ML)</u>	<b>A205880 001</b>	Aug 04, 2016	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-385 (of 436)

SODIUM BICARBONATE

INJECTABLE; INJECTION

SODIUM BICARBONATE

<u>AP</u>	!	HOSPIRA	<u>0.9MEQ/ML</u>	<u>A077394</u> <u>001</u>	Nov 09, 2005
<u>AP</u>	!		<u>1MEQ/ML</u>	<u>A077394</u> <u>002</u>	Nov 09, 2005
<u>AP</u>		HOSPIRA INC	<u>0.9MEQ/ML</u>	<u>A202494</u> <u>001</u>	Mar 06, 2017
<u>AP</u>			<u>1MEQ/ML</u>	<u>A202432</u> <u>001</u>	Sep 26, 2017
<u>AP</u>			<u>1MEQ/ML</u>	<u>A202494</u> <u>002</u>	Mar 06, 2017
<u>AP</u>		INTL MEDICATION SYS	<u>1MEQ/ML</u>	<u>A203449</u> <u>001</u>	Sep 19, 2017
		HOSPIRA INC	0.5MEQ/ML	A202679 001	Mar 07, 2017
			0.5MEQ/ML	A202981 001	Mar 04, 2016
			1MEQ/ML	A202495 001	Mar 06, 2017

SODIUM CHLORIDE

INJECTABLE; INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>		FRESENIUS KABI USA	<u>9MG/ML</u>	<u>A088911</u> <u>001</u>	Feb 07, 1985
<u>AP</u>	+	HOSPIRA	<u>9MG/ML</u>	<u>N018800</u> <u>001</u>	Oct 29, 1982
<u>SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	B BRAUN	<u>450MG/100ML</u>	<u>N019635</u> <u>001</u>	Mar 09, 1988
<u>AP</u>		BAXTER HLTHCARE	<u>450MG/100ML</u>	<u>N018016</u> <u>001</u>	
<u>AP</u>		HOSPIRA	<u>450MG/100ML</u>	<u>N019759</u> <u>001</u>	Jun 08, 1988
<u>AP</u>	+	ICU MEDICAL INC	<u>450MG/100ML</u>	<u>N018090</u> <u>001</u>	
<u>SODIUM CHLORIDE 0.9%</u>					
<u>AP</u>		SPECTRA MDCL	<u>9MG/ML</u>	<u>A206171</u> <u>001</u>	Jul 21, 2017
		DEVICES			
<u>AP</u>		WEST-WARD PHARMS	<u>9MG/ML</u>	<u>A201850</u> <u>001</u>	Jan 20, 2012
		INT			
<u>SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	B BRAUN	<u>900MG/100ML</u>	<u>N017464</u> <u>001</u>	
<u>AP</u>	+		<u>900MG/100ML</u>	<u>N019635</u> <u>002</u>	Mar 09, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>9MG/ML</u>	<u>N016677</u> <u>004</u>	Oct 30, 1985
<u>AP</u>	+		<u>9MG/ML</u>	<u>N020178</u> <u>002</u>	Dec 07, 1992
<u>AP</u>	+		<u>900MG/100ML</u>	<u>N016677</u> <u>001</u>	
<u>AP</u>	+		<u>900MG/100ML</u>	<u>N020178</u> <u>001</u>	Dec 07, 1992
<u>AP</u>	!	FRESENIUS KABI USA	<u>9MG/ML</u>	<u>A088912</u> <u>001</u>	Jan 10, 1985
<u>AP</u>			<u>900MG/100ML</u>	<u>A207310</u> <u>001</u>	Sep 19, 2017
<u>AP</u>		FRESENIUS MEDCL	<u>900MG/100ML</u>	<u>A078177</u> <u>001</u>	Apr 12, 2007
<u>AP</u>		HAEMONETICS	<u>900MG/100ML</u>	<u>A076316</u> <u>001</u>	Oct 27, 2004
<u>AP</u>	+	HOSPIRA	<u>9MG/ML</u>	<u>N018803</u> <u>001</u>	Oct 29, 1982
<u>AP</u>	+		<u>9MG/ML</u>	<u>N019217</u> <u>001</u>	Jul 13, 1984
<u>AP</u>	+		<u>9MG/ML</u>	<u>N019465</u> <u>002</u>	Jul 15, 1985
<u>AP</u>	+		<u>900MG/100ML</u>	<u>N019465</u> <u>001</u>	Jul 15, 1985
<u>AP</u>	+		<u>900MG/100ML</u>	<u>N019480</u> <u>001</u>	Sep 17, 1985
<u>AP</u>	+	ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N016366</u> <u>001</u>	
<u>AP</u>		JUBILANT	<u>9MG/ML</u>	<u>A203352</u> <u>001</u>	May 18, 2016
<u>AP</u>		HOLLISTRSTR			
<u>AP</u>		LABORATORIOS	<u>900MG/100ML</u>	<u>A207956</u> <u>001</u>	May 25, 2017
<u>AP</u>		GRIFOLS			
<u>AP</u>	!	TARO	<u>9MG/ML</u>	<u>A077407</u> <u>001</u>	Aug 11, 2006
<u>SODIUM CHLORIDE 3% IN PLASTIC CONTAINER</u>					
<u>AP</u>		B BRAUN	<u>3GM/100ML</u>	<u>N019635</u> <u>003</u>	Mar 09, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>3GM/100ML</u>	<u>N019022</u> <u>001</u>	Nov 01, 1983
<u>SODIUM CHLORIDE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>		B BRAUN	<u>5GM/100ML</u>	<u>N019635</u> <u>004</u>	Mar 09, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N019022</u> <u>002</u>	Nov 01, 1983
SODIUM CHLORIDE 0.9%					
	+	B BRAUN	900MG/100ML	N019635 005	Aug 11, 2016
		MEDEFIL INC	90MG/10ML (9MG/ML)	N202832 006	Jan 06, 2012
SODIUM CHLORIDE 0.9%					
		WEST-WARD PHARMS	9MG/ML	A201833 001	Sep 24, 2013
		INT			
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER					
	+	LIEBEL-FLARSHEIM	405MG/50ML (9MG/ML)	N021569 001	Jul 27, 2006
	+		1012.5MG/125ML (9MG/ML)	N021569 002	Jul 27, 2006
SODIUM CHLORIDE IN PLASTIC CONTAINER					
	+	HOSPIRA	2.5MEQ/ML	N018897 001	Jul 20, 1984
SOLUTION; IRRIGATION					
<u>SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>					
<u>AT</u>		B BRAUN	<u>900MG/100ML</u>	<u>N016733</u> <u>001</u>	
<u>AT</u>		BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N017427</u> <u>001</u>	
<u>AT</u>			<u>900MG/100ML</u>	<u>N017867</u> <u>001</u>	
<u>AT</u>		ICU MEDICAL INC	<u>900MG/100ML</u>	<u>N017514</u> <u>001</u>	
<u>AT</u>			<u>900MG/100ML</u>	<u>N018314</u> <u>001</u>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-386 (of 436)

SODIUM CHLORIDE

SOLUTION FOR SLUSH;IRRIGATION  
 SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER  
 BAXTER HLTHCARE 900MG/100ML

N019319 002 May 17, 1985

SODIUM FERRIC GLUCONATE COMPLEX

INJECTABLE;INJECTION

**FERRLECIT**

**AB** +! SANOFI AVENTIS US **62.5MG/5ML**  
**SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE**  
**AB** WEST-WARD PHARMS **62.5MG/5ML**  
 INT

**N020955 001** Feb 18, 1999  
**A078215 001** Mar 31, 2011

SODIUM FLUORIDE F-18

INJECTABLE;INTRAVENOUS

**SODIUM FLUORIDE F-18**

<b>AP</b>	3D IMAGING DRUG	<b>10-200mCi/ML</b>	<b>A203777 001</b>	Oct 19, 2015
<b>AP</b>	BIOMEDCL RES FDN	<b>10-200mCi/ML</b>	<b>A204351 001</b>	Jan 09, 2015
<b>AP</b>	CARDINAL HEALTH 414	<b>10-200mCi/ML</b>	<b>A203780 001</b>	Jul 30, 2015
<b>AP</b>	ESSENTIAL ISOTOPES	<b>10-200mCi/ML</b>	<b>A204541 001</b>	Oct 29, 2014
<b>AP</b>	GLOBAL ISOTOPES LLC	<b>10-200mCi/ML</b>	<b>A204464 001</b>	Oct 21, 2014
<b>AP</b>	HOT SHOTS NM LLC	<b>10-200mCi/ML</b>	<b>A204530 001</b>	Jul 29, 2015
<b>AP</b> !	HOUSTON CYCLOTRON	<b>10-200mCi/ML</b>	<b>A203544 001</b>	Dec 26, 2012
<b>AP</b>	JUBILANT DRAXIMAGE	<b>10-200mCi/ML</b>	<b>A203968 001</b>	Oct 23, 2015
<b>AP</b>	KREITCHMAN PET CTR	<b>10-200mCi/ML</b>	<b>A203936 001</b>	May 19, 2016
<b>AP</b>	MIDWEST MEDCL	<b>10-200mCi/ML</b>	<b>A204440 001</b>	Nov 17, 2015
<b>AP</b>	MIPS CRF	<b>10-200mCi/ML</b>	<b>A204517 001</b>	Jul 21, 2015
<b>AP</b>	NCM USA BRONX LLC	<b>10-200mCi/ML</b>	<b>A204513 001</b>	Nov 28, 2014
<b>AP</b>	PETNET	<b>10-200mCi/ML</b>	<b>A203890 001</b>	Sep 28, 2015
<b>AP</b>	PRECISION NUCLEAR	<b>10-200mCi/ML</b>	<b>A204542 001</b>	Feb 27, 2015
<b>AP</b>	SHERTECH LABS LLC	<b>10-200mCi/ML</b>	<b>A204315 001</b>	Sep 22, 2014
<b>AP</b>	SPECTRON MRC LLC	<b>10-200mCi/ML</b>	<b>A203912 001</b>	Apr 22, 2015
<b>AP</b>	UCSF RODIOPHARM	<b>10-200mCi/ML</b>	<b>A204437 001</b>	Mar 13, 2014
<b>AP</b>	UNIV TX MD ANDERSON	<b>10-200mCi/ML</b>	<b>A203247 001</b>	Dec 23, 2013
<b>AP</b>	UNIV UTAH CYCLOTRON	<b>10-200mCi/ML</b>	<b>A204497 001</b>	Apr 20, 2015
<b>AP</b>	ZEVACOR PHARMA INC	<b>10-200mCi/ML</b>	<b>A203592 001</b>	Aug 18, 2015
!	MCPRF	10-91.5mCi/ML	A203605 001	Jun 28, 2013
	THE FEINSTEIN INST	20-600mCi/ML	A204328 001	Nov 19, 2014

SODIUM IODIDE I-123

CAPSULE;ORAL

**SODIUM IODIDE I 123**

<b>AA</b> +!	CARDINAL HEALTH 418	<b>100uCi</b>	<b>N018671 001</b>	May 27, 1982
<b>AA</b> +!		<b>200uCi</b>	<b>N018671 002</b>	May 27, 1982
<b>AA</b>	MALLINKRODT NUCLEAR	<b>100uCi</b>	<b>A071909 001</b>	Feb 28, 1989
<b>AA</b>		<b>200uCi</b>	<b>A071910 001</b>	Feb 28, 1989

SODIUM IODIDE I-131

CAPSULE;ORAL

SODIUM IODIDE I 131  
 + JUBILANT DRAXIMAGE 0.009-0.1mCi  
 +! MALLINKRODT NUCLEAR 0.8-100mCi

N021305 006 May 19, 2005  
 N016517 001

SOLUTION;ORAL

HICON  
 +! JUBILANT DRAXIMAGE 250-1000mCi  
 SODIUM IODIDE I 131  
 +! MALLINKRODT NUCLEAR 3.5-150mCi/VIAL

N021305 007 Dec 05, 2011  
 N016515 001

SODIUM LACTATE

INJECTABLE;INJECTION

SODIUM LACTATE IN PLASTIC CONTAINER  
 +! HOSPIRA 5MEQ/ML

N018947 001 Sep 05, 1984

SODIUM NITRITE

SOLUTION;INTRAVENOUS  
 SODIUM NITRITE  
 +! HOPE PHARMS 300MG/10ML (30MG/ML)

N203922 001 Feb 14, 2012

SODIUM NITRITE; SODIUM THIOSULFATE

SOLUTION, SOLUTION;INTRAVENOUS, INTRAVENOUS

NITHIODOTE

+! HOPE PHARMS 300MG/10ML (30MG/ML), N/A;N/A,12.5GM/50ML  
 (250MG/ML)

N201444 001 Jan 14, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-387 (of 436)

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

**NITROPRESS**

<b>AP</b>	! HOSPIRA	<u>25MG/ML</u>	<b>A071961 001</b>	Aug 01, 1988
<b>SODIUM NITROPRUSSIDE</b>				
<b>AP</b>	AKORN	<u>25MG/ML</u>	<b>A208635 001</b>	May 04, 2017
<b>AP</b>	AMNEAL PHARMS CO	<u>25MG/ML</u>	<b>A209493 001</b>	Nov 07, 2017
<b>AP</b>	AMPHASSTAR PHARMS INC	<u>25MG/ML</u>	<b>A209832 001</b>	Dec 18, 2017
<b>AP</b>	MICRO LABS	<u>25MG/ML</u>	<b>A209352 001</b>	Dec 08, 2017
<b>AP</b>	NAMIGEN LLC	<u>25MG/ML</u>	<b>A207426 001</b>	Dec 08, 2016
<b>AP</b>	NEXUS PHARMS	<u>25MG/ML</u>	<b>A207499 001</b>	May 25, 2017

SOLUTION; IV (INFUSION)

NIPRIDE RTU IN SODIUM CHLORIDE 0.9%

+!	EXELA PHARMA SCS LLC	10MG/50ML (0.2MG/ML)	N209387 002	Dec 07, 2017
+!		50MG/100ML (0.5MG/ML)	N209387 001	Mar 08, 2017

SODIUM OXYBATE

SOLUTION; ORAL

**SODIUM OXYBATE**

<b>AA</b>	WEST-WARD PHARMS INT	<u>500MG/ML</u>	<b>A202090 001</b>	Jan 17, 2017
-----------	----------------------	-----------------	--------------------	--------------

**XYREM**

<b>AA</b>	+! JAZZ PHARMS	<u>500MG/ML</u>	<b>N021196 001</b>	Jul 17, 2002
-----------	----------------	-----------------	--------------------	--------------

SODIUM PHENYLBUTYRATE

POWDER; ORAL

**BUPHENYL**

<b>AB</b>	+! HORIZON PHARMA INC	<u>3GM/TEASPOONFUL</u>	<b>N020573 001</b>	Apr 30, 1996
-----------	-----------------------	------------------------	--------------------	--------------

**SODIUM PHENYLBUTYRATE**

<b>AB</b>	PAR PHARM	<u>3GM/TEASPOONFUL</u>	<b>A202819 001</b>	Mar 22, 2013
TABLET; ORAL				

**BUPHENYL**

<b>AB</b>	+! HORIZON PHARMA INC	<u>500MG</u>	<b>N020572 001</b>	May 13, 1996
-----------	-----------------------	--------------	--------------------	--------------

**SODIUM PHENYLBUTYRATE**

<b>AB</b>	ALVOGEN MALTA	<u>500MG</u>	<b>A090910 001</b>	Nov 18, 2011
<b>AB</b>	PAR PHARM	<u>500MG</u>	<b>A204395 001</b>	Apr 15, 2016

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

OSMOPREP

+!	SALIX PHARMS	0.398GM;1.102GM	N021892 001	Mar 16, 2006
----	--------------	-----------------	-------------	--------------

SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

SODIUM PHOSPHATES IN PLASTIC CONTAINER

+!	HOSPIRA	142MG/ML;276MG/ML	N018892 001	May 10, 1983
----	---------	-------------------	-------------	--------------

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

**KALEXATE**

<b>AA</b>	! KVK TECH	<u>454GM/BOT</u>	<b>A040905 001</b>	Mar 30, 2009
-----------	------------	------------------	--------------------	--------------

**KIONEX**

<b>AA</b>	PADDOCK LLC	<u>454GM/BOT</u>	<b>A040029 001</b>	Feb 06, 1998
-----------	-------------	------------------	--------------------	--------------

**SODIUM POLYSTYRENE SULFONATE**

<b>AA</b>	AILEX PHARMS LLC	<u>454GM/BOT</u>	<b>A206815 001</b>	Feb 18, 2016
<b>AA</b>	BELCHER PHARMS LLC	<u>454GM/BOT</u>	<b>A205727 001</b>	Feb 23, 2016
<b>AA</b>	CMP PHARMA INC	<u>454GM/BOT</u>	<b>A089910 001</b>	Jan 19, 1989
<b>AA</b>	ECI PHARMS LLC	<u>453.6GM/BOT</u>	<b>A090313 001</b>	Dec 21, 2011
<b>AA</b>	EPIC PHARMA LLC	<u>453.6GM/BOT</u>	<b>A202333 001</b>	Mar 19, 2014

**NUVO PHARM INC**

**454GM/BOT**

**A204071 001** Nov 28, 2014

KALEXATE

KVK TECH

A040905 002 Apr 03, 2015

SODIUM POLYSTYRENE SULFONATE

NUVO PHARM INC

A204071 002 Nov 28, 2014

SUSPENSION; ORAL, RECTAL

**KIONEX**

<b>AA</b>	PADDOCK LLC	<u>15GM/60ML</u>	<b>A040028 001</b>	Sep 17, 2007
-----------	-------------	------------------	--------------------	--------------

**SODIUM POLYSTYRENE SULFONATE**

PADDOCK LLC

**A090590 001** May 13, 2011

<b>AA</b>	WEST-WARD PHARMS INT	<u>15GM/60ML</u>	<b>A089049 001</b>	Nov 17, 1986
-----------	----------------------	------------------	--------------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-388 (of 436)

SODIUM POLYSTYRENE SULFONATE

SUSPENSION;ORAL, RECTAL

**SPS**

**AA ! CMP PHARMA INC 15GM/60ML**

**A087859 001 Dec 08, 1982**

SODIUM TETRADECYL SULFATE

INJECTABLE;INJECTION

SOTRADECOL

! MYLAN INSTITUTIONAL 20MG/2ML (10MG/ML)  
! 60MG/2ML (30MG/ML)

A040541 001 Nov 12, 2004  
A040541 002 Nov 12, 2004

SODIUM THIOSULFATE

SOLUTION;INTRAVENOUS

SODIUM THIOSULFATE

+! HOPE PHARMS 12.5GM/50ML (250MG/ML)

N203923 001 Feb 14, 2012

SOFOSBUVIR

TABLET;ORAL

SOVALDI

+! GILEAD SCIENCES INC 400MG

N204671 001 Dec 06, 2013

SOFOSBUVIR; VELPATASVIR

TABLET;ORAL

EPCLUSA

+! GILEAD SCIENCES INC 400MG;100MG

N208341 001 Jun 28, 2016

SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR

TABLET;ORAL

VOSEVI

+! GILEAD SCIENCES INC 400MG;100MG;100MG

N209195 001 Jul 18, 2017

SOLIFENACIN SUCCINATE

TABLET;ORAL

**SOLIFENACIN SUCCINATE**

**AB TEVA PHARMS USA 5MG**

**A091464 001 Apr 02, 2014**

**AB 10MG**

**A091464 002 Apr 02, 2014**

**VESICARE**

**AB + ASTELLAS 5MG**

**N021518 001 Nov 19, 2004**

**AB +! 10MG**

**N021518 002 Nov 19, 2004**

SOMATROPIN RECOMBINANT

INJECTABLE;INJECTION

GENOTROPIN

BX +! PHARMACIA AND 5.8MG/VIAL  
UPJOHN

N020280 006 Aug 24, 1995

GENOTROPIN PRESERVATIVE FREE

BX + PHARMACIA AND 1.5MG/VIAL  
UPJOHN

N020280 004 Aug 24, 1995

HUMATROPE

BX +! LILLY 5MG/VIAL

N019640 004 Mar 08, 1987

BX + 6MG/VIAL

N019640 005 Feb 04, 1999

NORDITROPIN FLEXPRO

BX NOVO NORDISK INC 5MG/1.5ML

N021148 008 Mar 01, 2010

BX 10MG/1.5ML

N021148 009 Mar 01, 2010

OMNITROPE

BX SANDOZ 1.5MG/VIAL

N021426 002 May 30, 2006

BX 5MG/1.5ML

N021426 003 Jan 16, 2008

BX 5.8MG/VIAL

N021426 001 May 30, 2006

BX 10MG/1.5ML

N021426 004 Aug 25, 2008

SAIZEN

BX + EMD SERONO 5MG/VIAL

N019764 002 Oct 08, 1996

SEROSTIM

BX EMD SERONO 5MG/VIAL

N020604 002 Aug 23, 1996

BX 6MG/VIAL

N020604 001 Aug 23, 1996

VALTROPIN

BX LG CHEM LTD 5MG/VIAL

N021905 001 Apr 19, 2007

ZOMACTON

BX +! FERRING 5MG/VIAL

N019774 002 Jan 04, 2002

GENOTROPIN

+! PHARMACIA AND 13.8MG/VIAL

N020280 007 Oct 23, 1996

GENOTROPIN PRESERVATIVE FREE

+ PHARMACIA AND 0.2MG/VIAL

N020280 001 Jan 27, 1998

+ UPJOHN 0.4MG/VIAL

N020280 002 Jan 27, 1998

+ 0.6MG/VIAL

N020280 003 Jan 27, 1998

+ 0.8MG/VIAL

N020280 005 Jan 27, 1998

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-389 (of 436)

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

GENOTROPIN PRESERVATIVE FREE

+ + + + + + !	1MG/VIAL 1.2MG/VIAL 1.4MG/VIAL 1.6MG/VIAL 1.8MG/VIAL 2MG/VIAL	N020280 008 N020280 009 N020280 010 N020280 011 N020280 012 N020280 013	Jan 27, 1998 Jan 27, 1998 Jan 27, 1998 Jan 27, 1998 Jan 27, 1998 Jan 27, 1998
HUMATROPE			
+! LILLY	12MG/VIAL	N019640 006	Feb 04, 1999
+!	24MG/VIAL	N019640 007	Feb 04, 1999
NORDITROPIN FLEXPRO			
NOVO NORDISK INC	15MG/1.5ML 30MG/3ML	N021148 010 N021148 011	Mar 01, 2010 Jan 23, 2015
NUTROPIN AQ NUSPIN			
+! GENENTECH	5MG/2ML (2.5MG/ML)	N020522 003	Jan 03, 2008
+!	10MG/2ML (5MG/ML)	N020522 005	Jan 03, 2008
+!	20MG/2ML (10MG/ML)	N020522 004	Jan 03, 2008
NUTROPIN AQ PEN			
+! GENENTECH	10MG/2ML (5MG/ML)	N020522 002	Apr 22, 2002
+!	20MG/2ML (10MG/ML)	N020522 006	Jan 03, 2008
SAIZEN			
+! EMD SERONO	8.8MG/VIAL	N019764 003	Aug 29, 2000
SEROSTIM			
EMD SERONO	4MG/VIAL	N020604 003	Jul 25, 1997
ZOMACTON			
+! FERRING	10MG/VIAL	N019774 003	Mar 07, 2012
ZORBTIVE			
+! EMD SERONO	8.8MG/VIAL	N021597 004	Dec 01, 2003

SONIDEGIB PHOSPHATE

CAPSULE; ORAL

ODOMZO

+! SUN PHARMA GLOBAL	EQ 200MG BASE	N205266 001	Jul 24, 2015
----------------------	---------------	-------------	--------------

SORAFENIB TOSYLATE

TABLET; ORAL

NEXAVAR

+! BAYER HLTHCARE	EQ 200MG BASE	N021923 001	Dec 20, 2005
-------------------	---------------	-------------	--------------

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE	3GM/100ML	N017863 001
-----------------	-----------	-------------

SORBITOL 3.3% IN PLASTIC CONTAINER

B BRAUN	3.3GM/100ML	N016741 001
---------	-------------	-------------

SOTALOL HYDROCHLORIDE

SOLUTION; INTRAVENOUS

SOTALOL HYDROCHLORIDE

+! ALTATHERA PHARMS LLC	150MG/10ML (15MG/ML)	N022306 001	Jul 02, 2009
-------------------------	----------------------	-------------	--------------

SOLUTION; ORAL

SOTYLIZE

+! ARBOR PHARMS LLC	5MG/ML (5MG/ML)	N205108 001	Oct 22, 2014
---------------------	-----------------	-------------	--------------

TABLET; ORAL

BETAPACE

<b>AB1</b> + COVIS PHARMA BV	<b>80MG</b>
<b>AB1</b> +	<b>120MG</b>
<b>AB1</b> +!	<b>160MG</b>
<b>AB1</b> +	<b>240MG</b>

**N019865 001** Oct 30, 1992

**N019865 005** Apr 20, 1994

**N019865 002** Oct 30, 1992

**N019865 003** Oct 30, 1992

SORINE

<b>AB1</b> UPSHER-SMITH LABS	<b>80MG</b>
<b>AB1</b>	<b>120MG</b>
<b>AB1</b>	<b>160MG</b>
<b>AB1</b>	<b>240MG</b>

**A075500 001** Apr 27, 2001

**A075500 004** Apr 27, 2001

**A075500 002** Apr 27, 2001

**A075500 003** Apr 27, 2001

SOTALOL HYDROCHLORIDE

<b>AB1</b> APOTEX INC	<b>80MG</b>
<b>AB1</b>	<b>120MG</b>
<b>AB1</b>	<b>160MG</b>
<b>AB1</b>	<b>240MG</b>
<b>AB1</b> BEXIMCO PHARMS USA	<b>80MG</b>
<b>AB1</b>	<b>120MG</b>
<b>AB1</b>	<b>160MG</b>

**A076140 001** Sep 26, 2002

**A076140 002** Sep 26, 2002

**A076140 003** Sep 26, 2002

**A076140 004** Sep 26, 2002

**A207428 001** Oct 21, 2016

**A207428 002** Oct 21, 2016

**A207428 003** Oct 21, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PREScription DRUG PRODUCT LIST**

3-390 (of 436)

## SOTALOL HYDROCHLORIDE

**TABLET; ORAL**

## SOTALOL HYDROCHLORIDE

<b>AB1</b>	TEVA	<b>80MG</b>	<b>A075429_001</b>	May 01, 2000
<b>AB1</b>		<b><u>120MG</u></b>	<b>A075429_002</b>	May 01, 2000
<b>AB1</b>		<b><u>160MG</u></b>	<b>A075429_003</b>	May 01, 2000
<b>AB1</b>		<b><u>240MG</u></b>	<b>A075429_004</b>	May 01, 2000
<b>AB1</b>	UPSHER-SMITH LABS	<b>80MG</b>	<b>A075366_001</b>	May 01, 2000
<b>AB1</b>		<b><u>120MG</u></b>	<b>A075366_002</b>	May 01, 2000
<b>AB1</b>		<b><u>160MG</u></b>	<b>A075366_003</b>	May 01, 2000
<b>AB1</b>		<b><u>240MG</u></b>	<b>A075366_004</b>	May 01, 2000
<b>AB1</b>	VINTAGE PHARMS	<b>80MG</b>	<b>A075563_001</b>	Nov 07, 2003
<b>AB1</b>		<b><u>120MG</u></b>	<b>A075563_002</b>	Nov 07, 2003
<b>AB1</b>		<b><u>160MG</u></b>	<b>A075563_003</b>	Nov 07, 2003
<b>AB1</b>		<b><u>240MG</u></b>	<b>A075563_004</b>	Nov 07, 2003

## BETAPACE AF

**AB2** + COVIS PHARMA BV **80MG** **N021151 001** Feb 22, 2000  
**AB2** + **120MG** **N021151 002** Feb 22, 2000  
**AB2** +! **160MG** **N021151 003** Feb 22, 2000

## SOTALOL HYDROCHLORIDE

<u><b>AB2</b></u>	APOTEX	<u><b>80MG</b></u>	<u><b>A076214 001</b></u>	Aug 27, 2003
<u><b>AB2</b></u>		<u><b>120MG</b></u>	<u><b>A076214 002</b></u>	Aug 27, 2003
<u><b>AB2</b></u>		<u><b>160MG</b></u>	<u><b>A076214 003</b></u>	Aug 27, 2003
<u><b>AB2</b></u>	EPIC PHARMA INC	<u><b>80MG</b></u>	<u><b>A077070 001</b></u>	Nov 04, 2005
<u><b>AB2</b></u>		<u><b>120MG</b></u>	<u><b>A077070 002</b></u>	Nov 04, 2005
<u><b>AB2</b></u>		<u><b>160MG</b></u>	<u><b>A077070 003</b></u>	Nov 04, 2005
<u><b>AB2</b></u>	MYLAN	<u><b>80MG</b></u>	<u><b>A077616 001</b></u>	Feb 07, 2007
<u><b>AB2</b></u>		<u><b>120MG</b></u>	<u><b>A077616 002</b></u>	Feb 07, 2007
<u><b>AB2</b></u>		<u><b>160MG</b></u>	<u><b>A077616 003</b></u>	Feb 07, 2007

## SOYBEAN OIL

## INJECTABLE; INJECTION

## INTRALIPID 10%

<u>AP</u>	+!	FRESENIUS	<u>10%</u>	<u>N017643</u>	<u>001</u>
		<u>INTRALIPID 20%</u>			
<u>AP</u>	+!	FRESENIUS	<u>20%</u>	<u>N018449</u>	<u>001</u>
<u>AP</u>	+!		<u>20%</u>	<u>N020248</u>	<u>001</u>
		<u>NUTRILIPID 10%</u>			Aug 07, 1996
<u>AP</u>	+!	B BRAUN	<u>10%</u>	<u>N019531</u>	<u>001</u>
		<u>NUTRILIPID 20%</u>			May 28, 1993
<u>AP</u>	+!	B BRAUN	<u>20%</u>	<u>N019531</u>	<u>002</u>
		INTRALIPID 30%			May 28, 1993
	+!	FRESENIUS	30%	N019942	001 Dec 30, 1993

## SPINOSAD

## SUSPENSION; TOPICAL

## NATROBA

+! PARAPRO LLC

0.9%

N022408 001 Jan 18, 2011

## SPIRONOLACTONE

## SUSPENSION; ORAL

CAROSPIR

+! CMP DEV LLC

25MG/5ML

N209478 001 Aug 04, 2017

**TABLET; ORAL**

## ALDACTONE

<u>AB</u>	+	GD SEARLE LLC	<u>25MG</u>	<u>N012151</u>	<u>009</u>	Dec 30, 1983
<u>AB</u>	+		<u>50MG</u>	<u>N012151</u>	<u>008</u>	Dec 30, 1982
<u>AB</u>	+!		<u>100MG</u>	<u>N012151</u>	<u>010</u>	Dec 30, 1983

## SPIRONOLACTONE

<u><b>AB</b></u>	ACCORD HLTHCARE	<u><b>25MG</b></u>	<u><b>A203512</b></u>	<u><b>001</b></u>	Sep 19, 2016
<u><b>AB</b></u>		<u><b>50MG</b></u>	<u><b>A203512</b></u>	<u><b>002</b></u>	Sep 19, 2016
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A203512</b></u>	<u><b>003</b></u>	Sep 19, 2016
<u><b>AB</b></u>	ACTAVIS ELIZABETH	<u><b>25MG</b></u>	<u><b>A040353</b></u>	<u><b>003</b></u>	Mar 15, 2006
<u><b>AB</b></u>		<u><b>50MG</b></u>	<u><b>A040353</b></u>	<u><b>001</b></u>	Jul 29, 1999
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A040353</b></u>	<u><b>002</b></u>	Jul 29, 1999
<u><b>AB</b></u>	AMNEAL PHARMS	<u><b>25MG</b></u>	<u><b>A091426</b></u>	<u><b>001</b></u>	Jul 02, 2010
<u><b>AB</b></u>		<u><b>50MG</b></u>	<u><b>A091426</b></u>	<u><b>002</b></u>	Jul 02, 2010
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A091426</b></u>	<u><b>003</b></u>	Jul 02, 2010
<u><b>AB</b></u>	AUROBINDO PHARMA LTD	<u><b>25MG</b></u>	<u><b>A202187</b></u>	<u><b>001</b></u>	Mar 06, 2014
<u><b>AB</b></u>		<u><b>50MG</b></u>	<u><b>A202187</b></u>	<u><b>002</b></u>	Mar 06, 2014
<u><b>AB</b></u>		<u><b>100MG</b></u>	<u><b>A202187</b></u>	<u><b>003</b></u>	Mar 06, 2014
<u><b>AB</b></u>	JUBILANT GENERICS	<u><b>25MG</b></u>	<u><b>A203253</b></u>	<u><b>001</b></u>	Apr 23, 2014
<u><b>AB</b></u>		<u><b>50MG</b></u>	<u><b>A203253</b></u>	<u><b>002</b></u>	Apr 23, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-391 (of 436)

SPIRONOLACTONE

TABLET;ORAL

SPIRONOLACTONE

<u>AB</u>		<u>100MG</u>	<u>A203253 003</u>	Apr 23, 2014
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A040424 001</u>	Aug 20, 2001
<u>AB</u>		<u>50MG</u>	<u>A040424 002</u>	Aug 20, 2001
<u>AB</u>		<u>100MG</u>	<u>A040424 003</u>	Aug 20, 2001
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>A086809 001</u>	
<u>AB</u>	SUN PHARM INDUSTRIES	<u>25MG</u>	<u>A089424 001</u>	Jul 23, 1986
<u>AB</u>		<u>50MG</u>	<u>A089424 002</u>	Aug 11, 1999
<u>AB</u>		<u>100MG</u>	<u>A089424 003</u>	Aug 11, 1999
<u>AB</u>	VINTAGE	<u>25MG</u>	<u>A040750 001</u>	Aug 29, 2006
<u>AB</u>		<u>50MG</u>	<u>A040750 002</u>	Aug 29, 2006
<u>AB</u>		<u>100MG</u>	<u>A040750 003</u>	Aug 29, 2006

STAVUDINE

CAPSULE;ORAL

STAVUDINE

<u>AB</u>	AUROBINDO PHARMA	<u>15MG</u>	<u>A077672 003</u>	Dec 29, 2008
<u>AB</u>		<u>20MG</u>	<u>A077672 004</u>	Dec 29, 2008
<u>AB</u>		<u>30MG</u>	<u>A077672 001</u>	Dec 29, 2008
<u>AB</u>		<u>40MG</u>	<u>A077672 002</u>	Dec 29, 2008
<u>AB</u>	HETERO LABS LTD III	<u>15MG</u>	<u>A078957 001</u>	Dec 29, 2008
<u>AB</u>		<u>20MG</u>	<u>A078957 002</u>	Dec 29, 2008
<u>AB</u>		<u>30MG</u>	<u>A078957 003</u>	Dec 29, 2008
<u>AB</u>		<u>40MG</u>	<u>A078957 004</u>	Dec 29, 2008
<u>AB</u>	MYLAN	<u>15MG</u>	<u>A079069 001</u>	Dec 29, 2008
<u>AB</u>		<u>20MG</u>	<u>A079069 002</u>	Dec 29, 2008
<u>AB</u>		<u>30MG</u>	<u>A079069 003</u>	Dec 29, 2008
<u>AB</u>		<u>40MG</u>	<u>A079069 004</u>	Dec 29, 2008

ZERIT

<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>15MG</u>	<u>N020412 002</u>	Jun 24, 1994
<u>AB</u>		<u>20MG</u>	<u>N020412 003</u>	Jun 24, 1994
<u>AB</u>	+!	<u>30MG</u>	<u>N020412 004</u>	Jun 24, 1994
<u>AB</u>	+!	<u>40MG</u>	<u>N020412 005</u>	Jun 24, 1994

FOR SOLUTION;ORAL

STAVUDINE

<u>AA</u>	CIPLA LTD	<u>1MG/ML</u>	<u>A078030 001</u>	Mar 20, 2009
<u>AA</u>	<u>ZERIT</u>			
<u>AA</u>	+! BRISTOL MYERS SQUIBB	<u>1MG/ML</u>	<u>N020413 001</u>	Sep 06, 1996

STERILE WATER FOR INJECTION

LIQUID;N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	+! HOSPIRA	<u>100%</u>	<u>N018802 001</u>	Oct 27, 1982
<u>AP</u>	<u>STERILE WATER FOR INJECTION</u>			
<u>AP</u>	FRESENIUS KABI USA	<u>100%</u>	<u>A209689 001</u>	Nov 24, 2017

<u>AP</u>	WEST-WARD PHARMS	<u>100%</u>	<u>A206369 001</u>	Sep 02, 2015
<u>AP</u>	INT			

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	+! B BRAUN	<u>100%</u>	<u>N019633 001</u>	Feb 29, 1988
<u>AP</u>	+! BAXTER HLTHCARE	<u>100%</u>	<u>N018632 001</u>	Jun 30, 1982
<u>AP</u>	+!	<u>100%</u>	<u>N018632 002</u>	Apr 19, 1988
<u>AP</u>	FRESENIUS KABI USA	<u>100%</u>	<u>A088400 001</u>	Jan 16, 1984
<u>AP</u>	+! HOSPIRA	<u>100%</u>	<u>N018801 001</u>	Oct 27, 1982
<u>AP</u>	+! ICU MEDICAL INC	<u>100%</u>	<u>N018233 001</u>	
<u>AP</u>	+!	<u>100%</u>	<u>N019869 001</u>	Dec 26, 1989
<u>AP</u>	TARO	<u>100%</u>	<u>A077393 001</u>	Aug 11, 2006

STERILE WATER FOR IRRIGATION

LIQUID;IRRIGATION

STERILE WATER

<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N017428 001</u>	
<u>AT</u>	<u>STERILE WATER IN PLASTIC CONTAINER</u>			

<u>AT</u>	B BRAUN	<u>100%</u>	<u>N016734 001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N017866 001</u>	
<u>AT</u>	ICU MEDICAL INC	<u>100%</u>	<u>N017513 001</u>	
<u>AT</u>		<u>100%</u>	<u>N018313 001</u>	

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-392 (of 436)

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

! X GEN PHARMS

EQ 1GM BASE/VIAL

A064210 001 Jun 30, 1998

STREPTOZOCIN

INJECTABLE; INJECTION

ZANOSAR

+! TEVA PHARMS USA

1GM/VIAL

N050577 001 May 07, 1982

STRONTIUM CHLORIDE SR-89

INJECTABLE; INJECTION

METASTRON

AP +! GE HEALTHCARE

1mCi/ML

N020134 001 Jun 18, 1993

STRONTIUM CHLORIDE SR-89

AP BIO NUCLEONICS

1mCi/ML

A075941 001 Jan 06, 2003

SUCCIMER

CAPSULE; ORAL

CHEMET

+! RECORDATI RARE

100MG

N019998 002 Jan 30, 1991

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

ANECTINE

AP +! SANDOZ INC

20MG/ML

N008453 002

QUELICIN

AP +! HOSPIRA

20MG/ML

N008845 006

QUELICIN PRESERVATIVE FREE

AP +! HOSPIRA

20MG/ML

N008845 001

SUCRALFATE

SUSPENSION; ORAL

CARAFATE

+! FOREST LABS INC

1GM/10ML

N019183 001 Dec 16, 1993

TABLET; ORAL

CARAFATE

AB +! FOREST LABS INC

1GM

N018333 001

SUCRALFATE

AB MYLAN IRELAND LTD

1GM

A074415 001 Jun 08, 1998

AB TEVA

1GM

A070848 001 Mar 29, 1996

SUCROFERRIC OXYHYDROXIDE

TABLET, CHEWABLE; ORAL

VELPHORO

+! VIFOR FRESENIUS

500MG

N205109 001 Nov 27, 2013

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA PRESERVATIVE FREE

AP +! AKORN

EQ 0.05MG BASE/ML

N019050 001 May 04, 1984

SUFENTANIL CITRATE

AP HOSPIRA

EQ 0.05MG BASE/ML

A074534 001 Dec 11, 1996

AP WEST-WARD PHARMS INT

EQ 0.05MG BASE/ML

A074413 001 Dec 15, 1995

SUGAMMADEX SODIUM

SOLUTION; INTRAVENOUS

BRIDION

+! ORGANON SUB MERCK

EQ 200MG BASE/2ML (EQ 100MG BASE/ML)

N022225 002 Dec 15, 2015

+

EQ 500MG BASE/5ML (EQ 100MG BASE/ML)

N022225 001 Dec 15, 2015

SULCONAZOLE NITRATE

CREAM; TOPICAL

EXELDERM

+! SUN PHARM INDs INC

1%

N018737 001 Feb 28, 1989

SOLUTION; TOPICAL

EXELDERM

+! SUN PHARM INDs INC

1%

N018738 001 Aug 30, 1985

SULFACETAMIDE SODIUM

LOTION; TOPICAL

KLARON

AB +! VALEANT PHARMS NORTH

10%

N019931 001 Dec 23, 1996

SULFACETAMIDE SODIUM

AB FOUGERA PHARMS

10%

A077015 001 Nov 17, 2006

AB PERRIGO CO

10%

A078649 001 Mar 23, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
 PRESCRIPTION DRUG PRODUCT LIST

3-393 (of 436)

SULFACETAMIDE SODIUM

LOTION;TOPICAL

SULFACETAMIDE SODIUM

TENNESSEE

AB TARO 10% A078668 001 May 20, 2009

OINTMENT;OPHTHALMIC

SULFACETAMIDE SODIUM

! PERRIGO CO 10%

TENNESSEE

SOLUTION/DROPS;OPHTHALMIC

BLEPH-10

AT ! ALLERGAN 10% A080028 001

SULFACETAMIDE SODIUM

AT AKORN 10% A040215 001 May 25, 1999

AT BAUSCH AND LOMB 10% A040066 001 Dec 28, 1994

AT SANDOZ INC 10% A089560 001 Oct 18, 1988

SULFADIAZINE

TABLET;ORAL

SULFADIAZINE

! SANDOZ 500MG

A040091 001 Jul 29, 1994

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE;INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AP MYLAN LABS LTD 80MG/ML;16MG/ML A206607 001 Aug 30, 2017

AP ! TEVA PHARMS USA 80MG/ML;16MG/ML A073303 001 Oct 31, 1991

SUSPENSION;ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB AUROBINDO PHARMA 200MG/5ML;40MG/5ML A091348 001 Jun 08, 2010

AB ! HI TECH PHARMA 200MG/5ML;40MG/5ML A074650 001 Dec 29, 1997

AB VINTAGE 200MG/5ML;40MG/5ML A077785 001 Jan 24, 2007

SULFATRIM PEDIATRIC

AB STI PHARMA LLC 200MG/5ML;40MG/5ML N018615 001 Jan 07, 1983

TABLET;ORAL

BACTRIM

AB + SUN PHARM INDUSTRIES 400MG;80MG N017377 001

BACTRIM DS

AB +! SUN PHARM INDUSTRIES 800MG;160MG N017377 002

SEPTRA

AB MONARCH PHARMS 400MG;80MG N017376 001

SEPTRA DS

AB MONARCH PHARMS 800MG;160MG N017376 002

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB AMNEAL PHARMS NY 400MG;80MG A076899 001 Jan 27, 2005

AB 800MG;160MG A076899 002 Jan 27, 2005

AB AUROBINDO PHARMA 400MG;80MG A090624 001 Feb 16, 2010

AB 800MG;160MG A090624 002 Feb 16, 2010

AB CHARTWELL MOLECULES 400MG;80MG A078060 002 Jan 25, 2007

AB 800MG;160MG A078060 001 Jan 25, 2007

AB GLENMARK GENERICS 400MG;80MG A090828 002 Dec 22, 2010

AB 800MG;160MG A090828 001 Dec 22, 2010

AB SUN PHARM INDUSTRIES 800MG;160MG A071017 001 Aug 25, 1986

AB VISTA PHARMS 400MG;80MG A076817 001 Oct 07, 2005

AB 800MG;160MG A076817 002 Oct 07, 2005

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

AB TEVA 800MG;160MG A070037 001 Jun 02, 1987

SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH

AB TEVA PHARMS 400MG;80MG A070030 001 Jun 02, 1987

SULFAMETHOXAZOLE AND TRIMETHOPRIM

SUN PHARM 400MG;80MG

INDUSTRIES

A071017 002 Aug 25, 1986

SULFANILAMIDE

CREAM;VAGINAL

AVC

+! MYLAN SPECIALITY LP 15%

N006530 003 Jan 27, 1987

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-394 (of 436)

**SULFASALAZINE**

SUSPENSION;ORAL

AZULFIDINE

! PHARMACIA AND  
UPJOHN

TABLET;ORAL

**AZULFIDINE**

<b>AB</b>	+!	PHARMACIA AND UPJOHN	<b><u>500MG</u></b>	<b><u>N007073 001</u></b>
<b>AB</b>		VINTAGE PHARMS	<b><u>500MG</u></b>	<b><u>A040349 001</u></b> Jan 11, 2002
<b>AB</b>		WATSON LABS	<b><u>500MG</u></b>	<b><u>A085828 001</u></b>
		TABLET, DELAYED RELEASE;ORAL		
		<b><u>AZULFIDINE EN-TABS</u></b>		
<b>AB</b>	+!	PHARMACIA AND UPJOHN	<b><u>500MG</u></b>	<b><u>N007073 002</u></b> Apr 06, 1983
		<b><u>SULFASALAZINE</u></b>		
<b>AB</b>		VINTAGE PHARMS	<b><u>500MG</u></b>	<b><u>A075339 001</u></b> Jan 11, 2002

**SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES**

FOR SUSPENSION;INTRAVENOUS

LUMASON

+! BRACCO

**60.7MG/25MG**

N203684 001 Oct 15, 2014

**SULINDAC**

TABLET;ORAL

**SULINDAC**

<b>AB</b>		EPIC PHARMA	<b><u>150MG</u></b>	<b><u>A072710 001</u></b> Mar 25, 1991
<b>AB</b>			<b><u>200MG</u></b>	<b><u>A072711 001</u></b> Mar 25, 1991
<b>AB</b>		EPIC PHARMA LLC	<b><u>150MG</u></b>	<b><u>A073262 002</u></b> Sep 06, 1991
<b>AB</b>			<b><u>200MG</u></b>	<b><u>A073262 001</u></b> Sep 06, 1991
<b>AB</b>		MYLAN	<b><u>150MG</u></b>	<b><u>A073039 002</u></b> Jun 22, 1993
<b>AB</b>			<b><u>200MG</u></b>	<b><u>A073039 001</u></b> Jun 22, 1993
<b>AB</b>		SUN PHARM INDUSTRIES	<b><u>150MG</u></b>	<b><u>A072050 001</u></b> Apr 17, 1991
<b>AB</b>			<b><u>200MG</u></b>	<b><u>A072051 001</u></b> Apr 17, 1991
<b>AB</b>		WATSON LABS	<b><u>150MG</u></b>	<b><u>A071891 001</u></b> Apr 03, 1990
<b>AB</b>	!		<b><u>200MG</u></b>	<b><u>A071795 001</u></b> Apr 03, 1990

**SUMATRIPTAN**

SPRAY;NASAL

**IMITREX**

<b>AB</b>	+!	GLAXOSMITHKLINE	<b><u>5MG/SPRAY</u></b>	<b><u>N020626 001</u></b> Aug 26, 1997
<b>AB</b>	+!		<b><u>20MG/SPRAY</u></b>	<b><u>N020626 003</u></b> Aug 26, 1997
		<b><u>SUMATRIPTAN</u></b>		
<b>AB</b>		LANNETT CO INC	<b><u>5MG/SPRAY</u></b>	<b><u>A204841 001</u></b> Feb 19, 2016
<b>AB</b>			<b><u>20MG/SPRAY</u></b>	<b><u>A204841 002</u></b> Feb 19, 2016

**SUMATRIPTAN SUCCINATE**

INJECTABLE;SUBCUTANEOUS

**IMITREX STATDOSE**

<b>AB</b>	+!	GLAXOSMITHKLINE	<b><u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u></b>	<b><u>N020080 002</u></b> Feb 01, 2006
<b>AB</b>	+!		<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>N020080 003</u></b> Dec 23, 1996
		<b><u>SUMATRIPTAN SUCCINATE</u></b>		

<b>AB</b>		ANTARES PHARMA INC	<b><u>EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)</u></b>	<b><u>A078319 001</u></b> Dec 10, 2015
<b>AB</b>			<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A078319 002</u></b> Dec 10, 2015
<b>AB</b>		DR REDDYS LABS INC	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A090495 001</u></b> Jan 29, 2014
<b>AB</b>		SUN PHARMA GLOBAL	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A090358 001</u></b> Jun 21, 2011

**IMITREX**

<b>AP</b>	+!	GLAXOSMITHKLINE	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>N020080 001</u></b> Dec 28, 1992
		<b><u>SUMATRIPTAN SUCCINATE</u></b>		

<b>AP</b>		AUROBINDO PHARMA LTD	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A202758 001</u></b> Apr 23, 2013
<b>AP</b>		FRESENIUS KABI USA	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A079242 001</u></b> Mar 02, 2009
<b>AP</b>		HIKMA FARMACEUTICA	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A200183 001</u></b> Sep 16, 2013
<b>AP</b>		INJECTALIA	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A090310 001</u></b> Aug 11, 2010
<b>AP</b>		MYLAN LABS LTD	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A203322 001</u></b> Apr 14, 2014

<b>AP</b>		PAR PHARM	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A077332 001</u></b> Oct 09, 2009
<b>AP</b>		PAR STERILE PRODUCTS	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A077871 001</u></b> Jul 09, 2009
<b>AP</b>		SAGENT AGILA	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A090641 001</u></b> Jul 28, 2010
<b>AP</b>		SAGENT AGILA LLC	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A090314 001</u></b> Jun 10, 2010
<b>AP</b>		TEVA PHARMS USA	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A077907 001</u></b> Feb 06, 2009

<b>AP</b>		WEST-WARD PHARMS INT	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A079123 001</u></b> Feb 06, 2009
<b>AP</b>		WOCKHARDT	<b><u>EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)</u></b>	<b><u>A078593 001</u></b> Feb 06, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-395 (of 436)

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS		
SUMAVENT DOSEPRO		
BX +!	ENDO VENTURES LTD	EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)
BX +!		EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)
POWDER; INHALATION		
ONZETRA XSAIL		
+!	AVANIR PHARMS	EQ 11MG BASE
SOLUTION; SUBCUTANEOUS		
ZEMBRACE SYMTOUCH		
DR REDDYS LABS LTD		EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)
TABLET; ORAL		
<b>IMITREX</b>		
<b>AB</b> +	GLAXOSMITHKLINE	<b>EQ 25MG BASE</b>
<b>AB</b> +		<b>EQ 50MG BASE</b>
<b>AB</b> +!		<b>EQ 100MG BASE</b>
<b>SUMATRIPTAN SUCCINATE</b>		
<b>AB</b>	APOTEK INC	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	AUROBINDO PHARMA	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	DR REDDYS LABS INC	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	MYLAN	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	ORCHID HLTHCARE	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	SUN PHARM IND'S	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	SUN PHARM IND'S LTD	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>
<b>AB</b>	WATSON LABS	<b>EQ 25MG BASE</b>
<b>AB</b>		<b>EQ 50MG BASE</b>
<b>AB</b>		<b>EQ 100MG BASE</b>

SUNITINIB MALATE

CAPSULE; ORAL		
SUTENT		
+ CPPI CV		EQ 12.5MG BASE
+		EQ 25MG BASE
+		EQ 37.5MG BASE
+		EQ 50MG BASE

N021938 001 Jan 26, 2006  
N021938 002 Jan 26, 2006  
N021938 004 Mar 31, 2009  
N021938 003 Jan 26, 2006

SUVOREXANT

TABLET; ORAL		
BELSOMRA		
+ MERCK SHARP DOHME		5MG
+		10MG
+		15MG
+		20MG

N204569 001 Aug 13, 2014  
N204569 002 Aug 13, 2014  
N204569 003 Aug 13, 2014  
N204569 004 Aug 13, 2014

TACROLIMUS

CAPSULE; ORAL		
<b>PROGRAF</b>		
<b>AB</b> +	ASTELLAS	<b>EQ 0.5MG BASE</b>
<b>AB</b> +		<b>EQ 1MG BASE</b>
<b>AB</b> +!		<b>EQ 5MG BASE</b>
<b>TACROLIMUS</b>		
<b>AB</b>	ACCORD HLTHCARE	<b>EQ 0.5MG BASE</b>
<b>AB</b>		<b>EQ 1MG BASE</b>
<b>AB</b>		<b>EQ 5MG BASE</b>
<b>AB</b>	BELCHER PHARMS LLC	<b>EQ 0.5MG BASE</b>
<b>AB</b>		<b>EQ 1MG BASE</b>
<b>AB</b>		<b>EQ 5MG BASE</b>
<b>AB</b>	DR REDDYS LABS LTD	<b>EQ 0.5MG BASE</b>
<b>AB</b>		<b>EQ 1MG BASE</b>
<b>AB</b>		<b>EQ 5MG BASE</b>

**N050708 003** Aug 24, 1998  
**N050708 001** Apr 08, 1994  
**N050708 002** Apr 08, 1994

**A091195 001** Aug 31, 2011  
**A091195 002** Aug 31, 2011  
**A091195 003** Aug 31, 2011

**A206651 001** Nov 30, 2017  
**A206651 002** Nov 30, 2017  
**A206651 003** Nov 30, 2017

**A090509 001** May 12, 2010  
**A090509 002** May 12, 2010  
**A090509 003** May 12, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-396 (of 436)

**TACROLIMUS**

CAPSULE;ORAL

**TACROLIMUS**

<b>AB</b>	MYLAN	<b>EQ 0.5MG BASE</b>	<b>A090596 001</b>	Sep 17, 2010
<b>AB</b>		<b>EQ 1MG BASE</b>	<b>A090596 002</b>	Sep 17, 2010
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A090596 003</b>	Sep 17, 2010
<b>AB</b>	PANACEA BIOTEC LTD	<b>EQ 0.5MG BASE</b>	<b>A090802 001</b>	Sep 28, 2012
<b>AB</b>		<b>EQ 1MG BASE</b>	<b>A090802 002</b>	Sep 28, 2012
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A090802 003</b>	Sep 28, 2012
<b>AB</b>	SANDOZ	<b>EQ 0.5MG BASE</b>	<b>A065461 001</b>	Aug 10, 2009
<b>AB</b>		<b>EQ 1MG BASE</b>	<b>A065461 002</b>	Aug 10, 2009
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A065461 003</b>	Aug 10, 2009
<b>AB</b>	STRIDES PHARMA	<b>EQ 0.5MG BASE</b>	<b>A090687 001</b>	Jul 22, 2014
<b>AB</b>		<b>EQ 1MG BASE</b>	<b>A090687 002</b>	Jul 22, 2014
<b>AB</b>		<b>EQ 5MG BASE</b>	<b>A090687 003</b>	Jul 22, 2014

CAPSULE, EXTENDED RELEASE;ORAL

ASTAGRAF XL

+ ASTELLAS

**EQ 0.5MG BASE**

N204096 001 Jul 19, 2013

+ ASTELLAS

**EQ 1MG BASE**

N204096 002 Jul 19, 2013

+!

**EQ 5MG BASE**

N204096 003 Jul 19, 2013

INJECTABLE;INJECTION

**PROGRAF**

<b>AP</b>	+! ASTELLAS	<b>EQ 5MG BASE/ML</b>	<b>N050709 001</b>	Apr 08, 1994
-----------	-------------	-----------------------	--------------------	--------------

**TACROLIMUS**

<b>AP</b>	HOSPIRA INC	<b>EQ 5MG BASE/ML</b>	<b>A203900 001</b>	Aug 25, 2017
-----------	-------------	-----------------------	--------------------	--------------

OINTMENT;TOPICAL

**PROTOPIC**

<b>AB</b>	+! LEO PHARMA AS	<b>0.03%</b>	<b>N050777 001</b>	Dec 08, 2000
-----------	------------------	--------------	--------------------	--------------

<b>AB</b>	+!	<b>0.1%</b>	<b>N050777 002</b>	Dec 08, 2000
-----------	----	-------------	--------------------	--------------

**TACROLIMUS**

<b>AB</b>	FOUGERA PHARMS INC	<b>0.03%</b>	<b>A200744 001</b>	Sep 09, 2014
-----------	--------------------	--------------	--------------------	--------------

<b>AB</b>		<b>0.1%</b>	<b>A200744 002</b>	Sep 09, 2014
-----------	--	-------------	--------------------	--------------

TABLET, EXTENDED RELEASE;ORAL

ENVARSUS XR

+ VELOXIS PHARMS INC

**EQ 0.75MG BASE**

N206406 001 Jul 10, 2015

+ VELOXIS PHARMS INC

**EQ 1MG BASE**

N206406 002 Jul 10, 2015

+!

**EQ 4MG BASE**

N206406 003 Jul 10, 2015

**Tadalafil**

TABLET;ORAL

ADCIRCA

+! ELI LILLY CO

**20MG**

N022332 001 May 22, 2009

CIALIS

+ LILLY

**2.5MG**

N021368 004 Jan 07, 2008

+

**5MG**

N021368 001 Nov 21, 2003

+

**10MG**

N021368 002 Nov 21, 2003

+

**20MG**

N021368 003 Nov 21, 2003

**Tafluprost**

SOLUTION/DROPS;OPHTHALMIC

ZIOPTAN

+! OAK PHARMS INC

**0.0015%**

N202514 001 Feb 10, 2012

**TALC**

AEROSOL, METERED;INTRAPLEURAL

SCLEROSOL

+! LYMOL MEDCL

**400MG/SPRAY**

N020587 001 Dec 24, 1997

POWDER;INTRAPLEURAL

STERITALC

+ NOVATECH SA

**2GM/VIAL**

N205555 001 May 01, 2017

+

**3GM/VIAL**

N205555 002 May 01, 2017

+

**4GM/VIAL**

N205555 003 May 01, 2017

TALC

+! LYMOL MEDCL

**5GM/BOT**

N021388 001 Dec 15, 2003

**Taliglucerase Alfa**

POWDER;IV (INFUSION)

ELELYSO

+! PFIZER

**200 UNITS/VIAL**

N022458 001 May 01, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-397 (of 436)

**TAMOXIFEN CITRATE**

SOLUTION;ORAL  
 SOLTAMOX  
 MIDATECH PHARMA US EQ 10MG BASE/5ML N021807 001 Oct 29, 2005

TABLET;ORAL

**TAMOXIFEN CITRATE**

<b>AB</b>	ACTAVIS LABS FL INC	<b>EQ 10MG BASE</b>	<b>A070929 001</b>	Feb 20, 2003
<b>AB</b>		<b>EQ 20MG BASE</b>	<b>A070929 002</b>	Feb 20, 2003
<b>AB</b>	APOTEX	<b>EQ 10MG BASE</b>	<b>A090878 001</b>	Sep 23, 2011
<b>AB</b>		<b>EQ 20MG BASE</b>	<b>A090878 002</b>	Sep 23, 2011
<b>AB</b>	MAYNE PHARMA	<b>EQ 10MG BASE</b>	<b>A075797 001</b>	Feb 20, 2003
<b>AB</b>	!	<b>EQ 20MG BASE</b>	<b>A075797 002</b>	Feb 20, 2003
<b>AB</b>	MYLAN	<b>EQ 10MG BASE</b>	<b>A074732 002</b>	Feb 20, 2003
<b>AB</b>		<b>EQ 20MG BASE</b>	<b>A074732 001</b>	Feb 20, 2003
<b>AB</b>	ZYDUS PHARMS USA INC	<b>EQ 10MG BASE</b>	<b>A206694 001</b>	Oct 27, 2017
<b>AB</b>		<b>EQ 20MG BASE</b>	<b>A206694 002</b>	Oct 27, 2017

**TAMSULOSIN HYDROCHLORIDE**

CAPSULE;ORAL

**FLOMAX**

<b>AB</b>	+!	BOEHRINGER INGELHEIM	<b>0 . 4MG</b>	<b>N020579 001</b>	Apr 15, 1997
-----------	----	----------------------	----------------	--------------------	--------------

**TAMSULOSIN HYDROCHLORIDE**

<b>AB</b>	ANCHEM PHARMS	<b>0 . 4MG</b>	<b>A202010 001</b>	Jan 04, 2013
<b>AB</b>	AUROBINDO PHARMA LTD	<b>0 . 4MG</b>	<b>A202433 001</b>	Apr 30, 2013
<b>AB</b>	IMPAKX LABS	<b>0 . 4MG</b>	<b>A090377 001</b>	Mar 02, 2010
<b>AB</b>	MACLEODS PHARMS LTD	<b>0 . 4MG</b>	<b>A204645 001</b>	Jan 20, 2017
<b>AB</b>	MYLAN	<b>0 . 4MG</b>	<b>A090408 001</b>	Apr 27, 2010
<b>AB</b>	SANDOZ	<b>0 . 4MG</b>	<b>A078015 001</b>	Apr 27, 2010
<b>AB</b>	SUN PHARM INDs LTD	<b>0 . 4MG</b>	<b>A090931 001</b>	Jul 15, 2010
<b>AB</b>	SYNTHON PHARMS	<b>0 . 4MG</b>	<b>A078801 001</b>	Apr 27, 2010
<b>AB</b>	TEVA PHARMS	<b>0 . 4MG</b>	<b>A077630 001</b>	Apr 27, 2010
<b>AB</b>	WOCKHARDT	<b>0 . 4MG</b>	<b>A078938 001</b>	Apr 27, 2010
<b>AB</b>	ZYDUS PHARMS USA INC	<b>0 . 4MG</b>	<b>A078225 001</b>	Apr 27, 2010

**TAMSULOSIN HYDROCHLORIDE**

<b>AB</b>	ALKEM LABS LTD	<b>0 . 4MG</b>	<b>A207405 001</b>	Aug 11, 2017
-----------	----------------	----------------	--------------------	--------------

**TAPENTADOL HYDROCHLORIDE**

SOLUTION;ORAL

NUCYNTA				
+!	DEPOMED INC	EQ 20MG BASE/ML	N203794 001	Oct 15, 2012

TABLET;ORAL

NUCYNTA				
+!	DEPOMED INC	EQ 50MG BASE	N022304 001	Nov 20, 2008
+		EQ 75MG BASE	N022304 002	Nov 20, 2008
+		EQ 100MG BASE	N022304 003	Nov 20, 2008

TABLET, EXTENDED RELEASE;ORAL

NUCYNTA ER				
+!	DEPOMED INC	EQ 50MG BASE	N200533 001	Aug 25, 2011
+		EQ 100MG BASE	N200533 002	Aug 25, 2011
+		EQ 150MG BASE	N200533 003	Aug 25, 2011
+		EQ 200MG BASE	N200533 004	Aug 25, 2011
+		EQ 250MG BASE	N200533 005	Aug 25, 2011

**TASIMELTEON**

CAPSULE;ORAL				
HETLIOZ				
+!	VANDA PHARMS INC	20MG	N205677 001	Jan 31, 2014

**TAVABOROLE**

SOLUTION;TOPICAL				
KERYDIN				
+!	ANACOR PHARMS INC	5%	N204427 001	Jul 07, 2014

**TAZAROTENE**

AEROSOL, FOAM;TOPICAL				
FABIOR				
+!	MAYNE PHARMA	0 . 1%	N202428 001	May 11, 2012

CREAM;TOPICAL

<b>AVAGE</b>					
<b>AB</b>	+!	ALLERGAN	<b>0 . 1%</b>	<b>N021184 003</b>	Sep 30, 2002

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-398 (of 436)

TAZAROTENE

CREAM;TOPICAL

**TAZAROTENE**

<b>AB</b>	G AND W LABS INC	<b>0.1%</b>	<b>A208662 001</b>	Dec 22, 2017
<b>AB</b>	TARO	<b>0.1%</b>	<b>A208258 001</b>	Apr 03, 2017
	<b>TAZORAC</b>			
<b>AB</b>	+! ALLERGAN	<b>0.1%</b>	<b>N021184 002</b>	Sep 29, 2000
	+!	0.05%	N021184 001	Sep 29, 2000
	GEL;TOPICAL			
	TAZORAC			
	+! ALLERGAN	0.05%	N020600 001	Jun 13, 1997
	+!	0.1%	N020600 002	Jun 13, 1997

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE;INJECTION

PULMOLITE

BS	+! JUBILANT DRAXIMAGE	N/A	N017776 001
BS	+! DRAXIMAGE	N/A	N017881 001 Dec 30, 1987

TECHNETIUM TC-99M BICISATE KIT

INJECTABLE;INJECTION

NEUROLITE

+!	LANTHEUS MEDCL	N/A
----	----------------	-----

N020256 001 Nov 23, 1994

TECHNETIUM TC-99M DISOFENIN KIT

INJECTABLE;INJECTION

HEPATOLITE

PHARMALUCENCE	N/A
---------------	-----

N018467 001 Mar 16, 1982

TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE;INJECTION

CERETEC

+!	GE HEALTHCARE	N/A
----	---------------	-----

N019829 001 Dec 30, 1988

POWDER;INTRAVENOUS

DRAX EXAMETAZIME

JUBILANT DRAXIMAGE	N/A
--------------------	-----

N208870 001 Aug 17, 2017

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE;INJECTION

**CHOLETEC**

<b>AP</b>	+! BRACCO	<b>N/A</b>	<b>N018963 001</b>	Jan 21, 1987
<b>AP</b>	PHARMALUCENCE	<b>N/A</b>	<b>A078242 001</b>	Jan 29, 2008

TECHNETIUM TC-99M MEDRONATE

INJECTABLE;INJECTION

DRAXIMAGE MDP-25

+!	JUBILANT DRAXIMAGE	N/A
----	--------------------	-----

N018035 002 Feb 27, 2004

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE;INJECTION

**CIS-MDP**

<b>AP</b>	PHARMALUCENCE	<b>N/A</b>	<b>N018124 001</b>
<b>AP</b>	CARDINAL HEALTH 414	<b>N/A</b>	<b>N018107 001</b>

TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE;INJECTION

TECHNESCAN MAG3

+!	MALLINKRODT NUCLEAR	N/A
----	---------------------	-----

N019882 001 Jun 15, 1990

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE;INJECTION

TECHNESCAN

+!	MALLINKRODT NUCLEAR	N/A
----	---------------------	-----

N018321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE;INJECTION

DTPA

+!	DRAXIMAGE	N/A
----	-----------	-----

N018511 001 Dec 29, 1989

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-399 (of 436)

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYRO

AP PHARMALUCENCE N/A

N019039 001 Jun 30, 1987

AP MALLINKRODT NUCLEAR N/A

N017538 001

TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

+! MALLINKRODT NUCLEAR N/A

N019981 001 Jun 10, 1991

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE

AP +! LANTHEUS MEDCL N/A

N019785 001 Dec 21, 1990

TECHNETIUM TC 99M SESTAMIBI

AP CARDINAL HEALTH 414 N/A

A078809 001 Apr 28, 2009

AP JUBILANT DRAXIMAGE N/A

A078806 001 Apr 29, 2009

AP PHARMALUCENCE 10-30mCi

A079157 001 Jul 10, 2009

TECHNETIUM TC-99M SESTAMIBI

AP MALLINKRODT NUCLEAR N/A

A078098 001 Sep 22, 2008

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INTRAVENOUS

TECHNELITE

+! LANTHEUS MEDCL 1-20 CI/GENERATOR

N017771 002 Feb 12, 2014

ULTRA-TECHNEKOW FM

+! MALLINKRODT NUCLEAR 1-19 CI/GENERATOR

N017243 003 Feb 18, 2014

SOLUTION; INTRAVENOUS, ORAL

TECHNETIUM TC 99M GENERATOR

+! GE HEALTHCARE 68-2703mCi/GENERATOR

N017693 002 Dec 13, 2013

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOID

+! PHARMALUCENCE N/A

N017858 001

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW 30ML

+! GE HEALTHCARE N/A

N020372 002 Jul 07, 2005

TECHNETIUM TC-99M TILMANOCEPT

INJECTABLE; INJECTION

LYMPHOSEEK KIT

+! CARDINAL HEALTH 414 N/A

N202207 001 Mar 13, 2013

TEDIZOLID PHOSPHATE

POWDER; IV (INFUSION)

SIVEXTRO

+! CUBIST PHARMS LLC 200MG/VIAL

N205436 001 Jun 20, 2014

TABLET; ORAL

SIVEXTRO

+! CUBIST PHARMS LLC 200MG

N205435 001 Jun 20, 2014

TEDUGLUTIDE RECOMBINANT

POWDER; SUBCUTANEOUS

GATTEX KIT

+! NPS PHARMS INC 5MG/VIAL

N203441 001 Dec 21, 2012

TELAVANCIN HYDROCHLORIDE

POWDER; IV (INFUSION)

VIBATIV

+! THERAVANCE EQ 750MG BASE/VIAL  
BIOPHARMA

N022110 002 Sep 11, 2009

TELMISARTAN

TABLET; ORAL

MICARDIS

AB + BOEHRINGER 20MG

N020850 003 Apr 04, 2000

INGELHEIM

AB + 40MG

N020850 001 Nov 10, 1998

AB +! 80MG

N020850 002 Nov 10, 1998

TELMISARTAN

AB ALEMBIC PHARMS LTD 20MG

A202130 001 Jul 07, 2014

AB 40MG

A202130 002 Jul 07, 2014

AB 80MG

A202130 003 Jul 07, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-400 (of 436)

TELMISARTAN

TABLET;ORAL

TELMISARTAN

<u>AB</u>	AMNEAL PHARMS	<u>20MG</u>	<u>A204415 001</u>	Sep 08, 2015
<u>AB</u>		<u>40MG</u>	<u>A204415 002</u>	Sep 08, 2015
<u>AB</u>		<u>80MG</u>	<u>A204415 003</u>	Sep 08, 2015
<u>AB</u>	AUROBINDO PHARMA LTD	<u>20MG</u>	<u>A206511 001</u>	Sep 03, 2015
<u>AB</u>		<u>40MG</u>	<u>A206511 002</u>	Sep 03, 2015
<u>AB</u>		<u>80MG</u>	<u>A206511 003</u>	Sep 03, 2015
<u>AB</u>	CADILA PHARMS LTD	<u>20MG</u>	<u>A208605 001</u>	Jul 25, 2017
<u>AB</u>		<u>40MG</u>	<u>A208605 002</u>	Jul 25, 2017
<u>AB</u>		<u>80MG</u>	<u>A208605 003</u>	Jul 25, 2017
<u>AB</u>	GLENMARK PHARMS LTD	<u>20MG</u>	<u>A090032 001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A090032 002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A090032 003</u>	Jul 07, 2014
<u>AB</u>	HETERO LABS LTD V	<u>20MG</u>	<u>A205901 001</u>	Apr 22, 2016
<u>AB</u>		<u>40MG</u>	<u>A205901 002</u>	Apr 22, 2016
<u>AB</u>		<u>80MG</u>	<u>A205901 003</u>	Apr 22, 2016
<u>AB</u>	INVENTIA HLTHCARE	<u>20MG</u>	<u>A205150 001</u>	Oct 30, 2015
<u>AB</u>		<u>40MG</u>	<u>A205150 002</u>	Oct 30, 2015
<u>AB</u>		<u>80MG</u>	<u>A205150 003</u>	Oct 30, 2015
<u>AB</u>	JUBILANT GENERICS	<u>20MG</u>	<u>A204164 001</u>	Aug 22, 2016
<u>AB</u>		<u>40MG</u>	<u>A204164 002</u>	Aug 22, 2016
<u>AB</u>		<u>80MG</u>	<u>A204164 003</u>	Aug 22, 2016
<u>AB</u>	MICRO LABS	<u>20MG</u>	<u>A207016 001</u>	Oct 03, 2017
<u>AB</u>		<u>40MG</u>	<u>A207016 002</u>	Oct 03, 2017
<u>AB</u>		<u>80MG</u>	<u>A207016 003</u>	Oct 03, 2017
<u>AB</u>	MYLAN PHARMS INC	<u>20MG</u>	<u>A202397 001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A202397 002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A202397 003</u>	Jul 07, 2014
<u>AB</u>	PRINSTON INC	<u>20MG</u>	<u>A207882 001</u>	May 03, 2017
<u>AB</u>		<u>40MG</u>	<u>A207882 002</u>	May 03, 2017
<u>AB</u>		<u>80MG</u>	<u>A207882 003</u>	May 03, 2017
<u>AB</u>	SANDOZ INC	<u>20MG</u>	<u>A203867 001</u>	Nov 03, 2014
<u>AB</u>		<u>40MG</u>	<u>A203867 002</u>	Nov 03, 2014
<u>AB</u>		<u>80MG</u>	<u>A203867 003</u>	Nov 03, 2014
<u>AB</u>	TORRENT PHARMS LTD	<u>20MG</u>	<u>A203171 001</u>	Jul 07, 2014
<u>AB</u>		<u>40MG</u>	<u>A203171 002</u>	Jul 07, 2014
<u>AB</u>		<u>80MG</u>	<u>A203171 003</u>	Jul 07, 2014
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>A078710 001</u>	Jan 08, 2014
<u>AB</u>		<u>40MG</u>	<u>A078710 002</u>	Jan 08, 2014
<u>AB</u>		<u>80MG</u>	<u>A078710 003</u>	Jan 08, 2014
<u>AB</u>	ZYDUS PHARMS USA INC	<u>20MG</u>	<u>A203325 001</u>	Aug 26, 2014
<u>AB</u>		<u>40MG</u>	<u>A203325 002</u>	Aug 26, 2014
<u>AB</u>		<u>80MG</u>	<u>A203325 003</u>	Aug 26, 2014

TELOTRISTAT ETIPRATE

TABLET;ORAL

XERMELO

+! LEXICON PHARMS INC EQ 250MG BASE

N208794 001 Feb 28, 2017

TEMAZEPEPAM

CAPSULE;ORAL

RESTORIL

<u>AB</u>	+	SPECGX LLC	<u>7.5MG</u>	<u>N018163 003</u>	Oct 25, 1991
<u>AB</u>	+		<u>15MG</u>	<u>N018163 001</u>	
<u>AB</u>	+		<u>22.5MG</u>	<u>N018163 004</u>	Nov 02, 2004
<u>AB</u>	+		<u>30MG</u>	<u>N018163 002</u>	

TEMAZEPEPAM

<u>AB</u>	ACTAVIS ELIZABETH	<u>15MG</u>	<u>A071620 002</u>	Aug 07, 1987
<u>AB</u>		<u>30MG</u>	<u>A071620 001</u>	Aug 07, 1987
<u>AB</u>	AMNEAL PHARMS	<u>7.5MG</u>	<u>A203482 001</u>	May 23, 2016
<u>AB</u>		<u>15MG</u>	<u>A203482 002</u>	May 23, 2016
<u>AB</u>		<u>22.5MG</u>	<u>A203482 003</u>	May 23, 2016
<u>AB</u>		<u>30MG</u>	<u>A203482 004</u>	May 23, 2016
<u>AB</u>	MYLAN	<u>7.5MG</u>	<u>A070920 002</u>	May 21, 2010
<u>AB</u>		<u>15MG</u>	<u>A070920 004</u>	Jul 07, 1986
<u>AB</u>		<u>22.5MG</u>	<u>A070920 003</u>	Jun 12, 2009
<u>AB</u>		<u>30MG</u>	<u>A070920 001</u>	Jul 10, 1986
<u>AB</u>	NOVEL LABS INC	<u>7.5MG</u>	<u>A071457 002</u>	Jun 22, 2012
<u>AB</u>		<u>15MG</u>	<u>A071456 001</u>	Apr 21, 1987
<u>AB</u>		<u>22.5MG</u>	<u>A071457 003</u>	Jun 22, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-401 (of 436)

TEMAZEPAM

CAPSULE; ORAL

TEMAZEPAM

<u>AB</u>		<u>30MG</u>	<u>A071457 001</u>	Apr 21, 1987
<u>AB</u>	PRINSTON INC	<u>7.5MG</u>	<u>A201781 001</u>	Jun 04, 2015
<u>AB</u>		<u>15MG</u>	<u>A201781 002</u>	Jun 04, 2015
<u>AB</u>		<u>22.5MG</u>	<u>A201781 003</u>	Jun 04, 2015
<u>AB</u>		<u>30MG</u>	<u>A201781 004</u>	Jun 04, 2015
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>A071427 001</u>	Jan 12, 1988
<u>AB</u>		<u>30MG</u>	<u>A071428 001</u>	Jan 12, 1988
<u>AB</u>	SUN PHARM INDUSTRIES	<u>7.5MG</u>	<u>A078581 001</u>	Sep 08, 2009
<u>AB</u>		<u>22.5MG</u>	<u>A071175 002</u>	Sep 14, 2009

TEMOZOLOMIDE

CAPSULE; ORAL

TEMODAR

<u>AB</u> +	MERCK SHARP DOHME	<u>5MG</u>	<u>N021029 001</u>	Aug 11, 1999
<u>AB</u> +		<u>20MG</u>	<u>N021029 002</u>	Aug 11, 1999
<u>AB</u> +		<u>100MG</u>	<u>N021029 003</u>	Aug 11, 1999
<u>AB</u> +		<u>140MG</u>	<u>N021029 005</u>	Oct 19, 2006
<u>AB</u> +		<u>180MG</u>	<u>N021029 006</u>	Oct 19, 2006
<u>AB</u> +!		<u>250MG</u>	<u>N021029 004</u>	Aug 11, 1999

TEMOZOLOMIDE

<u>AB</u>	ACCORD HLTHCARE	<u>5MG</u>	<u>A201528 001</u>	Feb 27, 2017
<u>AB</u>		<u>20MG</u>	<u>A201528 002</u>	Feb 27, 2017
<u>AB</u>		<u>100MG</u>	<u>A201528 003</u>	Feb 27, 2017
<u>AB</u>		<u>140MG</u>	<u>A201528 004</u>	Feb 27, 2017
<u>AB</u>		<u>180MG</u>	<u>A201528 005</u>	Feb 27, 2017
<u>AB</u>		<u>250MG</u>	<u>A201528 006</u>	Feb 27, 2017
<u>AB</u>	AMERIGEN PHARMS LTD	<u>5MG</u>	<u>A203490 001</u>	Jul 13, 2016
<u>AB</u>		<u>20MG</u>	<u>A203490 002</u>	Jul 13, 2016
<u>AB</u>		<u>100MG</u>	<u>A203490 003</u>	Jul 13, 2016
<u>AB</u>		<u>140MG</u>	<u>A203490 004</u>	Jul 13, 2016
<u>AB</u>		<u>180MG</u>	<u>A203490 005</u>	Jul 13, 2016
<u>AB</u>		<u>250MG</u>	<u>A203490 006</u>	Jul 13, 2016
<u>AB</u>	AMNEAL PHARMS	<u>5MG</u>	<u>A203691 001</u>	May 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203691 002</u>	May 08, 2015
<u>AB</u>		<u>100MG</u>	<u>A203691 003</u>	May 08, 2015
<u>AB</u>		<u>140MG</u>	<u>A203691 004</u>	May 08, 2015
<u>AB</u>		<u>180MG</u>	<u>A203691 005</u>	May 08, 2015
<u>AB</u>		<u>250MG</u>	<u>A203691 006</u>	May 08, 2015
<u>AB</u>	BARR	<u>5MG</u>	<u>A203691 001</u>	May 08, 2015
<u>AB</u>		<u>20MG</u>	<u>A203691 002</u>	May 08, 2015
<u>AB</u>		<u>100MG</u>	<u>A203691 003</u>	May 08, 2015
<u>AB</u>		<u>140MG</u>	<u>A203691 004</u>	May 08, 2015
<u>AB</u>		<u>180MG</u>	<u>A203691 005</u>	May 08, 2015
<u>AB</u>		<u>250MG</u>	<u>A203691 006</u>	May 08, 2015
<u>AB</u>	CHEMI SPA	<u>5MG</u>	<u>A204639 001</u>	Nov 23, 2016
<u>AB</u>		<u>20MG</u>	<u>A204639 002</u>	Nov 23, 2016
<u>AB</u>		<u>100MG</u>	<u>A204639 003</u>	Nov 23, 2016
<u>AB</u>		<u>140MG</u>	<u>A204639 004</u>	Nov 23, 2016
<u>AB</u>		<u>180MG</u>	<u>A204639 005</u>	Nov 23, 2016
<u>AB</u>		<u>250MG</u>	<u>A204639 006</u>	Nov 23, 2016
<u>AB</u>	DEVA HOLDING AS	<u>5MG</u>	<u>A207658 001</u>	Apr 26, 2017
<u>AB</u>		<u>20MG</u>	<u>A207658 002</u>	Apr 26, 2017
<u>AB</u>		<u>100MG</u>	<u>A207658 003</u>	Apr 26, 2017
<u>AB</u>		<u>140MG</u>	<u>A207658 004</u>	Apr 26, 2017
<u>AB</u>		<u>180MG</u>	<u>A207658 005</u>	Apr 26, 2017
<u>AB</u>		<u>250MG</u>	<u>A207658 006</u>	Apr 26, 2017
<u>AB</u>	IDT AUSTRALIA LTD	<u>5MG</u>	<u>A206413 001</u>	Apr 12, 2016
<u>AB</u>		<u>20MG</u>	<u>A206413 002</u>	Apr 12, 2016
<u>AB</u>		<u>100MG</u>	<u>A206413 003</u>	Apr 12, 2016
<u>AB</u>		<u>140MG</u>	<u>A206413 004</u>	Apr 12, 2016
<u>AB</u>		<u>180MG</u>	<u>A206413 005</u>	Apr 12, 2016
<u>AB</u>		<u>250MG</u>	<u>A206413 006</u>	Apr 12, 2016
<u>AB</u>	KREMERS URBAN PHARMS	<u>5MG</u>	<u>A203898 001</u>	Feb 10, 2016
<u>AB</u>		<u>20MG</u>	<u>A203898 002</u>	Feb 10, 2016
<u>AB</u>		<u>100MG</u>	<u>A203898 003</u>	Feb 10, 2016
<u>AB</u>		<u>140MG</u>	<u>A203898 004</u>	Feb 10, 2016
<u>AB</u>		<u>180MG</u>	<u>A203898 005</u>	Feb 10, 2016
<u>AB</u>		<u>250MG</u>	<u>A203898 006</u>	Feb 10, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-402 (of 436)

TEMOZOLOMIDE

CAPSULE; ORAL

TEMOZOLOMIDE

<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A205227 001</u>	Jun 29, 2016
<u>AB</u>		<u>20MG</u>	<u>A205227 002</u>	Jun 29, 2016
<u>AB</u>		<u>100MG</u>	<u>A205227 003</u>	Jun 29, 2016
<u>AB</u>		<u>140MG</u>	<u>A205227 004</u>	Jun 29, 2016
<u>AB</u>		<u>180MG</u>	<u>A205227 005</u>	Jun 29, 2016
<u>AB</u>		<u>250MG</u>	<u>A205227 006</u>	Jun 29, 2016
<u>AB</u>	RISING PHARMS INC	<u>5MG</u>	<u>A206309 001</u>	Apr 27, 2016
<u>AB</u>		<u>20MG</u>	<u>A206309 002</u>	Apr 27, 2016
<u>AB</u>		<u>100MG</u>	<u>A206309 003</u>	Apr 27, 2016
<u>AB</u>		<u>140MG</u>	<u>A206309 004</u>	Apr 27, 2016
<u>AB</u>		<u>180MG</u>	<u>A206309 005</u>	Apr 27, 2016
<u>AB</u>		<u>250MG</u>	<u>A206309 006</u>	Apr 27, 2016
<u>AB</u>	SUN PHARMA GLOBAL	<u>5MG</u>	<u>A201742 001</u>	Feb 12, 2014
<u>AB</u>		<u>20MG</u>	<u>A201742 002</u>	Feb 12, 2014
<u>AB</u>		<u>100MG</u>	<u>A201742 003</u>	Feb 12, 2014
<u>AB</u>		<u>140MG</u>	<u>A201742 004</u>	Feb 12, 2014
<u>AB</u>		<u>180MG</u>	<u>A201742 005</u>	Feb 12, 2014
<u>AB</u>		<u>250MG</u>	<u>A201742 006</u>	Feb 12, 2014
<u>AB</u>	WATSON LABS TEVA	<u>5MG</u>	<u>A203959 001</u>	Apr 18, 2017
<u>AB</u>		<u>20MG</u>	<u>A203959 002</u>	Apr 18, 2017
<u>AB</u>		<u>100MG</u>	<u>A203959 003</u>	Apr 18, 2017
<u>AB</u>		<u>140MG</u>	<u>A203959 004</u>	Apr 18, 2017
<u>AB</u>		<u>250MG</u>	<u>A203959 005</u>	Apr 18, 2017
<u>AB</u>	ZYDUS PHARMS USA INC	<u>5MG</u>	<u>A206750 001</u>	Jul 31, 2017
<u>AB</u>		<u>20MG</u>	<u>A206750 002</u>	Jul 31, 2017
<u>AB</u>		<u>100MG</u>	<u>A206750 003</u>	Jul 31, 2017
<u>AB</u>		<u>140MG</u>	<u>A206750 004</u>	Jul 31, 2017
<u>AB</u>		<u>180MG</u>	<u>A206750 005</u>	Jul 31, 2017
<u>AB</u>		<u>250MG</u>	<u>A206750 006</u>	Jul 31, 2017

POWDER; INTRAVENOUS

TEMODAR

+! MERCK SHARP DOHME 100MG/VIAL

N022277 001 Feb 27, 2009

TEMSIROLIMUS

SOLUTION; INTRAVENOUS

TORISEL

+! PF PRISM CV 25MG/ML (25MG/ML)

N022088 001 May 30, 2007

TENOFOVIR ALAFENAMIDE FUMARATE

TABLET; ORAL

VEMLIDY

+! GILEAD SCIENCES INC EQ 25MG BASE

N208464 001 Nov 10, 2016

TENOFOVIR DISOPROXIL FUMARATE

POWDER; ORAL

VIREAD

+! GILEAD SCIENCES INC 40MG/SCOOPFUL

N022577 001 Jan 18, 2012

TABLET; ORAL

TENOFOVIR DISOPROXIL FUMARATE

<u>AB</u>	TEVA PHARMS USA	<u>300MG</u>	<u>A091612 001</u>	Mar 18, 2015
<u>AB</u>	<u>VIREAD</u>			
<u>AB</u>	+! GILEAD SCIENCES INC	<u>300MG</u>	<u>N021356 001</u>	Oct 26, 2001
	+	150MG	N021356 002	Jan 18, 2012
	+	200MG	N021356 003	Jan 18, 2012
	+	250MG	N021356 004	Jan 18, 2012

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

<u>AB</u>	APOTEX	<u>EQ 1MG BASE</u>	<u>A075498 001</u>	Apr 12, 2001
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075498 002</u>	Apr 12, 2001
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075498 003</u>	Apr 12, 2001
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075498 004</u>	Apr 12, 2001
<u>AB</u>	IVAX SUB TEVA PHARMS	<u>EQ 1MG BASE</u>	<u>A075614 002</u>	Jan 30, 2001
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075614 001</u>	Jan 30, 2001
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075614 003</u>	Jan 30, 2001
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075614 004</u>	Jan 30, 2001
<u>AB</u>	JUBILANT CADISTA	<u>EQ 1MG BASE</u>	<u>A075317 001</u>	Dec 20, 2004
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075317 002</u>	Dec 20, 2004
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075317 003</u>	Dec 20, 2004

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-403 (of 436)

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

TERAZOSIN HYDROCHLORIDE

<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075317 004</u>	Dec 20, 2004
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 1MG BASE</u>	<u>A075140 002</u>	Feb 11, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>A075140 003</u>	Feb 11, 2000
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A075140 001</u>	Feb 11, 2000
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A075140 004</u>	Feb 11, 2000
<u>AB</u>	SANDOZ	<u>EQ 1MG BASE</u>	<u>A074823 001</u>	Mar 30, 1998
<u>AB</u>	!	<u>EQ 2MG BASE</u>	<u>A074823 002</u>	Mar 30, 1998
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>A074823 003</u>	Mar 30, 1998
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>A074823 004</u>	Mar 30, 1998

TERBINAFINE HYDROCHLORIDE

TABLET; ORAL

LAMISIL

<u>AB</u>	+!	NOVARTIS	<u>EQ 250MG BASE</u>	<u>N020539 001</u>	May 10, 1996
		<u>TERBINAFINE HYDROCHLORIDE</u>			
<u>AB</u>		APOTEX	<u>EQ 250MG BASE</u>	<u>A078199 001</u>	Jul 02, 2007
<u>AB</u>		AUROBINDO PHARMA	<u>EQ 250MG BASE</u>	<u>A078297 001</u>	Jul 02, 2007
<u>AB</u>		BRECKENRIDGE PHARM	<u>EQ 250MG BASE</u>	<u>A077714 001</u>	Jun 04, 2010
<u>AB</u>		CIPLA LTD	<u>EQ 250MG BASE</u>	<u>A077137 001</u>	Jul 02, 2007
<u>AB</u>		DR REDDYS LABS INC	<u>EQ 250MG BASE</u>	<u>A076390 001</u>	Jul 02, 2007
<u>AB</u>		GLENMARK GENERICS	<u>EQ 250MG BASE</u>	<u>A078157 001</u>	Jul 02, 2007
<u>AB</u>		HARRIS PHARM	<u>EQ 250MG BASE</u>	<u>A077919 001</u>	Jul 02, 2007
<u>AB</u>		INVAGEN PHARMS	<u>EQ 250MG BASE</u>	<u>A077533 001</u>	Jul 02, 2007
<u>AB</u>		ORCHID HLTHCARE	<u>EQ 250MG BASE</u>	<u>A078163 001</u>	Jul 02, 2007
<u>AB</u>		TEVA	<u>EQ 250MG BASE</u>	<u>A076377 001</u>	Jul 02, 2007

TERBUTALINE SULFATE

INJECTABLE; INJECTION

TERBUTALINE SULFATE

<u>AP</u>		AKORN	<u>1MGM/ML</u>	<u>A078151 001</u>	Jan 07, 2008
<u>AP</u>	!	ATHENEX INC	<u>1MGM/ML</u>	<u>A076770 001</u>	Apr 23, 2004
<u>AP</u>		FRESENIUS KABI USA	<u>1MGM/ML</u>	<u>A076887 001</u>	May 26, 2004
<u>AP</u>		HIKMA FARMACEUTICA	<u>1MGM/ML</u>	<u>A078630 001</u>	May 20, 2009
<u>AP</u>		UNITED BIOMEDCL	<u>1MGM/ML</u>	<u>A200122 001</u>	Nov 08, 2013

TABLET; ORAL

TERBUTALINE SULFATE

<u>AB</u>		IMPAK LABS	<u>2.5MG</u>	<u>A075877 001</u>	Jun 26, 2001
<u>AB</u>			<u>5MG</u>	<u>A075877 002</u>	Jun 26, 2001
<u>AB</u>		LANNETT	<u>2.5MG</u>	<u>A077152 001</u>	Mar 25, 2005
<u>AB</u>	!		<u>5MG</u>	<u>A077152 002</u>	Mar 25, 2005

TERCONAZOLE

CREAM; VAGINAL

TERAZOL 7

<u>AB</u>	+!	JANSSEN PHARMS	<u>0.4%</u>	<u>N019579 001</u>	Dec 31, 1987
		<u>TERCONAZOLE</u>			

<u>AB</u>		FOUGERA PHARMS	<u>0.4%</u>	<u>A076712 001</u>	Feb 18, 2005
<u>AB</u>		TARO	<u>0.4%</u>	<u>A076043 001</u>	Jan 19, 2005
BX	+	NYCOMED US	0.8%	N021735 001	Oct 01, 2004
BX	!	TARO	0.8%	A075953 001	Apr 06, 2004

SUPPOSITORY; VAGINAL

TERAZOL 3

<u>AB</u>	+!	JANSSEN PHARMS	<u>80MG</u>	<u>N019641 001</u>	May 24, 1988
		<u>TERCONAZOLE</u>			

<u>AB</u>		PERRIGO NEW YORK	<u>80MG</u>	<u>A077149 001</u>	Mar 17, 2006
<u>AB</u>		TARO	<u>80MG</u>	<u>A077553 001</u>	Mar 09, 2007

TERIFLUONOMIDE

TABLET; ORAL

AUBAGIO

+	SANOFI AVENTIS US	7MG	N202992 001	Sep 12, 2012
+		14MG	N202992 002	Sep 12, 2012

TERIPARATIDE RECOMBINANT HUMAN

INJECTABLE; SUBCUTANEOUS

FORTEO

+	LILLY	0.6MG/2.4ML (0.25MG/ML)	N021318 002	Jun 25, 2008
---	-------	-------------------------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-404 (of 436)

TESAMORELIN ACETATE

POWDER;SUBCUTANEOUS  
 EGRIFTA  
 +! THERATECHNOLOGIES EQ 1MG BASE/VIAL

N022505 001 Nov 10, 2010

TESTOSTERONE

FILM, EXTENDED RELEASE;TRANSDERMAL  
 ANDRODERM

+! ALLERGAN SALES LLC 2MG/24HR  
 +! 4MG/24HR

N020489 003 Oct 20, 2011  
 N020489 004 Oct 20, 2011

GEL;TRANSDERMAL

ANDROGEL

**AB1** + ABBVIE 25MG/2.5GM PACKET  
**AB1** +! 50MG/5GM PACKET

**N021015 001** Feb 28, 2000  
**N021015 002** Feb 28, 2000

TESTOSTERONE

**AB1** ACTAVIS LABS UT INC 25MG/2.5GM PACKET  
**AB1** 50MG/5GM PACKET  
**AB1** PAR PHARM 25MG/2.5GM PACKET  
**AB1** 50MG/5GM PACKET  
**AB1** PERRIGO ISRAEL 25MG/2.5GM PACKET  
**AB1** 50MG/5GM PACKET

**A076737 001** Jan 27, 2006  
**A076737 002** Jan 27, 2006  
**A076744 001** May 23, 2007  
**A076744 002** May 23, 2007  
**N203098 002** Jan 31, 2013  
**N203098 003** Jan 31, 2013

ANDROGEL

**AB2** + ABBVIE 1.62% (20.25MG/1.25GM PACKET)  
**AB2** +! 1.62% (40.5MG/2.5GM PACKET)

**N022309 002** Sep 07, 2012  
**N022309 003** Sep 07, 2012

TESTIM

**AB2** +! AUXILIUM PHARMS LLC 50MG/5GM PACKET

**N021454 001** Oct 31, 2002

TESTOSTERONE

**AB2** ACTAVIS LABS UT INC 50MG/5GM PACKET  
**AB2** PERRIGO UK FINCO 1.62% (20.25MG/1.25GM PACKET)  
**AB2** 1.62% (40.5MG/2.5GM PACKET)

**A091073 001** Sep 18, 2017  
**A205781 001** Jul 12, 2017  
**A205781 002** Jul 12, 2017

VOGELXO

**AB2** UPSHER-SMITH LABS 50MG/5GM PACKET  
 TESTOSTERONE

**N204399 002** Jun 04, 2014

BX ANI PHARMS INC 25MG/2.5GM PACKET  
 BX 50MG/5GM PACKET  
 GEL, METERED;NASAL  
 NATESTO

N202763 001 Feb 14, 2012  
 N202763 002 Feb 14, 2012

AYTU BIOSCIENCE INC 5.5MG/0.122GM ACTUATION

N205488 001 May 28, 2014

GEL, METERED;TRANSDERMAL

ANDROGEL

**AB** +! ABBVIE 1.62% (20.25MG/1.25GM ACTUATION)  
**AB** +! 12.5MG/1.25GM ACTUATION

**N022309 001** Apr 29, 2011  
**N021015 003** Sep 26, 2003

FORTESTA

**AB** +! ENDO PHARMS 10MG/0.5GM ACTUATION

**N021463 001** Dec 29, 2010

TESTOSTERONE

**AB** ACTAVIS LABS UT INC 10MG/0.5GM ACTUATION  
**AB** 12.5MG/1.25GM ACTUATION  
**AB** PERRIGO ISRAEL 12.5MG/1.25GM ACTUATION  
**AB** 1.62% (20.25MG/1.25GM ACTUATION)

**A204571 001** Aug 05, 2015  
**A076737 003** Mar 09, 2015  
**N203098 001** Jan 31, 2013  
**A204268 001** Aug 04, 2015

VOGELXO

BX UPSHER-SMITH LABS 12.5MG/1.25GM ACTUATION  
 PELLET;IMPLANTATION

N204399 003 Jun 04, 2014

TESTOPEL

! AUXILIUM PHARMS INC 75MG

A080911 001

SOLUTION, METERED;TRANSDERMAL

TESTOSTERONE

**AT** ACTAVIS LABS UT INC 30MG/1.5ML ACTUATION  
**AT** LUPIN LTD 30MG/1.5ML ACTUATION  
**AT** ! PERRIGO ISRAEL 30MG/1.5ML ACTUATION

**A205328 001** Aug 07, 2017  
**A208061 001** Oct 23, 2017  
**A204255 001** Feb 28, 2017

TABLET, EXTENDED RELEASE;BUCCAL

STRIANT

+! AUXILIUM PHARMS LLC 30MG

N021543 001 Jun 19, 2003

TESTOSTERONE CYPIONATE

INJECTABLE;INJECTION

DEPO-TESTOSTERONE

**AO** ! PHARMACIA AND UPJOHN 100MG/ML  
**AO** ! 200MG/ML

**A085635 002**

**A085635 003**

TESTOSTERONE CYPIONATE

**AO** HIKMA FARMACEUTICA 200MG/ML  
**AO** LUITPOLD PHARMS INC 200MG/ML  
**AO** MYLAN INSTITUTIONAL 200MG/ML  
**AO** PADDOCK LLC 200MG/ML

**A091244 001** May 01, 2012  
**A207742 001** Jun 16, 2017  
**A040652 001** Dec 11, 2006  
**A040530 001** Jan 31, 2005

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-405 (of 436)

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

**TESTOSTERONE CYPIONATE**

<u>AO</u>	SANDOZ INC	<u>100MG/ML</u>	<u>A040615 001</u>	Aug 10, 2006
<u>AO</u>		<u>200MG/ML</u>	<u>A040615 002</u>	Aug 10, 2006
<u>AO</u>	SUN PHARM IND LTD	<u>100MG/ML</u>	<u>A201720 001</u>	Jun 03, 2013
<u>AO</u>		<u>200MG/ML</u>	<u>A201720 002</u>	Jun 03, 2013
<u>AO</u>	WATSON PHARMS INC	<u>200MG/ML</u>	<u>A086030 001</u>	
<u>AO</u>	WEST-WARD PHARMS INT	<u>100MG/ML</u>	<u>A090387 001</u>	Jul 15, 2010
<u>AO</u>		<u>200MG/ML</u>	<u>A090387 002</u>	Jul 15, 2010

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

**DELATESTRYL**

<u>AO</u> +!	ENDO PHARMS	<u>200MG/ML</u>	<u>N009165 003</u>	
<u>AO</u>	HIKMA FARMACEUTICA	<u>200MG/ML</u>	<u>A091120 001</u>	Sep 18, 2012
<u>AO</u>	MYLAN INSTITUTIONAL	<u>200MG/ML</u>	<u>A040647 001</u>	Oct 05, 2009
<u>AO</u>	PADDOCK LLC	<u>200MG/ML</u>	<u>A040575 001</u>	Jun 14, 2006
<u>AO</u>	WATSON PHARMS INC	<u>200MG/ML</u>	<u>A085598 001</u>	

TESTOSTERONE UNDECANOATE

INJECTABLE; INTRAMUSCULAR

AVEED

+! ENDO PHARMS INC

750MG/3ML (250MG/ML)

N022219 001 Mar 05, 2014

TETRABENAZINE

TABLET; ORAL

**TETRABENAZINE**

<u>AB</u>	ACTAVIS LABS FL INC	<u>25MG</u>	<u>A206686 001</u>	Jul 07, 2017
<u>AB</u>	APICORE US	<u>12.5MG</u>	<u>A207682 001</u>	Jan 31, 2017
<u>AB</u>		<u>25MG</u>	<u>A207682 002</u>	Jan 31, 2017
<u>AB</u>	BIONPHARMA INC	<u>12.5MG</u>	<u>A208826 001</u>	Dec 18, 2017
<u>AB</u>		<u>25MG</u>	<u>A208826 002</u>	Dec 18, 2017
<u>AB</u>	DR REDDYS LABS LTD	<u>12.5MG</u>	<u>A209284 001</u>	Jan 08, 2018
<u>AB</u>		<u>25MG</u>	<u>A209284 002</u>	Jan 08, 2018
<u>AB</u>	HETERO LABS LTD V	<u>12.5MG</u>	<u>A204574 001</u>	Feb 03, 2016
<u>AB</u>		<u>25MG</u>	<u>A204574 002</u>	Feb 03, 2016
<u>AB</u>	SUN PHARMA GLOBAL	<u>12.5MG</u>	<u>A206129 001</u>	Aug 17, 2015
<u>AB</u>		<u>25MG</u>	<u>A206129 002</u>	Aug 17, 2015
<u>AB</u>	XENAZINE			
<u>AB</u> +	VALEANT PHARMS NORTH	<u>12.5MG</u>	<u>N021894 001</u>	Aug 15, 2008
<u>AB</u> +!		<u>25MG</u>	<u>N021894 002</u>	Aug 15, 2008

TETRACAIN HYDROCHLORIDE

SOLUTION; OPHTHALMIC

TETRACAIN HYDROCHLORIDE

+! NOVARTIS PHARMS CORP

N208135 001 Feb 29, 2016

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

**ACHROMYCIN V**

<u>AB</u> +	HERITAGE PHARMA	<u>250MG</u>	<u>N050278 003</u>	
<u>AB</u> +!		<u>500MG</u>	<u>N050278 001</u>	

**TETRACYCLINE HYDROCHLORIDE**

<u>AB</u>	CHARTWELL TETRA	<u>250MG</u>	<u>A062752 001</u>	Aug 12, 1988
<u>AB</u>		<u>500MG</u>	<u>A062752 002</u>	Aug 12, 1988
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>A061837 001</u>	
<u>AB</u>		<u>500MG</u>	<u>A061837 002</u>	

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION; NASAL

TYZINE

! FOUGERA PHARMS

0.05%

0.1%

A086576 002

A086576 001

SPRAY; NASAL

TYZINE

! FOUGERA PHARMS

0.1%

A086576 003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-406 (of 436)

THALIDOMIDE

CAPSULE; ORAL

THALOMID

+ CELGENE	50MG	N020785 001 Jul 16, 1998
+	100MG	N020785 002 Jan 17, 2003
+	150MG	N020785 004 Jan 10, 2007
+!	200MG	N020785 003 Jan 17, 2003

THALLOUS CHLORIDE TL-201

INJECTABLE; INJECTION

**THALLOUS CHLORIDE TL 201**

<b>AP +!</b>	GE HEALTHCARE	<b>1mCi/ML</b>	<b>N018110 002</b> Feb 27, 1996
<b>AP +!</b>	LANTHEUS MEDCL	<b>1mCi/ML</b>	<b>N017806 001</b>
<b>AP +!</b>	MALLINKRODT NUCLEAR	<b>1mCi/ML</b>	<b>N018150 001</b>

INJECTABLE; INTRAVENOUS

**THALLOUS CHLORIDE TL 201**

<b>AP +!</b>	LANTHEUS MEDCL	<b>2mCi/ML</b>	<b>N017806 002</b> Oct 09, 1998
--------------	----------------	----------------	---------------------------------

THEOPHYLLINE

CAPSULE, EXTENDED RELEASE; ORAL

THEO-24

ACTIENT PHARMS	100MG	A087942 001 Aug 22, 1983
! AUXILIUM PHARMS INC	400MG	A081034 001 Feb 28, 1992
AUXILIUM PHARMS LLC	200MG	A087943 001 Aug 22, 1983
	300MG	A087944 001 Aug 22, 1983

INJECTABLE; INJECTION

**THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER**

<b>AP +!</b>	B BRAUN	<b>40MG/100ML</b>	<b>N019826 001</b> Aug 14, 1992
<b>AP +!</b>	B BRAUN	<b>160MG/100ML</b>	<b>N019826 003</b> Aug 14, 1992

**THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER**

<b>AP +!</b>	B BRAUN	<b>320MG/100ML</b>	<b>N019826 006</b> Aug 14, 1992
<b>AP +!</b>	HOSPIRA INC	<b>40MG/100ML</b>	<b>N019211 001</b> Dec 14, 1984
<b>AP +!</b>		<b>160MG/100ML</b>	<b>N019211 003</b> Dec 14, 1984

<b>AP +!</b>		<b>320MG/100ML</b>	<b>N019211 006</b> Jan 20, 1988
THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER			N019826 002 Aug 14, 1992

+! B BRAUN 80MG/100ML

THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER

+! HOSPIRA INC 4MG/ML

SOLUTION; ORAL

**THEOPHYLLINE**

<b>AA !</b>	SILARK	<b>80MG/15ML</b>	<b>A091156 001</b> Apr 13, 2011
<b>AA</b>	TRIS PHARMA INC	<b>80MG/15ML</b>	<b>A091586 001</b> Jun 15, 2012

SOLUTION, ELIXIR; ORAL

**ELIXOPHYLLIN**

<b>AA !</b>	NOSTRUM LABS INC	<b>80MG/15ML</b>	<b>A085186 001</b>
-------------	------------------	------------------	--------------------

**THEOPHYLLINE**

<b>AA</b>	PHARM ASSOC	<b>80MG/15ML</b>	<b>A206344 001</b> Dec 16, 2016
-----------	-------------	------------------	---------------------------------

TABLET, EXTENDED RELEASE; ORAL

**THEOPHYLLINE**

<b>AB</b>	ALEMBIC PHARMS LTD	<b>300MG</b>	<b>A090430 001</b> Oct 27, 2010
<b>AB</b>	GLENMARK GENERICS	<b>400MG</b>	<b>A090355 001</b> Jul 13, 2010

**600MG**

<b>AB</b>	MYLAN IRELAND LTD	<b>400MG</b>	<b>A090355 002</b> Jul 13, 2010
<b>AB</b>		<b>600MG</b>	<b>A040560 003</b> Apr 21, 2006

**600MG**

<b>AB !</b>	PLIVA	<b>100MG</b>	<b>A040560 002</b> Apr 21, 2006
<b>AB !</b>		<b>200MG</b>	<b>A089807 001</b> Apr 30, 1990

**300MG**

<b>AB</b>	RHODES PHARMS	<b>400MG</b>	<b>A089763 001</b> Apr 30, 1990
<b>AB</b>		<b>600MG</b>	<b>A087571 001</b> Sep 01, 1982

**600MG**

THEOCHRON

NOSTRUM PHARMS LLC 100MG

A087400 003 Feb 21, 1985

200MG

A087400 004 Feb 21, 1985

THEOPHYLLINE

! ALEMBIC PHARMS LTD 450MG

A090430 002 Oct 27, 2010

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

**THIAMINE HYDROCHLORIDE**

<b>AP !</b>	FRESENIUS KABI USA	<b>100MG/ML</b>	<b>A080556 001</b>
<b>AP</b>	MYLAN INSTITUTIONAL	<b>100MG/ML</b>	<b>A091623 001</b> Jun 25, 2012

<b>AP</b>	SAGENT PHARMS	<b>100MG/ML</b>	<b>A206106 001</b> Dec 01, 2017
-----------	---------------	-----------------	---------------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-407 (of 436)

THIOGUANINE

TABLET;ORAL  
 THIOGUANINE  
 +! ASPEN GLOBAL INC 40MG N012429 001

THIORIDAZINE HYDROCHLORIDE

TABLET;ORAL  
THIORIDAZINE HYDROCHLORIDE  
AB MYLAN 10MG A088004 002 Mar 15, 1983  
AB 25MG A088004 003 Mar 15, 1983  
AB 50MG A088004 004 Mar 15, 1983  
AB ! 100MG A088004 001 Nov 18, 1983  
AB SUN PHARM INDUSTRIES 10MG A089953 004 Aug 01, 1986  
AB 25MG A089953 003 Aug 01, 1986  
AB 50MG A089953 002 Aug 01, 1986  
AB 100MG A089953 001 Oct 07, 1988

THIOTEP A

INJECTABLE;INJECTION  
 THIOTEP A  
 ! WEST-WARD PHARMS 15MG/VIAL A075547 001 Apr 02, 2001  
 INT  
 POWDER;IV (INFUSION)  
 TEPADINA  
 +! ADIENNE SA 15MG/VIAL N208264 001 Jan 26, 2017  
 +! 100MG/VIAL N208264 002 Jan 26, 2017

THIOTHIXENE

CAPSULE;ORAL  
 THIOTHIXENE  
 MYLAN 1MG A071093 002 Jun 23, 1987  
 2MG A071093 003 Jun 23, 1987  
 ! 5MG A071093 004 Jun 23, 1987  
 10MG A071093 001 Jun 23, 1987

THYROTROPIN ALFA

INJECTABLE;INJECTION  
 THYROID  
 +! GENZYME 1.1MG/VIAL N020898 001 Nov 30, 1998

TIAGABINE HYDROCHLORIDE

TABLET;ORAL  
GABITRIL  
AB + CEPHALON 2MG N020646 005 Apr 16, 1999  
AB +! 4MG N020646 001 Sep 30, 1997  
AB + 12MG N020646 002 Sep 30, 1997  
AB + 16MG N020646 003 Sep 30, 1997  
TIAGABINE HYDROCHLORIDE  
AB AMNEAL PHARMS CO 2MG A208181 001 Dec 08, 2017  
AB 4MG A208181 002 Dec 08, 2017  
AB 12MG A208181 003 Dec 08, 2017  
AB 16MG A208181 004 Dec 08, 2017  
AB SUN PHARM IND S 2MG A077555 001 Nov 04, 2011  
AB 4MG A077555 002 Nov 04, 2011  
AB WILSHIRE PHARMS INC 2MG A206857 001 Oct 13, 2017  
AB 4MG A206857 002 Oct 13, 2017  
AB 12MG A206857 003 Oct 13, 2017  
AB 16MG A206857 004 Oct 13, 2017

TICAGRELOR

TABLET;ORAL  
 BRILINTA  
 + ASTRAZENECA PHARMS 60MG N022433 002 Sep 03, 2015  
 +! 90MG N022433 001 Jul 20, 2011

TICLOPIDINE HYDROCHLORIDE

TABLET;ORAL  
TICLOPIDINE HYDROCHLORIDE  
AB APOTEX 250MG A075089 001 Jul 01, 1999  
AB SUN PHARM IND S INC 250MG A075526 001 Sep 26, 2002  
AB ! TEVA 250MG A075149 001 Aug 20, 1999

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-408 (of 436)

**TIGECYCLINE**

POWDER;IV (INFUSION)

**TIGECYCLINE**

<b>AP</b>	FRESENIUS KABI USA	<b>50MG/VIAL</b>	<b>N205645 001</b>	Dec 01, 2016
<b>AP</b>	SANDOZ INC	<b>50MG/VIAL</b>	<b>A091620 001</b>	May 27, 2015

**TYGACIL**

<b>AP</b>	+! PF PRISM CV	<b>50MG/VIAL</b>	<b>N021821 001</b>	Jun 15, 2005
-----------	----------------	------------------	--------------------	--------------

**TIMOLOL**

SOLUTION/DROPS;OPHTHALMIC

**BETIMOL**

<b>AT</b>	+! OAK PHARMS INC	<b>EQ 0.25% BASE</b>	<b>N020439 001</b>	Mar 31, 1995
<b>AT</b>	+!	<b>EQ 0.5% BASE</b>	<b>N020439 002</b>	Mar 31, 1995

**TIMOLOL**

<b>AT</b>	AKORN	<b>EQ 0.25% BASE</b>	<b>A205309 001</b>	Sep 30, 2016
<b>AT</b>		<b>EQ 0.5% BASE</b>	<b>A205309 002</b>	Sep 30, 2016

**TIMOLOL MALEATE**

SOLUTION, GEL FORMING/DROPS;OPHTHALMIC

**TIMOLOL MALEATE**

<b>AB</b>	SANDOZ INC	<b>EQ 0.25% BASE</b>	<b>N020963 001</b>	Oct 21, 1998
<b>AB</b>		<b>EQ 0.5% BASE</b>	<b>N020963 002</b>	Oct 21, 1998

**TIMOPTIC-XE**

<b>AB</b>	+! VALEANT PHARMS LLC	<b>EQ 0.25% BASE</b>	<b>N020330 001</b>	Nov 04, 1993
<b>AB</b>	+!	<b>EQ 0.5% BASE</b>	<b>N020330 002</b>	Nov 04, 1993

SOLUTION/DROPS;OPHTHALMIC

**TIMOLOL MALEATE**

<b>AT</b>	BAUSCH AND LOMB	<b>EQ 0.25% BASE</b>	<b>A074778 001</b>	Mar 25, 1997
<b>AT</b>	FDC LTD	<b>EQ 0.25% BASE</b>	<b>A077259 001</b>	Apr 30, 2008
<b>AT</b>	PACIFIC PHARMA	<b>EQ 0.25% BASE</b>	<b>A074746 001</b>	Mar 25, 1997
<b>AT</b>	! SANDOZ INC	<b>EQ 0.25% BASE</b>	<b>A074261 001</b>	Apr 28, 1995
<b>AT</b>	WOCKHARDT	<b>EQ 0.25% BASE</b>	<b>A078771 001</b>	Sep 28, 2009
<b>AT1</b>	AKORN	<b>EQ 0.5% BASE</b>	<b>A074466 001</b>	Mar 25, 1997
<b>AT1</b>		<b>EQ 0.5% BASE</b>	<b>A074516 001</b>	Mar 25, 1997
<b>AT1</b>	BAUSCH AND LOMB	<b>EQ 0.5% BASE</b>	<b>A074776 001</b>	Mar 25, 1997
<b>AT1</b>	FDC LTD	<b>EQ 0.5% BASE</b>	<b>A077259 002</b>	Apr 30, 2008
<b>AT1</b>	HI TECH PHARMA	<b>EQ 0.5% BASE</b>	<b>A075163 001</b>	Sep 10, 2002
<b>AT1</b>	PACIFIC PHARMA	<b>EQ 0.5% BASE</b>	<b>A074747 001</b>	Mar 25, 1997
<b>AT1</b>	! SANDOZ INC	<b>EQ 0.5% BASE</b>	<b>A074262 001</b>	Apr 28, 1995
<b>AT1</b>	WOCKHARDT	<b>EQ 0.5% BASE</b>	<b>A078771 002</b>	Sep 28, 2009

**ISTALOL**

<b>AT2</b>	+! BAUSCH AND LOMB	<b>EQ 0.5% BASE</b>	<b>N021516 001</b>	Jun 04, 2004
------------	--------------------	---------------------	--------------------	--------------

**TIMOLOL MALEATE**

<b>AT2</b>	APOTEX INC	<b>EQ 0.5% BASE</b>	<b>A204936 001</b>	Apr 17, 2015
	TIMOPTIC IN OCUDOSE			
	+! ATON	EQ 0.25% BASE	N019463 001	Nov 05, 1986

+!

EQ 0.5% BASE

N019463 002 Nov 05, 1986

+!

EQ 0.5% BASE

N019463 001 Jun 08, 1990

TABLET;ORAL

TIMOLOL MALEATE

MYLAN	5MG	A072668 002	Jun 08, 1990
	10MG	A072668 003	Jun 08, 1990
!	20MG	A072668 001	Jun 08, 1990

**TINIDAZOLE**

TABLET;ORAL

**TINDAMAX**

<b>AB</b>	+ MISSION PHARMA	<b>250MG</b>	<b>N021618 001</b>	May 17, 2004
<b>AB</b>	+!	<b>500MG</b>	<b>N021618 002</b>	May 17, 2004

**TINIDAZOLE**

<b>AB</b>	EDENBRIDGE PHARMS	<b>250MG</b>	<b>A203808 001</b>	Aug 04, 2015
<b>AB</b>		<b>500MG</b>	<b>A203808 002</b>	Aug 04, 2015
<b>AB</b>	NOVEL LABS INC	<b>250MG</b>	<b>A202044 001</b>	Apr 30, 2012
<b>AB</b>		<b>500MG</b>	<b>A202044 002</b>	Apr 30, 2012
<b>AB</b>	UNIQUE PHARM LABS	<b>250MG</b>	<b>A202489 001</b>	Oct 09, 2013
<b>AB</b>		<b>500MG</b>	<b>A202489 002</b>	Oct 09, 2013
<b>AB</b>	WEST-WARD PHARMS INT	<b>250MG</b>	<b>A201172 001</b>	Apr 30, 2012
<b>AB</b>		<b>500MG</b>	<b>A201172 002</b>	Apr 30, 2012

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-409 (of 436)

TIOPRONIN

TABLET;ORAL  
 THIOLA  
 +! MISSION PHARMA 100MG N019569 001 Aug 11, 1988

TIOTROPIUM BROMIDE

POWDER;INHALATION  
 SPIRIVA  
 +! BOEHRINGER INGELHEIM EQ 0.018MG BASE/INH N021395 001 Jan 30, 2004  
 SPRAY, METERED;INHALATION  
 SPIRIVA RESPIMAT  
 + BOEHRINGER INGELHEIM EQ 0.00125MG BASE/INH N021936 002 Sep 15, 2015  
 +! EQ 0.0025MG BASE/INH N021936 001 Sep 24, 2014

TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE

TABLET;ORAL  
 LONSURF  
 + TAIHO ONCOLOGY EQ 6.14MG BASE;15MG N207981 001 Sep 22, 2015  
 +! EQ 8.19MG BASE;20MG N207981 002 Sep 22, 2015

TIPRANAVIR

CAPSULE;ORAL  
 APTIVUS  
 +! BOEHRINGER INGELHEIM 250MG N021814 001 Jun 22, 2005  
 SOLUTION;ORAL  
 APTIVUS  
 +! BOEHRINGER INGELHEIM 100MG/ML N022292 001 Jun 23, 2008

TIROFIBAN HYDROCHLORIDE

INJECTABLE;INJECTION  
 AGGRASTAT  
 + MEDICURE EQ 5MG BASE/100ML (EQ 0.05MG BASE/ML) N020913 002 May 17, 2002  
 +! EQ 12.5MG BASE/250ML (EQ 0.05MG BASE/ML) N020913 003 Apr 20, 2000  
 SOLUTION;INJECTION  
 AGGRASTAT  
 +! MEDICURE EQ 3.75MG BASE/15ML (EQ 0.25MG BASE/ML) N020912 002 Aug 31, 2016

TIZANIDINE HYDROCHLORIDE

CAPSULE;ORAL  
TIZANIDINE HYDROCHLORIDE  
AB APOTEX INC EQ 2MG BASE A078868 001 Feb 03, 2012  
AB EQ 4MG BASE A078868 002 Feb 03, 2012  
AB EQ 6MG BASE A078868 003 Feb 03, 2012  
AB JUBILANT GENERICS EQ 2MG BASE A209605 001 Aug 04, 2017  
AB EQ 4MG BASE A209605 002 Aug 04, 2017  
AB EQ 6MG BASE A209605 003 Aug 04, 2017  
AB MYLAN PHARMS INC EQ 2MG BASE A091502 001 Nov 09, 2012  
AB EQ 4MG BASE A091502 002 Nov 09, 2012  
AB EQ 6MG BASE A091502 003 Nov 09, 2012  
AB PAR PHARM INC EQ 2MG BASE A207199 001 Mar 14, 2017  
AB EQ 4MG BASE A207199 002 Mar 14, 2017  
AB EQ 6MG BASE A207199 003 Mar 14, 2017  
AB ZYDUS PHARMS USA INC EQ 2MG BASE A208622 001 Mar 03, 2017  
AB EQ 4MG BASE A208622 002 Mar 03, 2017  
AB EQ 6MG BASE A208622 003 Mar 03, 2017

ZANAFLEX

AB + COVIS PHARMA BV EQ 2MG BASE N021447 001 Aug 29, 2002  
AB + EQ 4MG BASE N021447 002 Aug 29, 2002  
AB +! EQ 6MG BASE N021447 003 Aug 29, 2002

TABLET;ORAL

TIZANIDINE HYDROCHLORIDE

AB APOTEX EQ 2MG BASE A076533 001 Jan 16, 2004  
AB EQ 4MG BASE A076533 002 Jan 16, 2004  
AB DR REDDYS LABS INC EQ 2MG BASE A076286 001 Jul 03, 2002  
AB EQ 4MG BASE A076286 002 Jul 03, 2002  
AB EPIC PHARMA LLC EQ 2MG BASE A076347 001 Oct 11, 2002  
AB EQ 4MG BASE A076347 002 Oct 11, 2002  
AB MYLAN EQ 2MG BASE A076354 001 Mar 28, 2003  
AB EQ 4MG BASE A076354 002 Mar 28, 2003  
AB OXFORD PHARMS EQ 2MG BASE A076281 001 Oct 20, 2003

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-410 (of 436)

TIZANIDINE HYDROCHLORIDE

TABLET;ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076281 002</u>	Oct 20, 2003
<u>AB</u>	PAR PHARM INC	<u>EQ 2MG BASE</u>	<u>A207170 001</u>	Jan 26, 2017
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A207170 002</u>	Jan 26, 2017
<u>AB</u>	SANDOZ INC	<u>EQ 2MG BASE</u>	<u>A076280 001</u>	Nov 26, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076280 002</u>	Jun 27, 2002
<u>AB</u>	SUN PHARM INDS INC	<u>EQ 2MG BASE</u>	<u>A076416 001</u>	Sep 29, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076416 002</u>	Sep 29, 2003
<u>AB</u>	TEVA	<u>EQ 2MG BASE</u>	<u>A076284 001</u>	Jul 03, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A076284 002</u>	Jul 03, 2002
<u>AB</u>	UNICHEM LABS LTD	<u>EQ 2MG BASE</u>	<u>A091283 001</u>	Nov 28, 2012
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>A091283 002</u>	Nov 28, 2012

ZANAFLEX

<u>AB</u>	+!	COVIS PHARMA BV	<u>EQ 4MG BASE</u>	<u>N020397 001</u>	Nov 27, 1996
-----------	----	-----------------	--------------------	--------------------	--------------

TOBRAMYCIN

OINTMENT;OPHTHALMIC

TOBREX

+! NOVARTIS PHARMS CORP

POWDER;INHALATION

TOBI PODHALER

+! NOVARTIS

28MG

N201688 001 Mar 22, 2013

SOLUTION;INHALATION

KITABIS PAK

<u>AN</u>	PULMOFLOW INC	<u>300MG/5ML</u>	<u>N205433 001</u>	Dec 02, 2014
-----------	---------------	------------------	--------------------	--------------

TOBI

<u>AN</u>	+!	NOVARTIS PHARMS	<u>300MG/5ML</u>	<u>N050753 001</u>	Dec 22, 1997
-----------	----	-----------------	------------------	--------------------	--------------

TOBRAMYCIN

<u>AN</u>	AKORN INC	<u>300MG/5ML</u>	<u>A201422 001</u>	May 28, 2014
-----------	-----------	------------------	--------------------	--------------

<u>AN</u>	AMNEAL PHARMS	<u>300MG/5ML</u>	<u>A205501 001</u>	Jul 13, 2015
-----------	---------------	------------------	--------------------	--------------

<u>AN</u>	LUPIN ATLANTIS	<u>300MG/5ML</u>	<u>A208964 001</u>	Mar 22, 2017
-----------	----------------	------------------	--------------------	--------------

<u>AN</u>	MYLAN PHARMS INC	<u>300MG/5ML</u>	<u>A209554 001</u>	Oct 13, 2017
-----------	------------------	------------------	--------------------	--------------

<u>AN</u>	TEVA PHARMS USA	<u>300MG/5ML</u>	<u>A091589 001</u>	Oct 10, 2013
-----------	-----------------	------------------	--------------------	--------------

BETHKIS

+! CHIESI USA INC

300MG/4ML

N201820 001 Oct 12, 2012

SOLUTION/DROPS;OPHTHALMIC

AKTOB

<u>AT</u>	AKORN	<u>0.3%</u>	<u>A064096 001</u>	Jan 31, 1996
-----------	-------	-------------	--------------------	--------------

TOBRAMYCIN

<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>A064052 001</u>	Nov 29, 1993
-----------	-----------------	-------------	--------------------	--------------

<u>AT</u>	FERA PHARMS	<u>0.3%</u>	<u>A065026 001</u>	Sep 11, 2001
-----------	-------------	-------------	--------------------	--------------

<u>AT</u>	SOMERSET THERAPS LLC	<u>0.3%</u>	<u>A207444 001</u>	Jun 28, 2017
-----------	----------------------	-------------	--------------------	--------------

TOBREX

<u>AT</u>	+!	NOVARTIS PHARMS CORP	<u>0.3%</u>	<u>N050541 001</u>
-----------	----	----------------------	-------------	--------------------

<u>AT</u>	SANDOZ INC	<u>0.3%</u>	<u>A062535 001</u>	Dec 13, 1984
-----------	------------	-------------	--------------------	--------------

TOBRAMYCIN SULFATE

INJECTABLE;INJECTION

TOBRAMYCIN SULFATE

<u>AP</u>	AKORN	<u>EQ 40MG BASE/ML</u>	<u>A205179 001</u>	Sep 16, 2014
-----------	-------	------------------------	--------------------	--------------

<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 40MG BASE/ML</u>	<u>A206965 001</u>	Jul 01, 2016
-----------	----------------------	------------------------	--------------------	--------------

<u>AP</u>	FRESENIUS KABI USA	<u>EQ 10MG BASE/ML</u>	<u>A065122 001</u>	Nov 29, 2002
-----------	--------------------	------------------------	--------------------	--------------

<u>AP</u>	!	<u>EQ 40MG BASE/ML</u>	<u>A065122 002</u>	Nov 29, 2002
-----------	---	------------------------	--------------------	--------------

<u>AP</u>	!	<u>EQ 1.2GM BASE/VIAL</u>	<u>N050789 001</u>	Jul 13, 2004
-----------	---	---------------------------	--------------------	--------------

<u>AP</u>	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>A063112 001</u>	Apr 30, 1991
-----------	---------	------------------------	--------------------	--------------

<u>AP</u>	!	<u>EQ 40MG BASE/ML</u>	<u>A065407 001</u>	Mar 11, 2008
-----------	---	------------------------	--------------------	--------------

<u>AP</u>	MYLAN LABS LTD	<u>EQ 40MG BASE/ML</u>	<u>A063100 001</u>	Jan 30, 1992
-----------	----------------	------------------------	--------------------	--------------

<u>AP</u>	TEVA PHARMS USA	<u>EQ 40MG BASE/ML</u>	<u>A065120 001</u>	Nov 29, 2002
-----------	-----------------	------------------------	--------------------	--------------

<u>AP</u>	WEST-WARD PHARMS INT	<u>EQ 40MG BASE/ML</u>	<u>A063117 001</u>	Apr 26, 1991
-----------	----------------------	------------------------	--------------------	--------------

<u>AP</u>	!	<u>EQ 1.2GM BASE/VIAL</u>	<u>A065013 001</u>	Aug 17, 2001
-----------	---	---------------------------	--------------------	--------------

<u>AP</u>	X GEN PHARMS	<u>EQ 1.2GM BASE/VIAL</u>	<u>A205685 001</u>	Sep 16, 2014
-----------	--------------	---------------------------	--------------------	--------------

TOBRAMYCIN SULFATE (PHARMACY BULK)

! FRESENIUS KABI USA EQ 40MG BASE/ML

A065120 001 Nov 29, 2002

TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

! HOSPIRA EQ 1.2MG BASE/ML

A063081 003 Jul 31, 1990

! EQ 1.6MG BASE/ML

A063081 006 Jun 02, 1993

! EQ 80MG BASE/100ML

A063081 001 Jul 31, 1990

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-411 (of 436)

TOFACITINIB CITRATE

TABLET;ORAL	XELJANZ	+! PF PRISM CV	EQ 5MG BASE	N203214 001 Nov 06, 2012
TABLET, EXTENDED RELEASE;ORAL	XELJANZ XR	+! PFIZER INC	EQ 11MG BASE	N208246 001 Feb 23, 2016

TOLAZAMIDE

TABLET;ORAL	TOLAZAMIDE	MYLAN PHARMS INC	250MG	A070259 001 Jan 02, 1986
	!		500MG	A070259 003 Mar 17, 1986

TOLBUTAMIDE

TABLET;ORAL	TOLBUTAMIDE	! MYLAN PHARMS INC	500MG	A086445 001
-------------	-------------	--------------------	-------	-------------

TOLCAPONE

TABLET;ORAL	<b>TASMAR</b>	<b>AB +! VALEANT PHARMS LLC</b>	<b>100MG</b>	<b>N020697 001</b> Jan 29, 1998
	<b>TOLCAPONE</b>	<b>AB PAR PHARM INC</b>	<b>100MG</b>	<b>A204584 001</b> Mar 26, 2015

TOLMETIN SODIUM

CAPSULE;ORAL	<b>TOLMETIN SODIUM</b>	<b>AB MYLAN</b>	<b>EQ 400MG BASE</b>	<b>A073393 001</b> May 27, 1993
		<b>AB ! TEVA</b>	<b>EQ 400MG BASE</b>	<b>A073290 001</b> Nov 27, 1991
TABLET;ORAL	TOLMETIN SODIUM	! MYLAN	EQ 600MG BASE	A074473 001 Aug 30, 1994

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL	<b>DETROL LA</b>	<b>AB + PHARMACIA AND UPJOHN</b>	<b>2MG</b>	<b>N021228 001</b> Dec 22, 2000
		<b>AB +! TEVA</b>	<b>4MG</b>	<b>N021228 002</b> Dec 22, 2000

TOLTERODINE TARTRATE

HETERO LABS LTD III	<b>AB 2MG</b>	<b>A206419 001</b> Dec 12, 2017
	<b>AB 4MG</b>	<b>A206419 002</b> Dec 12, 2017
MYLAN PHARMS INC	<b>AB 2MG</b>	<b>A201486 001</b> Oct 31, 2013
	<b>AB 4MG</b>	<b>A201486 002</b> Oct 31, 2013
TEVA PHARMS USA	<b>AB 2MG</b>	<b>A079141 001</b> Nov 22, 2016
	<b>AB 4MG</b>	<b>A079141 002</b> Nov 22, 2016
TORRENT PHARMS LTD	<b>AB 2MG</b>	<b>A203016 001</b> Aug 11, 2015
	<b>AB 4MG</b>	<b>A203016 002</b> Aug 11, 2015

TABLET;ORAL

<b>DETROL</b>	<b>AB + PHARMACIA AND UPJOHN</b>	<b>1MG</b>	<b>N020771 001</b> Mar 25, 1998
	<b>AB +! TEVA</b>	<b>2MG</b>	<b>N020771 002</b> Mar 25, 1998

TOLTERODINE TARTRATE

APOTEX CORP	<b>AB 1MG</b>	<b>A200164 001</b> Sep 25, 2012
	<b>AB 2MG</b>	<b>A200164 002</b> Sep 25, 2012
IVAX SUB TEVA PHARMS	<b>AB 1MG</b>	<b>A077006 001</b> Feb 23, 2015
	<b>AB 2MG</b>	<b>A077006 002</b> Feb 23, 2015
MACLEODS PHARMS LTD	<b>AB 1MG</b>	<b>A203409 001</b> Aug 31, 2015
	<b>AB 2MG</b>	<b>A203409 002</b> Aug 31, 2015
MYLAN PHARMS INC	<b>AB 1MG</b>	<b>A202641 001</b> Nov 27, 2012
	<b>AB 2MG</b>	<b>A202641 002</b> Nov 27, 2012

TOLVAPTAN

TABLET;ORAL	SAMSCA	+ OTSUKA AMERICA PHARM	15MG	N022275 001 May 19, 2009
		+!	30MG	N022275 002 May 19, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-412 (of 436)

TOPIRAMATE

CAPSULE; ORAL

TOPAMAX

<u>AB</u>	+	JANSSEN PHARMS	<u>15MG</u>	<u>N020844 001</u>	Oct 26, 1998
<u>AB</u>	+!		<u>25MG</u>	<u>N020844 002</u>	Oct 26, 1998

TOPIRAMATE

<u>AB</u>		TEVA	<u>15MG</u>	<u>A076575 001</u>	Apr 17, 2009
<u>AB</u>			<u>25MG</u>	<u>A076575 002</u>	Apr 17, 2009
<u>AB</u>		WATSON LABS	<u>15MG</u>	<u>A077868 001</u>	Apr 15, 2009
<u>AB</u>			<u>25MG</u>	<u>A077868 002</u>	Apr 15, 2009
<u>AB</u>		ZYDUS PHARMS USA INC	<u>15MG</u>	<u>A078877 001</u>	Oct 14, 2009
<u>AB</u>			<u>25MG</u>	<u>A078877 002</u>	Oct 14, 2009

CAPSULE, EXTENDED RELEASE; ORAL

TOPIRAMATE

<u>AB</u>		ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A207382 001</u>	Nov 24, 2017
<u>AB</u>			<u>50MG</u>	<u>A207382 002</u>	Nov 24, 2017
<u>AB</u>			<u>100MG</u>	<u>A207382 003</u>	Nov 24, 2017

TROKENDI XR

<u>AB</u>	+	SUPERNUS PHARMS	<u>25MG</u>	<u>N201635 001</u>	Aug 16, 2013
<u>AB</u>	+		<u>50MG</u>	<u>N201635 002</u>	Aug 16, 2013
<u>AB</u>	+		<u>100MG</u>	<u>N201635 003</u>	Aug 16, 2013

QUDEXY XR

+	UPSHER-SMITH LABS	25MG	N205122 001	Mar 11, 2014
+		50MG	N205122 002	Mar 11, 2014
+		100MG	N205122 003	Mar 11, 2014
+		150MG	N205122 004	Mar 11, 2014
+		200MG	N205122 005	Mar 11, 2014

TROKENDI XR

+	SUPERNUS PHARMS	200MG	N201635 004	Aug 16, 2013
---	-----------------	-------	-------------	--------------

TABLET; ORAL

TOPAMAX

<u>AB</u>	+	JANSSEN PHARMS	<u>25MG</u>	<u>N020505 004</u>	Dec 24, 1996
<u>AB</u>	+		<u>50MG</u>	<u>N020505 005</u>	Dec 24, 1996
<u>AB</u>	+!		<u>100MG</u>	<u>N020505 001</u>	Dec 24, 1996
<u>AB</u>	+		<u>200MG</u>	<u>N020505 002</u>	Dec 24, 1996

TOPIRAMATE

<u>AB</u>		ACCORD HLTHCARE	<u>25MG</u>	<u>A076311 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A076311 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076311 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076311 004</u>	Mar 27, 2009
<u>AB</u>		APOTEX INC	<u>25MG</u>	<u>A077733 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A077733 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A077733 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A077733 004</u>	Mar 27, 2009
<u>AB</u>		AUROBINDO PHARMA	<u>25MG</u>	<u>A078462 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A078462 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A078462 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A078462 004</u>	Mar 27, 2009
<u>AB</u>		CIPLA LTD	<u>25MG</u>	<u>A076343 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A076343 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076343 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076343 004</u>	Mar 27, 2009
<u>AB</u>		GLENMARK GENERICS	<u>25MG</u>	<u>A077627 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A077627 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A077627 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A077627 004</u>	Mar 27, 2009
<u>AB</u>		INVAGEN PHARMS	<u>25MG</u>	<u>A079162 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A079162 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A079162 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A079162 004</u>	Mar 27, 2009
<u>AB</u>		LUPIN	<u>25MG</u>	<u>A078410 001</u>	Sep 11, 2013
<u>AB</u>			<u>50MG</u>	<u>A078410 002</u>	Sep 11, 2013
<u>AB</u>			<u>100MG</u>	<u>A078410 003</u>	Sep 11, 2013
<u>AB</u>			<u>200MG</u>	<u>A078410 004</u>	Sep 11, 2013
<u>AB</u>		SUN PHARM INDs LTD	<u>25MG</u>	<u>A076327 001</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A076327 002</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A076327 003</u>	Mar 27, 2009
<u>AB</u>		SUN PHARMA GLOBAL	<u>25MG</u>	<u>A090278 001</u>	Mar 27, 2009
<u>AB</u>			<u>50MG</u>	<u>A090278 002</u>	Mar 27, 2009
<u>AB</u>			<u>100MG</u>	<u>A090278 003</u>	Mar 27, 2009
<u>AB</u>			<u>200MG</u>	<u>A090278 004</u>	Mar 27, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-413 (of 436)

TOPIRAMATE

TABLET;ORAL

TOPIRAMATE

<u>AB</u>	TEVA	<u>25MG</u>	<u>A076317 001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A076317 002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A076317 003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A076317 004</u>	Mar 27, 2009
<u>AB</u>	TORRENT PHARMS	<u>25MG</u>	<u>A079153 001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A079153 002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A079153 003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A079153 004</u>	Mar 27, 2009
<u>AB</u>	UNICHEM LABS LTD	<u>25MG</u>	<u>A090162 001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A090162 002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A090162 003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A090162 004</u>	Feb 19, 2013
<u>AB</u>	UPSHER-SMITH LABS	<u>25MG</u>	<u>A078499 001</u>	Jan 07, 2010
<u>AB</u>		<u>50MG</u>	<u>A078499 002</u>	Jan 07, 2010
<u>AB</u>		<u>100MG</u>	<u>A078499 003</u>	Jan 07, 2010
<u>AB</u>		<u>200MG</u>	<u>A078499 004</u>	Jan 07, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>25MG</u>	<u>A078235 001</u>	Mar 27, 2009
<u>AB</u>		<u>50MG</u>	<u>A078235 002</u>	Mar 27, 2009
<u>AB</u>		<u>100MG</u>	<u>A078235 003</u>	Mar 27, 2009
<u>AB</u>		<u>200MG</u>	<u>A078235 004</u>	Mar 27, 2009

TOPOTECAN HYDROCHLORIDE

CAPSULE;ORAL

HYCAMTIN

+ NOVARTIS PHARMS CORP	EQ 0.25MG BASE	N020981 001	Oct 11, 2007
+!	EQ 1MG BASE	N020981 002	Oct 11, 2007

INJECTABLE;INJECTION

HYCAMTIN

<u>AP</u> +! NOVARTIS PHARMS CORP	<u>EQ 4MG BASE/VIAL</u>	<u>N020671 001</u>	May 28, 1996
-----------------------------------	-------------------------	--------------------	--------------

TOPOTECAN HYDROCHLORIDE

<u>AP</u> ACCORD HLTHCARE	<u>EQ 4MG BASE/VIAL</u>	<u>A202351 001</u>	Jun 26, 2013
<u>AP</u> ACTAVIS TOTOWA	<u>EQ 4MG BASE/VIAL</u>	<u>A090620 001</u>	Dec 02, 2010
<u>AP</u> CIPLA LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A091199 001</u>	Dec 01, 2010
<u>AP</u> DR REDDYS LABS LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A201191 001</u>	Mar 09, 2011
<u>AP</u> FRESENIUS KABI USA	<u>EQ 4MG BASE/VIAL</u>	<u>A091089 001</u>	Nov 29, 2010
<u>AP</u> HONG KONG	<u>EQ 4MG BASE/VIAL</u>	<u>A201166 001</u>	Aug 08, 2012
<u>AP</u> MYLAN LABS LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A091542 001</u>	Aug 28, 2012
<u>AP</u> NOVAST LABS LTD	<u>EQ 4MG BASE/VIAL</u>	<u>A206962 001</u>	Nov 30, 2016
<u>AP</u> SAGENT PHARMS	<u>EQ 4MG BASE/VIAL</u>	<u>A091284 001</u>	Jan 26, 2011

SOLUTION;INTRAVENOUS

TOPOTECAN HYDROCHLORIDE

<u>AP</u> ACCORD HLTHCARE	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A204406 002</u>	Jul 06, 2017
<u>AP</u> +! HOSPIRA INC	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N200582 001</u>	Feb 02, 2011
<u>AP</u> MYLAN LABS LTD	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>A206074 001</u>	Nov 24, 2017
<u>AP</u> TEVA PHARMS USA	<u>EQ 4MG BASE/4ML (EQ 1MG BASE/ML)</u>	<u>N022453 001</u>	Dec 20, 2012
ACCORD HLTHCARE	EQ 1MG BASE/ML (EQ 1MG BASE/ML)	A204406 001	Jul 06, 2017

TOREMIFENE CITRATE

TABLET;ORAL

FARESTON

+! KYOWA KIRIN	EQ 60MG BASE	N020497 001	May 29, 1997
----------------	--------------	-------------	--------------

TORSEMIDE

TABLET;ORAL

DEMADEX

<u>AB</u> + MYLAN SPECIALITY LP	<u>5MG</u>	<u>N020136 001</u>	Aug 23, 1993
<u>AB</u> +	<u>10MG</u>	<u>N020136 002</u>	Aug 23, 1993
<u>AB</u> +!	<u>20MG</u>	<u>N020136 003</u>	Aug 23, 1993
<u>AB</u> +	<u>100MG</u>	<u>N020136 004</u>	Aug 23, 1993

TORSEMIDE

<u>AB</u> APOTEX INC	<u>5MG</u>	<u>A076894 001</u>	May 31, 2005
<u>AB</u>	<u>10MG</u>	<u>A076894 002</u>	May 31, 2005
<u>AB</u>	<u>20MG</u>	<u>A076894 003</u>	May 31, 2005
<u>AB</u>	<u>100MG</u>	<u>A076894 004</u>	May 31, 2005
<u>AB</u> AUROBINDO PHARMA	<u>5MG</u>	<u>A078249 001</u>	Oct 17, 2007
<u>AB</u>	<u>10MG</u>	<u>A078249 002</u>	Oct 17, 2007
<u>AB</u>	<u>20MG</u>	<u>A078249 003</u>	Oct 17, 2007
<u>AB</u>	<u>100MG</u>	<u>A078249 004</u>	Oct 17, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-414 (of 436)

**TORSEMIDE**

TABLET;ORAL

**TORSEMIDE**

<b>AB</b>	HETERO LABS LTD III	<b>5MG</b>	<b>A079234 001</b>	Jan 27, 2009
<b>AB</b>		<b>10MG</b>	<b>A079234 002</b>	Jan 27, 2009
<b>AB</b>		<b>20MG</b>	<b>A079234 003</b>	Jan 27, 2009
<b>AB</b>		<b>100MG</b>	<b>A079234 004</b>	Jan 27, 2009
<b>AB</b>	PAR PHARM	<b>5MG</b>	<b>A076226 001</b>	May 27, 2003
<b>AB</b>		<b>10MG</b>	<b>A076226 002</b>	May 27, 2003
<b>AB</b>		<b>20MG</b>	<b>A076226 003</b>	May 27, 2003
<b>AB</b>		<b>100MG</b>	<b>A076226 004</b>	May 27, 2003
<b>AB</b>	PLIVA PHARM IND	<b>5MG</b>	<b>A076346 001</b>	May 30, 2003
<b>AB</b>		<b>10MG</b>	<b>A076346 002</b>	May 30, 2003
<b>AB</b>		<b>20MG</b>	<b>A076346 003</b>	May 30, 2003
<b>AB</b>		<b>100MG</b>	<b>A076346 004</b>	Oct 19, 2004
<b>AB</b>	TEVA	<b>5MG</b>	<b>A076110 001</b>	May 14, 2002
<b>AB</b>		<b>10MG</b>	<b>A076110 002</b>	May 14, 2002
<b>AB</b>		<b>20MG</b>	<b>A076110 003</b>	May 14, 2002
<b>AB</b>		<b>100MG</b>	<b>A076110 004</b>	May 14, 2002
<b>AB</b>	VINTAGE PHARMS	<b>5MG</b>	<b>A090613 001</b>	Mar 22, 2011
<b>AB</b>		<b>10MG</b>	<b>A090613 002</b>	Mar 22, 2011
<b>AB</b>		<b>20MG</b>	<b>A090613 003</b>	Mar 22, 2011
<b>AB</b>		<b>100MG</b>	<b>A090613 004</b>	Mar 22, 2011
<b>AB</b>	WEST-WARD PHARMS INT	<b>5MG</b>	<b>A076943 001</b>	Mar 01, 2005
<b>AB</b>		<b>10MG</b>	<b>A076943 002</b>	Mar 01, 2005
<b>AB</b>		<b>20MG</b>	<b>A076943 003</b>	Mar 01, 2005

**TRABECTEDIN**

POWDER;IV (INFUSION)

YONDELIS

+! JANSSEN PRODS 1MG/VIAL

N207953 001 Oct 23, 2015

**TRAMADOL HYDROCHLORIDE**

CAPSULE, EXTENDED RELEASE;ORAL

CONZIP

+!	CIPHER PHARMS INC	100MG
+		150MG
+		200MG
+		300MG

N022370 001	May 07, 2010
N022370 004	Aug 01, 2011
N022370 002	May 07, 2010
N022370 003	May 07, 2010

TABLET;ORAL

**TRAMADOL HYDROCHLORIDE**

<b>AB</b>	ACI HEALTHCARE LTD	<b>50MG</b>	<b>A202075 001</b>	Nov 28, 2011
<b>AB</b>	AMNEAL PHARMS	<b>50MG</b>	<b>A076003 001</b>	Jun 20, 2002
<b>AB</b>	APOTEX	<b>50MG</b>	<b>A075981 001</b>	Jul 10, 2002
<b>AB</b>	AUROBINDO PHARMA LTD	<b>50MG</b>	<b>A203494 001</b>	Mar 31, 2014
<b>AB</b>	CSPC OUYI PHARM CO	<b>50MG</b>	<b>A091498 001</b>	Mar 29, 2013
<b>AB</b>	IPCA LABS LTD	<b>50MG</b>	<b>A201973 001</b>	Nov 16, 2012
<b>AB</b>	MACLEODS PHARMS LTD	<b>50MG</b>	<b>A205702 001</b>	Sep 25, 2015
<b>AB</b>	MYLAN	<b>50MG</b>	<b>A075986 001</b>	Jun 21, 2002
<b>AB</b>	PLIVA	<b>50MG</b>	<b>A075982 001</b>	Jul 01, 2002
<b>AB</b>	SPECGX LLC	<b>50MG</b>	<b>A075983 001</b>	Jun 25, 2002
<b>AB</b>	SUN PHARM INDS INC	<b>50MG</b>	<b>A075964 001</b>	Jun 19, 2002
<b>AB</b>	SUN PHARM INDUSTRIES	<b>50MG</b>	<b>A076100 001</b>	Jun 20, 2002
<b>AB</b>	TEVA	<b>50MG</b>	<b>A075977 001</b>	Jun 19, 2002
<b>AB</b>	ZYDUS PHARMS USA INC	<b>50MG</b>	<b>A090404 001</b>	Jan 31, 2011

**ULTRAM**

<b>AB</b>	+! JANSSEN PHARMS	<b>50MG</b>	<b>N020281 002</b>	Mar 03, 1995
	TABLET, EXTENDED RELEASE;ORAL			

**TRAMADOL HYDROCHLORIDE**

<b>AB1</b>	AUROBINDO PHARMA LTD	<b>100MG</b>	<b>A204421 001</b>	Oct 20, 2015
<b>AB1</b>		<b>200MG</b>	<b>A204421 002</b>	Oct 20, 2015
<b>AB1</b>		<b>300MG</b>	<b>A204421 003</b>	Oct 20, 2015
<b>AB1</b>	! LUPIN LTD	<b>100MG</b>	<b>A200503 001</b>	Aug 29, 2011
<b>AB1</b>		<b>200MG</b>	<b>A200503 002</b>	Aug 29, 2011
<b>AB1</b>		<b>300MG</b>	<b>A200503 003</b>	Aug 29, 2011
<b>AB1</b>	MYLAN PHARMS INC	<b>100MG</b>	<b>A205257 001</b>	Dec 22, 2015
<b>AB1</b>		<b>200MG</b>	<b>A205257 002</b>	Dec 22, 2015
<b>AB1</b>		<b>300MG</b>	<b>A205257 003</b>	Dec 22, 2015
<b>AB1</b>	PAR PHARM INC	<b>100MG</b>	<b>A078783 001</b>	Nov 13, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-415 (of 436)

TRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB1</u>		<u>200MG</u>	<u>A078783</u> <u>002</u>	Nov 13, 2009
<u>AB1</u>		<u>300MG</u>	<u>A078783</u> <u>003</u>	Sep 20, 2011
<u>AB1</u>	SUN PHARMA GLOBAL	<u>100MG</u>	<u>A201384</u> <u>001</u>	Dec 07, 2011
<u>AB1</u>		<u>200MG</u>	<u>A201384</u> <u>002</u>	Dec 07, 2011
<u>AB1</u>		<u>300MG</u>	<u>A201384</u> <u>003</u>	Dec 07, 2011
<u>AB2</u>	ACTAVIS ELIZABETH	<u>100MG</u>	<u>A091609</u> <u>001</u>	Jun 27, 2012
<u>AB2</u>		<u>200MG</u>	<u>A091609</u> <u>002</u>	Jun 27, 2012
<u>AB2</u>		<u>300MG</u>	<u>A091609</u> <u>003</u>	Jun 27, 2012
<u>AB2</u>	ANCHEN PHARMS	<u>100MG</u>	<u>A200491</u> <u>001</u>	Jun 27, 2012
<u>AB2</u>		<u>200MG</u>	<u>A200491</u> <u>002</u>	Jun 27, 2012
<u>AB2</u>		<u>300MG</u>	<u>A200491</u> <u>003</u>	Jun 27, 2012
<u>AB2</u> !	SUN PHARMA GLOBAL	<u>100MG</u>	<u>A091607</u> <u>001</u>	Dec 30, 2011
<u>AB2</u>		<u>200MG</u>	<u>A091607</u> <u>002</u>	Dec 30, 2011
<u>AB2</u>		<u>300MG</u>	<u>A091607</u> <u>003</u>	Dec 30, 2011

TRAMETINIB DIMETHYL SULFOXIDE

TABLET; ORAL

MEKINIST

+ NOVARTIS PHARMS CORP	EQ 0.5MG	N204114 001	May 29, 2013
+!	EQ 2MG	N204114 003	May 29, 2013

TRANDOLAPRIL

TABLET; ORAL

MAVIK

<u>AB</u> + ABBVIE	<u>1MG</u>	<u>N020528</u> <u>001</u>	Apr 26, 1996
<u>AB</u> +	<u>2MG</u>	<u>N020528</u> <u>002</u>	Apr 26, 1996
<u>AB</u> +!	<u>4MG</u>	<u>N020528</u> <u>003</u>	Apr 26, 1996

TRANDOLAPRIL

<u>AB</u> AUROBINDO PHARMA	<u>1MG</u>	<u>A078438</u> <u>001</u>	Jun 12, 2007
<u>AB</u>	<u>2MG</u>	<u>A078438</u> <u>002</u>	Jun 12, 2007
<u>AB</u>	<u>4MG</u>	<u>A078438</u> <u>003</u>	Jun 12, 2007
<u>AB</u> EPIC PHARMA	<u>1MG</u>	<u>A078508</u> <u>003</u>	Jun 18, 2008
<u>AB</u>	<u>2MG</u>	<u>A078508</u> <u>001</u>	Jun 18, 2008
<u>AB</u>	<u>4MG</u>	<u>A078508</u> <u>002</u>	Jun 18, 2008
<u>AB</u> LUPIN	<u>1MG</u>	<u>A077522</u> <u>001</u>	Jun 12, 2007
<u>AB</u>	<u>2MG</u>	<u>A077522</u> <u>002</u>	Jun 12, 2007
<u>AB</u>	<u>4MG</u>	<u>A077522</u> <u>003</u>	Jun 12, 2007
<u>AB</u> TEVA PHARMS	<u>1MG</u>	<u>A077489</u> <u>001</u>	Dec 12, 2006
<u>AB</u>	<u>2MG</u>	<u>A077489</u> <u>002</u>	Dec 12, 2006
<u>AB</u>	<u>4MG</u>	<u>A077489</u> <u>003</u>	Dec 12, 2006
<u>AB</u> WATSON LABS	<u>1MG</u>	<u>A077805</u> <u>001</u>	Jun 12, 2007
<u>AB</u>	<u>2MG</u>	<u>A077805</u> <u>002</u>	Jun 12, 2007
<u>AB</u>	<u>4MG</u>	<u>A077805</u> <u>003</u>	Jun 12, 2007

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARKA

<u>AB</u> + ABBVIE	<u>1MG;240MG</u>	<u>N020591</u> <u>003</u>	Oct 22, 1996
<u>AB</u> +	<u>2MG;180MG</u>	<u>N020591</u> <u>001</u>	Oct 22, 1996
<u>AB</u> +	<u>2MG;240MG</u>	<u>N020591</u> <u>004</u>	Oct 22, 1996
<u>AB</u> +!	<u>4MG;240MG</u>	<u>N020591</u> <u>002</u>	Oct 22, 1996

TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE

<u>AB</u> GLENMARK GENERICS	<u>1MG;240MG</u>	<u>A079135</u> <u>004</u>	Aug 30, 2010
<u>AB</u>	<u>2MG;180MG</u>	<u>A079135</u> <u>001</u>	May 26, 2010
<u>AB</u>	<u>2MG;240MG</u>	<u>A079135</u> <u>002</u>	May 26, 2010
<u>AB</u>	<u>4MG;240MG</u>	<u>A079135</u> <u>003</u>	May 05, 2010

TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON

<u>AP</u> +! PHARMACIA AND UPJOHN	<u>100MG/ML</u>	<u>N019281</u> <u>001</u>	Dec 30, 1986
-----------------------------------	-----------------	---------------------------	--------------

TRANEXAMIC ACID

<u>AP</u> ACIC FINE CHEMS	<u>100MG/ML</u>	<u>A202436</u> <u>001</u>	Feb 11, 2014
<u>AP</u> AKORN	<u>100MG/ML</u>	<u>A202373</u> <u>001</u>	Nov 17, 2011
<u>AP</u>	<u>100MG/ML</u>	<u>A206594</u> <u>001</u>	Sep 28, 2017
<u>AP</u>	<u>100MG/ML</u>	<u>A206634</u> <u>001</u>	Jun 09, 2016
<u>AP</u> AMNEAL PHARMS CO	<u>100MG/ML</u>	<u>A208840</u> <u>001</u>	Feb 28, 2017
<u>AP</u> AUROBINDO PHARMA LTD	<u>100MG/ML</u>	<u>A205035</u> <u>001</u>	Jan 14, 2016
<u>AP</u> EMCURE PHARMS LTD	<u>100MG/ML</u>	<u>A203521</u> <u>001</u>	Aug 12, 2014

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-416 (of 436)

TRANEXAMIC ACID

INJECTABLE; INJECTION

**TRANEXAMIC ACID**

<b>AP</b>	FRESENIUS KABI USA	<u>100MG/ML</u>	<b>A091596 001</b>	Mar 02, 2012
<b>AP</b>	GLAND PHARMA LTD	<u>100MG/ML</u>	<b>A207239 001</b>	Feb 13, 2017
<b>AP</b>	LUITPOLD	<u>100MG/ML</u>	<b>A201885 001</b>	Aug 10, 2011
<b>AP</b>	MYLAN INSTITUTIONAL	<u>100MG/ML</u>	<b>A091657 001</b>	Nov 03, 2011
<b>AP</b>	VIRTUS PHARMS	<u>100MG/ML</u>	<b>A202755 001</b>	Feb 25, 2016
<b>AP</b>	VIVA HLTHCARE	<u>100MG/ML</u>	<b>A206713 001</b>	Jun 27, 2017
<b>AP</b>	X-GEN PHARMS INC	<u>100MG/ML</u>	<b>A201580 001</b>	Jun 14, 2013
<b>AP</b>	ZYDUS PHARMS USA INC	<u>100MG/ML</u>	<b>A205228 001</b>	Jul 17, 2017

TABLET; ORAL

**LYSTEDA**

<b>AB</b>	+!	FERRING PHARMS INC	<u>650MG</u>	<b>N022430 001</b>	Nov 13, 2009

**TRANEXAMIC ACID**

<b>AB</b>	ACTAVIS LABS FL INC	<u>650MG</u>	<b>A202093 001</b>	Dec 27, 2012
<b>AB</b>	APOTEX INC	<u>650MG</u>	<b>A202286 001</b>	Jan 27, 2014
<b>AB</b>	MYLAN	<u>650MG</u>	<b>A205133 001</b>	Sep 21, 2015

TRANYLCYPROMINE SULFATE

TABLET; ORAL

**PARNATE**

<b>AB</b>	+!	CONCORDIA PHARMS INC	<u>EQ 10MG BASE</u>	<b>N012342 003</b>	Aug 16, 1985

**TRANYLCYPROMINE SULFATE**

<b>AB</b>	PAR PHARM	<u>EQ 10MG BASE</u>	<b>A040640 001</b>	Jun 29, 2006

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

**TRAVATAN Z**

<b>AT</b>	+!	NOVARTIS PHARMS CORP	<u>0.004%</u>	<b>N021994 001</b>	Sep 21, 2006

**TRAVOPROST**

<b>AT</b>	APOTEX INC	<u>0.004%</u>	<b>A203431 001</b>	Jul 10, 2015
<b>AT</b>	MYLAN PHARMS INC	<u>0.004%</u>	<b>A205050 001</b>	Jul 07, 2017
!	PAR PHARM	0.004%	A091340 001	Mar 01, 2013

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

**TRAZODONE HYDROCHLORIDE**

<b>AB</b>	ACCORD HLTHCARE	<u>50MG</u>	<b>A206923 001</b>	Sep 08, 2017
<b>AB</b>		<u>100MG</u>	<b>A206923 002</b>	Sep 08, 2017
<b>AB</b>		<u>150MG</u>	<b>A206923 003</b>	Sep 08, 2017
<b>AB</b>		<u>300MG</u>	<b>A206923 004</b>	Sep 08, 2017
<b>AB</b>	ALVOGEN	<u>50MG</u>	<b>A071636 001</b>	Apr 18, 1988
<b>AB</b>		<u>100MG</u>	<b>A071514 001</b>	Apr 18, 1988
<b>AB</b>	APOTEX	<u>50MG</u>	<b>A071258 001</b>	Mar 25, 1987
<b>AB</b>	! APOTEX INC	<u>100MG</u>	<b>A071196 001</b>	Mar 25, 1987
<b>AB</b>		<u>150MG</u>	<b>A071196 002</b>	Apr 26, 1999
<b>AB</b>		<u>300MG</u>	<b>A071196 003</b>	Apr 26, 1999
<b>AB</b>	PLIVA	<u>150MG</u>	<b>A071525 001</b>	Mar 09, 1988
<b>AB</b>	SUN PHARM INDUSTRIES	<u>50MG</u>	<b>A073137 002</b>	Mar 24, 1993
<b>AB</b>		<u>100MG</u>	<b>A073137 001</b>	Mar 24, 1993
<b>AB</b>		<u>150MG</u>	<b>A073137 003</b>	Dec 22, 1995
<b>AB</b>	TEVA PHARMS USA	<u>50MG</u>	<b>A071523 001</b>	Dec 11, 1987
<b>AB</b>		<u>100MG</u>	<b>A071524 001</b>	Dec 11, 1987
<b>AB</b>	TORRENT PHARMS LTD	<u>50MG</u>	<b>A202180 001</b>	Nov 27, 2013
<b>AB</b>		<u>100MG</u>	<b>A202180 002</b>	Nov 27, 2013
<b>AB</b>		<u>150MG</u>	<b>A202180 003</b>	Nov 27, 2013
<b>AB</b>		<u>300MG</u>	<b>A202180 004</b>	Nov 27, 2013
<b>AB</b>	VINTAGE	<u>50MG</u>	<b>A072192 001</b>	Feb 02, 1989
<b>AB</b>		<u>100MG</u>	<b>A072193 001</b>	Feb 02, 1989
<b>AB</b>	ZYDUS PHARMS USA INC	<u>50MG</u>	<b>A205253 001</b>	Oct 10, 2017
<b>AB</b>		<u>100MG</u>	<b>A205253 002</b>	Oct 10, 2017
<b>AB</b>		<u>150MG</u>	<b>A205253 003</b>	Oct 10, 2017
<b>AB</b>		<u>300MG</u>	<b>A205253 004</b>	Oct 10, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-417 (of 436)

TREPROSTINIL

INJECTABLE; IV (INFUSION), SUBCUTANEOUS

REMODULIN

<u>AP</u>	+!	UNITED THERAP	<u>1MG/ML</u>
<u>AP</u>	+!		<u>2 . 5MG/ML</u>
<u>AP</u>	+!		<u>5MG/ML</u>
<u>AP</u>	+!		<u>10MG/ML</u>

<u>N021272</u>	<u>001</u>	May 21, 2002
<u>N021272</u>	<u>002</u>	May 21, 2002
<u>N021272</u>	<u>003</u>	May 21, 2002
<u>N021272</u>	<u>004</u>	May 21, 2002

TREPROSTINIL

<u>AP</u>	SANDOZ INC	<u>1MG/ML</u>
<u>AP</u>		<u>2 . 5MG/ML</u>
<u>AP</u>		<u>5MG/ML</u>
<u>AP</u>		<u>10MG/ML</u>

<u>A203649</u>	<u>001</u>	Nov 30, 2017
<u>A203649</u>	<u>002</u>	Nov 30, 2017
<u>A203649</u>	<u>003</u>	Nov 30, 2017
<u>A203649</u>	<u>004</u>	Nov 30, 2017

SOLUTION; INHALATION

TYVASO

+! UNITED THERAP

0 . 6MG/ML

N022387 001 Jul 30, 2009

TREPROSTINIL DIOLAMINE

TABLET, EXTENDED RELEASE; ORAL

ORENITRAM

+ UNITED THERAP	EQ 0.125MG BASE
+	EQ 0.25MG BASE
+	EQ 1MG BASE
+!	EQ 2 . 5MG BASE
+	EQ 5MG BASE

N203496 001	Dec 20, 2013
N203496 002	Dec 20, 2013
N203496 003	Dec 20, 2013
N203496 004	Dec 20, 2013
N203496 005	Oct 07, 2016

TRETINOIN

CAPSULE; ORAL

TRETINOIN

<u>AB</u>	ANCHEN PHARMS	<u>10MG</u>
<u>AB</u>	! BARR LABS INC	<u>10MG</u>
<u>AB</u>	GLENMARK PHARMS LTD	<u>10MG</u>

<u>A201687</u>	<u>001</u>	Oct 24, 2012
<u>A077684</u>	<u>001</u>	Jun 22, 2007
<u>A208279</u>	<u>001</u>	Dec 23, 2016

CREAM; TOPICAL

AVITA

<u>AB</u>	MYLAN PHARMS INC	<u>0 . 025%</u>
-----------	------------------	-----------------

N020404 003 Jan 14, 1997

RETIN-A

<u>AB</u>	+! VALEANT BERMUDA	<u>0 . 025%</u>
<u>AB</u>	VALEANT PHARMS NORTH	<u>0 . 1%</u>

<u>N019049</u>	<u>001</u>	Sep 16, 1988
<u>N017340</u>	<u>001</u>	

TRETINOIN

<u>AB</u>	PERRIGO PHARMA INTL	<u>0 . 025%</u>
<u>AB</u>		<u>0 . 1%</u>

<u>A075264</u>	<u>001</u>	Dec 24, 1998
<u>A075213</u>	<u>001</u>	Dec 24, 1998

RETIN-A

<u>AB1</u>	+! VALEANT BERMUDA	<u>0 . 05%</u>
------------	--------------------	----------------

N017522 001

TRETINOIN

<u>AB1</u>	PERRIGO PHARMA INTL	<u>0 . 05%</u>
------------	---------------------	----------------

<u>A075265</u>	<u>001</u>	Dec 24, 1998
----------------	------------	--------------

RENOVA

<u>AB2</u>	+! VALEANT PHARMS NORTH	<u>0 . 05%</u>
------------	-------------------------	----------------

<u>N019963</u>	<u>001</u>	Dec 29, 1995
----------------	------------	--------------

TRETINOIN

<u>AB2</u>	ZO SKIN HEALTH RENOVA	<u>0 . 05%</u>
	+! VALEANT PHARMS NORTH	0 . 02%

<u>A076498</u>	<u>001</u>	Sep 15, 2005
N021108	001	Aug 31, 2000

GEL; TOPICAL

ATRALIN

<u>AB</u>	+! DOW PHARM	<u>0 . 05%</u>
-----------	--------------	----------------

<u>N022070</u>	<u>001</u>	Jul 26, 2007
<u>N017955</u>	<u>001</u>	

RETIN-A

<u>AB</u>	+! VALEANT INTL	<u>0 . 01%</u>
<u>AB</u>	+	<u>0 . 025%</u>

<u>N017579</u>	<u>002</u>	
----------------	------------	--

RETIN-A MICRO

<u>AB</u>	+! VALEANT INTL	<u>0 . 04%</u>
<u>AB</u>	+	<u>0 . 1%</u>

<u>N020475</u>	<u>002</u>	May 10, 2002
<u>N020475</u>	<u>001</u>	Feb 07, 1997

TRETINOIN

<u>AB</u>	MYLAN PHARMS INC	<u>0 . 04%</u>
<u>AB</u>		<u>0 . 05%</u>
<u>AB</u>		<u>0 . 1%</u>

<u>A202567</u>	<u>001</u>	Jul 17, 2013
<u>A207955</u>	<u>001</u>	Aug 13, 2015
<u>A202026</u>	<u>001</u>	Jul 17, 2013

<u>AB</u>	PERRIGO PHARMA INTL	<u>0 . 01%</u>
<u>AB</u>		<u>0 . 025%</u>

<u>A075589</u>	<u>001</u>	Jun 11, 2002
<u>A075529</u>	<u>001</u>	Feb 22, 2000

AVITA

BT	MYLAN	0 . 025%
----	-------	----------

N020400	001	Jan 29, 1998
---------	-----	--------------

RETIN-A-MICRO

+!	VALEANT INTL	0 . 06%
----	--------------	---------

N020475	004	Oct 23, 2017
---------	-----	--------------

+!		0 . 08%
----	--	---------

N020475	003	Jan 28, 2014
---------	-----	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-418 (of 436)

TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	ALKEM LABS LTD	<u>0.025%</u>	<u>A207651</u> <u>001</u>	Dec 26, 2017
<u>AT</u>		<u>0.1%</u>	<u>A207651</u> <u>002</u>	Dec 26, 2017
<u>AT</u>		<u>0.5%</u>	<u>A207651</u> <u>003</u>	Dec 26, 2017
<u>AT</u> !	FOUGERA PHARMS	<u>0.025%</u>	<u>A085692</u> <u>001</u>	
<u>AT</u> !		<u>0.1%</u>	<u>A085692</u> <u>003</u>	
<u>AT</u> !		<u>0.5%</u>	<u>A085692</u> <u>002</u>	
<u>AT</u>	G AND W LABS	<u>0.025%</u>	<u>A089797</u> <u>001</u>	May 31, 1991
<u>AT</u>		<u>0.1%</u>	<u>A089798</u> <u>001</u>	May 31, 1991
<u>AT</u>	GLENMARK PHARMS LTD	<u>0.1%</u>	<u>A207117</u> <u>001</u>	Aug 05, 2016
<u>AT</u>	LUPIN ATLANTIS	<u>0.025%</u>	<u>A208763</u> <u>001</u>	Feb 01, 2017
<u>AT</u>		<u>0.1%</u>	<u>A208763</u> <u>002</u>	Feb 01, 2017
<u>AT</u>		<u>0.5%</u>	<u>A208763</u> <u>003</u>	Feb 01, 2017
<u>AT</u> +	MYLAN PHARMS INC	<u>0.025%</u>	<u>N011601</u> <u>003</u>	
<u>AT</u> +		<u>0.1%</u>	<u>N011601</u> <u>006</u>	
<u>AT</u>	PERRIGO NEW YORK	<u>0.025%</u>	<u>A086415</u> <u>001</u>	
<u>AT</u>		<u>0.1%</u>	<u>A086414</u> <u>001</u>	
<u>AT</u>		<u>0.5%</u>	<u>A086413</u> <u>001</u>	
<u>AT</u>	TARO	<u>0.1%</u>	<u>A040039</u> <u>001</u>	Nov 26, 1997
<u>AT</u>	TELIGENT PHARMA INC	<u>0.1%</u>	<u>A208848</u> <u>001</u>	Sep 18, 2017
<u>AT</u>	VINTAGE	<u>0.025%</u>	<u>A040671</u> <u>001</u>	Jun 09, 2006
<u>AT</u>		<u>0.1%</u>	<u>A040671</u> <u>002</u>	Jun 09, 2006
<u>TRIDERM</u>				
<u>AT</u>	CROWN LABS	<u>0.025%</u>	<u>A088042</u> <u>002</u>	Mar 25, 2015
<u>AT</u>		<u>0.1%</u>	<u>A088042</u> <u>001</u>	Mar 19, 1984
<u>AT</u>		<u>0.5%</u>	<u>A088042</u> <u>003</u>	Mar 25, 2015

FOR SUSPENSION, EXTENDED RELEASE; INTRA-ARTICULAR

ZILRETTA

+! FLEXION THERAPS INC 32MG/VIAL

N208845 001 Oct 06, 2017

INJECTABLE; INJECTION

KENALOG-40

<u>AP</u> +!	APOTHECON	<u>40MG/ML</u>	<u>N014901</u> <u>001</u>
<u>TRIAMCINOLONE ACETONIDE</u>			
<u>AP</u>	AMNEAL PHARMS CO	<u>40MG/ML</u>	<u>A207550</u> <u>001</u>
	KENALOG-10		Dec 11, 2017
+	APOTHECON	10MG/ML	N012041 001
INJECTABLE; INTRAVITREAL			
TRIESENCE			
+!	NOVARTIS PHARMS CORP	40MG/ML (40MG/ML)	N022048 001 Nov 29, 2007

LOTION;TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	AKORN	<u>0.025%</u>	<u>A202374</u> <u>001</u>	May 08, 2013
<u>AT</u>		<u>0.1%</u>	<u>A202374</u> <u>002</u>	May 08, 2013
<u>AT</u>	FOUGERA PHARMS	<u>0.025%</u>	<u>A040467</u> <u>001</u>	Apr 21, 2003
<u>AT</u>		<u>0.1%</u>	<u>A040467</u> <u>002</u>	Apr 21, 2003
<u>AT</u>	G AND W LABS INC	<u>0.1%</u>	<u>A089129</u> <u>001</u>	Aug 14, 1986
<u>AT</u>	TELIGENT PHARMA INC	<u>0.025%</u>	<u>A204608</u> <u>001</u>	Jul 07, 2016
<u>AT</u>		<u>0.1%</u>	<u>A204606</u> <u>001</u>	Jul 07, 2016
<u>AT</u>	VINTAGE	<u>0.1%</u>	<u>A040672</u> <u>002</u>	Dec 13, 2006
<u>AT</u> !	WOCKHARDT BIO AG	<u>0.025%</u>	<u>A088450</u> <u>001</u>	Apr 01, 1985
<u>AT</u> !		<u>0.1%</u>	<u>A088451</u> <u>001</u>	Apr 03, 1985

OINTMENT;TOPICAL

TRIAMCINOLONE ACETONIDE

<u>AT</u>	FOUGERA PHARMS	<u>0.025%</u>	<u>A085691</u> <u>001</u>
<u>AT</u>		<u>0.1%</u>	<u>A085691</u> <u>003</u>
<u>AT</u>		<u>0.5%</u>	<u>A085691</u> <u>002</u>
<u>AT</u>	G AND W LABS	<u>0.025%</u>	<u>A089795</u> <u>001</u>
<u>AT</u>		<u>0.1%</u>	<u>A089796</u> <u>001</u>
<u>AT</u>	G AND W LABS INC	<u>0.5%</u>	<u>A208925</u> <u>001</u>
<u>AT</u>	GLENMARK PHARMS	<u>0.1%</u>	<u>A208320</u> <u>001</u>
<u>AT</u>	GLENMARK PHARMS LTD	<u>0.5%</u>	<u>A206379</u> <u>001</u>
<u>AT</u> !	PERRIGO NEW YORK	<u>0.025%</u>	<u>A087356</u> <u>001</u>
<u>AT</u> !		<u>0.1%</u>	<u>A087357</u> <u>001</u>
<u>AT</u> !		<u>0.5%</u>	<u>A087385</u> <u>001</u>
<u>AT</u>	TARO	<u>0.1%</u>	<u>A040037</u> <u>001</u>
<u>AT</u>	TELIGENT PHARMA INC	<u>0.1%</u>	<u>A205373</u> <u>001</u>
<u>AT</u>		<u>0.5%</u>	<u>A208590</u> <u>001</u>

TRIAMCINOLONE ACETONIDE IN ABSORB BASE

! CMP PHARMA INC 0.05%

A089595 001 Mar 23, 1995



38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-420 (of 436)

TRIHEXYPHENIDYL HYDROCHLORIDE

TABLET;ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

<u>AA</u>		<u>5MG</u>	<u>A040254 002</u> Dec 24, 1998
<u>AA</u>	!	WATSON LABS	<u>A084363 001</u>
<u>AA</u>	!	<u>5MG</u>	<u>A084364 001</u>

TRIMETHADIONE

TABLET;ORAL

TRIDIONE

+! ABBVIE 150MG

N005856 009

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE;ORAL

TIGAN

<u>AB</u>	+	KING PHARMS LLC	<u>300MG</u>	<u>N017531 006</u> Dec 13, 2001
-----------	---	-----------------	--------------	---------------------------------

TRIMETHOBENZAMIDE HYDROCHLORIDE

<u>AB</u>		GAVIS PHARMS	<u>300MG</u>	<u>A076546 001</u> Aug 20, 2003
-----------	--	--------------	--------------	---------------------------------

<u>AB</u>		SUN PHARM INDUSTRIES	<u>300MG</u>	<u>A076570 001</u> Aug 28, 2003
-----------	--	----------------------	--------------	---------------------------------

INJECTABLE;INJECTION

TIGAN

<u>AP</u>	+	PAR STERILE PRODUCTS	<u>100MG/ML</u>	<u>N017530 001</u>
-----------	---	----------------------	-----------------	--------------------

TRIMETHOBENZAMIDE HYDROCHLORIDE

<u>AP</u>		LUITPOLD	<u>100MG/ML</u>	<u>A091330 001</u> Mar 08, 2011
-----------	--	----------	-----------------	---------------------------------

TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>		LUITPOLD	<u>100MG/ML</u>	<u>A091329 001</u> Mar 08, 2011
-----------	--	----------	-----------------	---------------------------------

TRIMETHOPRIM

TABLET;ORAL

TRIMETHOPRIM

<u>AB</u>	+	MAYNE PHARMA	<u>100MG</u>	<u>N018679 001</u> Jul 30, 1982
<u>AB</u>		NOVEL LABS INC	<u>100MG</u>	<u>A091437 001</u> Jun 15, 2011
<u>AB</u>		WATSON LABS	<u>100MG</u>	<u>A070049 001</u> Jun 06, 1985

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION;ORAL

PRIMSOL

+! ALLEGIS EQ 50MG BASE/5ML

N074973 001 Jan 24, 2000

TRIMIPRAMINE MALEATE

CAPSULE;ORAL

SURMONTIL

<u>AB</u>	+	ODYSSEY PHARMS	<u>EQ 25MG BASE</u>	<u>N016792 001</u>
<u>AB</u>	+		<u>EQ 50MG BASE</u>	<u>N016792 002</u>
<u>AB</u>	+		<u>EQ 100MG BASE</u>	<u>N016792 003</u> Sep 15, 1982

TRIMIPRAMINE MALEATE

<u>AB</u>		CROSSMEDIKA SA	<u>EQ 25MG BASE</u>	<u>A208127 001</u> Apr 15, 2016
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A208127 002</u> Apr 15, 2016
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A208127 003</u> Apr 15, 2016

<u>AB</u>		ELITE LABS INC	<u>EQ 25MG BASE</u>	<u>A077361 001</u> Aug 02, 2006
<u>AB</u>			<u>EQ 50MG BASE</u>	<u>A077361 002</u> Aug 02, 2006
<u>AB</u>			<u>EQ 100MG BASE</u>	<u>A077361 003</u> Aug 02, 2006

TRIPTORELIN PAMOATE

FOR SUSPENSION, EXTENDED RELEASE;INTRAMUSCULAR

TRIPTODUR KIT

+! ARBOR PHARMS LLC EQ 22.5MG BASE/VIAL

N208956 001 Jun 29, 2017

INJECTABLE;INTRAMUSCULAR

TRELSTAR

+! ALLERGAN SALES LLC EQ 3.75MG BASE/VIAL

N020715 001 Jun 15, 2000

+! EQ 11.25MG BASE/VIAL

N021288 001 Jun 29, 2001

+! EQ 22.5MG BASE/VIAL

N022437 001 Mar 10, 2010

TROMETHAMINE

INJECTABLE;INJECTION

THAM

+! HOSPIRA 3.6GM/100ML

N013025 002

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

MYDRIACYL

<u>AT</u>	!	NOVARTIS PHARMS CORP	<u>1%</u>	<u>A084306 001</u>
<u>AT</u>	!	SANDOZ INC	<u>0.5%</u>	<u>A084305 001</u>

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-421 (of 436)

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC

TROPICACYL

<u>AT</u>	AKORN	<u>0.5%</u>	<u>A040314 001</u>	Sep 29, 2000
<u>AT</u>		<u>1%</u>	<u>A040315 001</u>	Sep 29, 2000
<u>TROPICAMIDE</u>				
<u>AT</u>	BAUSCH AND LOMB	<u>0.5%</u>	<u>A040067 001</u>	Jul 27, 1994
<u>AT</u>		<u>1%</u>	<u>A040064 001</u>	Jul 27, 1994

TROSPiUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

TROSPiUM CHLORIDE

<u>AB</u>	! ACTAVIS LABS FL INC	<u>60MG</u>	<u>A091289 001</u>	Oct 12, 2012
<u>AB</u>	PADDICK LLC	<u>60MG</u>	<u>A201291 001</u>	May 24, 2013

TABLET;ORAL

TROSPiUM CHLORIDE

<u>AB</u>	APOTEX	<u>20MG</u>	<u>A091513 001</u>	Dec 06, 2011
<u>AB</u>	! GLENMARK GENERICS	<u>20MG</u>	<u>A091575 001</u>	Aug 13, 2010
<u>AB</u>	HERITAGE PHARMS INC	<u>20MG</u>	<u>A204945 001</u>	Aug 30, 2016
<u>AB</u>	INVAGEN PHARMS	<u>20MG</u>	<u>A091688 001</u>	Aug 23, 2016
<u>AB</u>	PADDICK LLC	<u>20MG</u>	<u>A091573 001</u>	Nov 17, 2010

TRYPAN BLUE

SOLUTION;OPHTHALMIC

MEMBRANEBLUE

+!	DORC	0.15%	N022278 001	Feb 20, 2009
+!	DORC	0.06%	N021670 001	Dec 16, 2004

ULIPRISTAL ACETATE

TABLET;ORAL

ELLA

<u>AB</u>	+! LAB HRA PHARMA	<u>30MG</u>	<u>N022474 001</u>	Aug 13, 2010
<u>AB</u>	TEVA PHARMS USA	<u>30MG</u>	<u>A207952 001</u>	Feb 13, 2017

UMECLIDINIUM BROMIDE

POWDER;INHALATION

INCRUSE ELLIPTA

+!	GLAXO GRP ENGLAND	EQ 62.5MCG BASE/INH	N205382 001	Apr 30, 2014
----	-------------------	---------------------	-------------	--------------

UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE

POWDER;INHALATION

ANORO ELLIPTA

+!	GLAXOSMITHKLINE	EQ 0.0625MG BASE/INH;EQ 0.025MG BASE/INH	N203975 001	Dec 18, 2013
----	-----------------	--	-------------	--------------

UREA, C-14

CAPSULE;ORAL

PYTEST

+!	AVENT	1uCi	N020617 001	May 09, 1997
Pytest Kit				

+! AVENT

URIDINE TRIACETATE

GRANULE;ORAL

VISTOGARD

+!	WELLSTAT THERAP	10GM/PACKET	N208159 001	Dec 11, 2015
+!	WELLSTAT THERAP	2GM/PACKET	N208169 001	Sep 04, 2015

XURIDEN

+! WELLSTAT THERAP

UROFOLLITROPIN

INJECTABLE;INTRAMUSCULAR, SUBCUTANEOUS

BRAVELLE

+!	FERRING	75 IU/VIAL	N021289 001	May 06, 2002
----	---------	------------	-------------	--------------

URSODIOL

CAPSULE;ORAL

ACTIGALL

<u>AB</u>	+! ALLERGAN SALES LLC	<u>300MG</u>	<u>N019594 002</u>	Dec 31, 1987

URSODIOL

<u>AB</u>	EPIC PHARMA	<u>300MG</u>	<u>A075517 001</u>	Mar 14, 2000
<u>AB</u>	LANNETT	<u>300MG</u>	<u>A079082 001</u>	Dec 15, 2008
<u>AB</u>	MYLAN	<u>300MG</u>	<u>A090530 001</u>	Feb 17, 2010
<u>AB</u>	TEVA PHARMS	<u>300MG</u>	<u>A075592 001</u>	May 25, 2000

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
PRESCRIPTION DRUG PRODUCT LIST

3-422 (of 436)

URSODIOL

TABLET;ORAL

<u>URSO 250</u>		<u>250MG</u>	<u>N020675 001</u>	Dec 10, 1997
<u>AB</u>	+ FOREST LABS INC	<u>250MG</u>		
<u>URSO FORTE</u>			<u>N020675 002</u>	Jul 21, 2004
<u>AB</u>	+! FOREST LABS INC	<u>500MG</u>		
<u>URSODIOL</u>				
<u>AB</u>	GLENMARK GENERICS	<u>250MG</u>	<u>A090801 001</u>	Jul 12, 2011
<u>AB</u>		<u>500MG</u>	<u>A090801 002</u>	Jul 12, 2011
<u>AB</u>	IMPAX LABS INC	<u>250MG</u>	<u>A200826 001</u>	Dec 23, 2011
<u>AB</u>		<u>500MG</u>	<u>A200826 002</u>	Dec 23, 2011
<u>AB</u>	PAR PHARM	<u>250MG</u>	<u>A202540 001</u>	Feb 14, 2013
<u>AB</u>		<u>500MG</u>	<u>A202540 002</u>	Feb 14, 2013

VALACYCLOVIR HYDROCHLORIDE

TABLET;ORAL

<u>VALACYCLOVIR HYDROCHLORIDE</u>				
<u>AB</u>	APOTEX INC	<u>EQ 500MG BASE</u>	<u>A090500 001</u>	Apr 04, 2014
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090500 002</u>	Apr 04, 2014
<u>AB</u>	AUROBINDO PHARMA	<u>EQ 500MG BASE</u>	<u>A090682 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090682 002</u>	May 24, 2010
<u>AB</u>	CIPLA LTD	<u>EQ 500MG BASE</u>	<u>A077135 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A077135 002</u>	May 24, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 500MG BASE</u>	<u>A079012 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A079012 002</u>	May 24, 2010
<u>AB</u>	HETERO LABS LTD V	<u>EQ 500MG BASE</u>	<u>A203047 001</u>	Apr 08, 2015
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A203047 002</u>	Apr 08, 2015
<u>AB</u>	JUBILANT GENERICS	<u>EQ 500MG BASE</u>	<u>A201506 001</u>	Apr 03, 2012
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A201506 002</u>	Apr 03, 2012
<u>AB</u>	MYLAN PHARMS INC	<u>EQ 500MG BASE</u>	<u>A078518 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A078518 002</u>	May 24, 2010
<u>AB</u>	SANDOZ	<u>EQ 500MG BASE</u>	<u>A077478 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A077478 002</u>	May 24, 2010
<u>AB</u>	SUN PHARM INDs LTD	<u>EQ 500MG BASE</u>	<u>A076588 001</u>	Jan 31, 2007
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A076588 002</u>	Jan 31, 2007
<u>AB</u>	TEVA PHARMS	<u>EQ 500MG BASE</u>	<u>A077655 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A077655 002</u>	May 24, 2010
<u>AB</u>	WATSON LABS INC	<u>EQ 500MG BASE</u>	<u>A090370 001</u>	Mar 16, 2011
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090370 002</u>	Mar 16, 2011
<u>AB</u>	WEST-WARD PHARMS INT	<u>EQ 500MG BASE</u>	<u>A078656 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A078656 002</u>	May 24, 2010
<u>AB</u>	WOCKHARDT	<u>EQ 500MG BASE</u>	<u>A090216 001</u>	May 24, 2010
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A090216 002</u>	May 24, 2010
<u>AB</u>	ZYDUS PHARMS USA INC	<u>EQ 500MG BASE</u>	<u>A079137 001</u>	Dec 29, 2017
<u>AB</u>		<u>EQ 1GM BASE</u>	<u>A079137 002</u>	Dec 29, 2017
<u>VALTREX</u>				
<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 500MG BASE</u>	<u>N020487 001</u>	Jun 23, 1995
<u>AB</u>	+!	<u>EQ 1GM BASE</u>	<u>N020487 002</u>	Jun 23, 1995

VALBENAZINE TOSYLATE

CAPSULE;ORAL

INGREZZA

+! NEUROCRINE	40MG	<u>N209241 001</u>	Apr 11, 2017
+	80MG	<u>N209241 002</u>	Oct 04, 2017

VALGANCICLOVIR HYDROCHLORIDE

FOR SOLUTION;ORAL

<u>VALCYTE</u>				
<u>AB</u>	+! HOFFMANN LA ROCHE	<u>50MG/ML</u>	<u>N022257 001</u>	Aug 28, 2009
<u>VALGANCICLOVIR HYDROCHLORIDE</u>				
<u>AB</u>	ACTAVIS LABS FL INC	<u>50MG/ML</u>	<u>A205220 001</u>	Jul 18, 2016
TABLET;ORAL				
<u>VALCYTE</u>				
<u>AB</u>	+! HOFFMANN LA ROCHE	<u>EQ 450MG BASE</u>	<u>N021304 001</u>	Mar 29, 2001
<u>VALGANCICLOVIR HYDROCHLORIDE</u>				
<u>AB</u>	AUROBINDO PHARMA LTD	<u>EQ 450MG BASE</u>	<u>A204750 001</u>	Mar 31, 2016
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 450MG BASE</u>	<u>A203511 001</u>	Nov 04, 2014
<u>AB</u>		<u>EQ 450MG BASE</u>	<u>A206876 001</u>	Dec 12, 2017
<u>AB</u>	ENDO PHARMS INC	<u>EQ 450MG BASE</u>	<u>A200790 001</u>	Nov 04, 2014
<u>AB</u>	HETERO LABS LTD V	<u>EQ 450MG BASE</u>	<u>A205166 001</u>	Mar 18, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-423 (of 436)

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPACON

<u>AP</u>	+!	ABBVIE	<u>EQ 100MG BASE/ML</u>	<u>N020593 001</u>	Dec 30, 1996
		<u>VALPROATE SODIUM</u>			
<u>AP</u>		ATHENEX INC	<u>EQ 100MG BASE/ML</u>	<u>A076295 001</u>	Nov 14, 2002
<u>AP</u>		FRESENIUS KABI USA	<u>EQ 100MG BASE/ML</u>	<u>A076539 001</u>	Jun 26, 2003

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

<u>AB</u>	+!	ABBVIE	<u>250MG</u>	<u>N018081 001</u>	
		<u>VALPROIC ACID</u>			
<u>AB</u>		BIONPHARMA INC	<u>250MG</u>	<u>A073484 001</u>	Jun 29, 1993
<u>AB</u>		CATALENT	<u>250MG</u>	<u>A073229 001</u>	Oct 29, 1991

SYRUP; ORAL

DEPAKENE

<u>AA</u>	+!	ABBVIE	<u>250MG/5ML</u>	<u>N018082 001</u>	
		<u>VALPROIC ACID</u>			
<u>AA</u>		ANI PHARMS INC	<u>250MG/5ML</u>	<u>A073178 001</u>	Aug 25, 1992
<u>AA</u>		ECI PHARMS LLC	<u>250MG/5ML</u>	<u>A090517 001</u>	May 28, 2010
<u>AA</u>		HIGH TECH PHARMA	<u>250MG/5ML</u>	<u>A074060 001</u>	Jan 13, 1995
<u>AA</u>		PHARM ASSOC	<u>250MG/5ML</u>	<u>A075379 001</u>	Dec 15, 2000
<u>AA</u>		VINTAGE	<u>250MG/5ML</u>	<u>A077960 001</u>	Oct 13, 2006
<u>AA</u>		VISTAPHARM	<u>250MG/5ML</u>	<u>A075782 001</u>	Dec 22, 2000
<u>AA</u>		WOCKHARDT BIO AG	<u>250MG/5ML</u>	<u>A070868 001</u>	Jul 01, 1986

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

+! ENDO PHARM

40MG/ML

N020892 001 Sep 25, 1998

VALSARTAN

SOLUTION; ORAL

PREXXARTAN

+! CARMEL BIOSCIENCES

20MG/5ML

N209139 001 Dec 19, 2017

+!

80MG/20ML

N209139 002 Dec 19, 2017

TABLET; ORAL

DIOVAN

<u>AB</u>	+	NOVARTIS	<u>40MG</u>	<u>N021283 004</u>	Aug 14, 2002
<u>AB</u>	+		<u>80MG</u>	<u>N021283 001</u>	Jul 18, 2001
<u>AB</u>	+		<u>160MG</u>	<u>N021283 002</u>	Jul 18, 2001
<u>AB</u>	+		<u>320MG</u>	<u>N021283 003</u>	Jul 18, 2001

VALSARTAN

<u>AB</u>		ALEMBIC PHARMS LTD	<u>40MG</u>	<u>A091367 001</u>	Jan 05, 2015
<u>AB</u>			<u>80MG</u>	<u>A091367 002</u>	Jan 05, 2015
<u>AB</u>			<u>160MG</u>	<u>A091367 003</u>	Jan 05, 2015
<u>AB</u>			<u>320MG</u>	<u>A091367 004</u>	Jan 05, 2015
<u>AB</u>		AMNEAL PHARMS	<u>40MG</u>	<u>A204011 001</u>	Jan 11, 2016
<u>AB</u>			<u>80MG</u>	<u>A204011 002</u>	Jan 11, 2016
<u>AB</u>			<u>160MG</u>	<u>A204011 003</u>	Jan 11, 2016
<u>AB</u>			<u>320MG</u>	<u>A204011 004</u>	Jan 11, 2016
<u>AB</u>		AUROBINDO PHARMA LTD	<u>40MG</u>	<u>A202223 001</u>	Jan 05, 2015
<u>AB</u>			<u>80MG</u>	<u>A202223 002</u>	Jan 05, 2015
<u>AB</u>			<u>160MG</u>	<u>A202223 003</u>	Jan 05, 2015
<u>AB</u>			<u>320MG</u>	<u>A202223 004</u>	Jan 05, 2015
<u>AB</u>		HETERO LABS LTD V	<u>40MG</u>	<u>A203311 001</u>	Jan 05, 2015
<u>AB</u>			<u>80MG</u>	<u>A203311 002</u>	Jan 05, 2015
<u>AB</u>			<u>160MG</u>	<u>A203311 003</u>	Jan 05, 2015
<u>AB</u>			<u>320MG</u>	<u>A203311 004</u>	Jan 05, 2015
<u>AB</u>		IVAX PHARMS	<u>40MG</u>	<u>A077530 001</u>	Jan 04, 2016
<u>AB</u>			<u>80MG</u>	<u>A077530 002</u>	Jan 04, 2016
<u>AB</u>			<u>160MG</u>	<u>A077530 003</u>	Jan 04, 2016
<u>AB</u>			<u>320MG</u>	<u>A077530 004</u>	Jan 04, 2016
<u>AB</u>		JUBILANT GENERICS	<u>40MG</u>	<u>A203536 001</u>	Jan 05, 2015
<u>AB</u>			<u>80MG</u>	<u>A203536 002</u>	Jan 05, 2015
<u>AB</u>			<u>160MG</u>	<u>A203536 003</u>	Jan 05, 2015
<u>AB</u>			<u>320MG</u>	<u>A203536 004</u>	Jan 05, 2015
<u>AB</u>		LUPIN LTD	<u>40MG</u>	<u>A201677 001</u>	Jan 05, 2015
<u>AB</u>			<u>80MG</u>	<u>A201677 002</u>	Jan 05, 2015
<u>AB</u>			<u>160MG</u>	<u>A201677 003</u>	Jan 05, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-424 (of 436)

VALSARTAN

TABLET;ORAL

VALSARTAN

<u>AB</u>		<u>320MG</u>	<u>A201677 004</u>	Jan 05, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>40MG</u>	<u>A202696 001</u>	Sep 16, 2016
<u>AB</u>		<u>80MG</u>	<u>A202696 002</u>	Sep 16, 2016
<u>AB</u>		<u>160MG</u>	<u>A202696 003</u>	Sep 16, 2016
<u>AB</u>		<u>320MG</u>	<u>A202696 004</u>	Sep 16, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>40MG</u>	<u>A090866 001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A090866 002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A090866 003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A090866 004</u>	Jan 05, 2015
<u>AB</u>	OHM LABS INC	<u>40MG</u>	<u>A077492 001</u>	Jun 26, 2014
<u>AB</u>		<u>80MG</u>	<u>A077492 002</u>	Jun 26, 2014
<u>AB</u>		<u>160MG</u>	<u>A077492 003</u>	Jun 26, 2014
<u>AB</u>		<u>320MG</u>	<u>A077492 004</u>	Jun 26, 2014
<u>AB</u>	PRINSTON INC	<u>40MG</u>	<u>A204821 001</u>	Jun 09, 2015
<u>AB</u>		<u>80MG</u>	<u>A204821 002</u>	Jun 09, 2015
<u>AB</u>		<u>160MG</u>	<u>A204821 003</u>	Jun 09, 2015
<u>AB</u>		<u>320MG</u>	<u>A204821 004</u>	Jun 09, 2015
<u>AB</u>	TORRENT PHARMS LTD	<u>40MG</u>	<u>A202728 001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A202728 002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A202728 003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A202728 004</u>	Jan 05, 2015
<u>AB</u>	WATSON LABS INC	<u>40MG</u>	<u>A090642 001</u>	Jan 05, 2015
<u>AB</u>		<u>80MG</u>	<u>A090642 002</u>	Jan 05, 2015
<u>AB</u>		<u>160MG</u>	<u>A090642 003</u>	Jan 05, 2015
<u>AB</u>		<u>320MG</u>	<u>A090642 004</u>	Jan 05, 2015

VANCOMYCIN HYDROCHLORIDE

CAPSULE;ORAL

VANCOCIN HYDROCHLORIDE

<u>AB</u>	+	ANI PHARMS INC	<u>EQ 125MG BASE</u>	<u>N050606 001</u>	Apr 15, 1986
<u>AB</u>	!+		<u>EQ 250MG BASE</u>	<u>N050606 002</u>	Apr 15, 1986

VANCOMYCIN HYDROCHLORIDE

<u>AB</u>	AKORN	<u>EQ 125MG BASE</u>	<u>A065478 001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065478 002</u>	Apr 09, 2012
<u>AB</u>	FRESENIUS KABI USA	<u>EQ 125MG BASE</u>	<u>A065453 001</u>	Jun 18, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065453 002</u>	Jun 18, 2012
<u>AB</u>	LUPIN LTD	<u>EQ 125MG BASE</u>	<u>A090439 001</u>	Jan 28, 2015
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A090439 002</u>	Jan 28, 2015
<u>AB</u>	STRIDES PHARMA	<u>EQ 125MG BASE</u>	<u>A065490 001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065490 002</u>	Apr 09, 2012
<u>AB</u>	WATSON LABS	<u>EQ 125MG BASE</u>	<u>A065510 001</u>	Apr 09, 2012
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>A065510 002</u>	Apr 09, 2012

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A205780 001</u>	Mar 31, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A205780 002</u>	Mar 31, 2016
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A205779 001</u>	Mar 29, 2016
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A205779 002</u>	Mar 29, 2016
<u>AP</u>	CFT PHARMS LLC	<u>EQ 5GM BASE/VIAL</u>	<u>A204125 001</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A204125 002</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>A204107 001</u>	Dec 28, 2015
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A204107 002</u>	Dec 28, 2015
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A202275 001</u>	Oct 31, 2013
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A202275 002</u>	Oct 31, 2013
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A202464 001</u>	Oct 09, 2013
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A202274 001</u>	Oct 31, 2013
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A062663 001</u>	Mar 17, 1987
<u>AP</u>		<u>EQ 750MG BASE/VIAL</u>	<u>A062663 005</u>	Aug 17, 2016
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062663 002</u>	Jul 31, 1987
<u>AP</u>	!	<u>EQ 5GM BASE/VIAL</u>	<u>A062663 003</u>	Jun 03, 1988
<u>AP</u>	!	<u>EQ 10GM BASE/VIAL</u>	<u>A062663 004</u>	Nov 28, 1997
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A205694 001</u>	Jan 21, 2016
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A205694 002</u>	Jan 21, 2016
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A062911 001</u>	Aug 04, 1988
<u>AP</u>	!	<u>EQ 500MG BASE/VIAL</u>	<u>A062931 001</u>	Oct 29, 1992
<u>AP</u>	!	<u>EQ 750MG BASE/VIAL</u>	<u>A062912 002</u>	Jan 07, 2009
<u>AP</u>	!	<u>EQ 750MG BASE/VIAL</u>	<u>A062933 002</u>	May 27, 2009
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062912 001</u>	Aug 04, 1988
<u>AP</u>	!	<u>EQ 1GM BASE/VIAL</u>	<u>A062933 001</u>	Oct 29, 1992

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-425 (of 436)

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	!	<u>EQ 5GM BASE/VIAL</u>	<u>A063076 001</u>	Dec 21, 1990
<u>AP</u>	HOSPIRA INC	<u>EQ 10GM BASE/VIAL</u>	<u>A065455 001</u>	Apr 29, 2009
<u>AP</u>	MUSTAFA NEVZAT ILAC	<u>EQ 500MG BASE/VIAL</u>	<u>A065401 001</u>	Jun 30, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065401 002</u>	Jun 30, 2008
<u>AP</u>	MYLAN LABS LTD	<u>EQ 500MG BASE/VIAL</u>	<u>A065397 001</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A065397 002</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A065432 001</u>	Dec 30, 2008
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A091469 001</u>	Jul 01, 2011
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A091554 001</u>	Sep 19, 2011
<u>AP</u>	SAGENT PHARMS	<u>EQ 5GM BASE/VIAL</u>	<u>A200837 001</u>	Aug 10, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A200837 002</u>	Sep 02, 2014
<u>AP</u>	SANDOZ	<u>EQ 500MG BASE/VIAL</u>	<u>A090250 001</u>	Apr 27, 2010
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A090250 002</u>	Apr 27, 2010
<u>AP</u>	SANDOZ INC	<u>EQ 5GM BASE/VIAL</u>	<u>A201048 001</u>	Aug 10, 2012
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A201048 002</u>	Aug 10, 2012
<u>AP</u>	XELLIA PHARMS APS	<u>EQ 500MG BASE/VIAL</u>	<u>A091377 001</u>	Sep 09, 2015
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>A091377 002</u>	Sep 09, 2015
<u>AP</u>		<u>EQ 5GM BASE/VIAL</u>	<u>A206243 001</u>	Dec 23, 2015
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>A206243 002</u>	Dec 23, 2015

VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER

+!	BAXTER HLTHCARE	EQ 500MG BASE/100ML
+!		EQ 750MG BASE/150ML

N050671 001 Apr 29, 1993  
N050671 002 Dec 20, 2010

POWDER; IV (INFUSION)

VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER
SAMSON MEDCL

A091532 001 Jan 06, 2016

VANDETANIB

TABLET; ORAL

CAPRELSA

+ ! GENZYME CORP	100MG	N022405 001	Apr 06, 2011
+ !	300MG	N022405 002	Apr 06, 2011

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

+ ! BAYER HLTHCARE	2.5MG	N021400 003	Aug 19, 2003
+	5MG	N021400 001	Aug 19, 2003
+	10MG	N021400 002	Aug 19, 2003
+ !	20MG	N021400 004	Aug 19, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

STAXYN

+ ! BAYER HLTHCARE	10MG	N200179 001	Jun 17, 2010
--------------------	------	-------------	--------------

VARENICLINE TARTRATE

TABLET; ORAL

CHANTIX

+ ! PFIZER INC	EQ 0.5MG BASE	N021928 001	May 10, 2006
+ !	EQ 1MG BASE	N021928 002	May 10, 2006

VASOPRESSIN

SOLUTION; IV (INFUSION)

VASOSTRICT

+ ! PAR STERILE PRODUCTS	20UNITS/ML (20UNITS/ML)	N204485 001	Apr 17, 2014
+ !	200UNITS/10ML (20UNITS/ML)	N204485 002	Dec 17, 2016

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>	GLAND PHARMA LTD	<u>10MG/VIAL</u>	<u>A205390 001</u>	May 26, 2016
<u>AP</u>		<u>20MG/VIAL</u>	<u>A205390 002</u>	May 26, 2016
<u>AP</u>	HOSPIRA	<u>10MG/VIAL</u>	<u>A075164 001</u>	Oct 21, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A075164 002</u>	Oct 21, 1999
<u>AP</u>	MYLAN LABS LTD	<u>10MG/VIAL</u>	<u>A090243 001</u>	May 11, 2010
<u>AP</u>		<u>20MG/VIAL</u>	<u>A090243 002</u>	May 11, 2010
<u>AP</u>	SAGENT PHARMS	<u>10MG/VIAL</u>	<u>A078274 001</u>	Dec 29, 2008
<u>AP</u>		<u>20MG/VIAL</u>	<u>A078274 002</u>	Dec 29, 2008
<u>AP</u>	! SUN PHARMA GLOBAL	<u>10MG/VIAL</u>	<u>A079001 001</u>	Jun 17, 2009
<u>AP</u>	!	<u>20MG/VIAL</u>	<u>A079001 002</u>	Jun 17, 2009
<u>AP</u>	TEVA PHARMS USA	<u>10MG/VIAL</u>	<u>A074688 001</u>	Aug 25, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>A074688 002</u>	Aug 25, 1999
<u>AP</u>	WEST-WARD PHARMS	<u>10MG/VIAL</u>	<u>A075549 001</u>	Jun 13, 2000

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-426 (of 436)

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

INT

AP 20MG/VIAL

A075549 002 Jun 13, 2000

VELAGLUCERASE ALFA

INJECTABLE; IV (INFUSION)

VPRIV

SHIRE HUMAN GENETIC 400 UNITS/VIAL

N022575 001 Feb 26, 2010

VEMURAFENIB

TABLET; ORAL

ZELBORAF

+! HOFFMANN LA ROCHE 240MG

N202429 001 Aug 17, 2011

VENETOCLAX

TABLET; ORAL

VENCLEXTA

+ ABBVIE INC 10MG  
+ 50MG  
+! 100MG

N208573 001 Apr 11, 2016  
N208573 002 Apr 11, 2016  
N208573 003 Apr 11, 2016

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

AB + WYETH PHARMS INC EQ 37.5MG BASE  
AB + EQ 75MG BASE  
AB +! EQ 150MG BASE

N020699 001 Oct 20, 1997  
N020699 002 Oct 20, 1997  
N020699 004 Oct 20, 1997

VENLAFAXINE HYDROCHLORIDE

AB ANCHEN PHARMS EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A078087 001 Mar 16, 2012  
A078087 002 Mar 16, 2012  
A078087 003 Mar 16, 2012

AB AUROBINDO PHARMA LTD EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A200834 001 Apr 14, 2011  
A200834 002 Apr 14, 2011  
A200834 003 Apr 14, 2011

AB DR REDDYS LABS LTD EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A078421 001 May 06, 2011  
A078421 002 May 06, 2011  
A078421 003 May 06, 2011

AB MACLEODS PHARMS LTD EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A204889 001 Oct 05, 2017  
A204889 002 Oct 05, 2017  
A204889 003 Oct 05, 2017

AB ORCHID HLTHCARE EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A091123 001 Jul 11, 2011  
A091123 002 Jul 11, 2011  
A091123 003 Jul 11, 2011

AB TEVA EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A076565 001 Jun 28, 2010  
A076565 002 Jun 28, 2010  
A076565 003 Jun 28, 2010

AB TORRENT PHARMS LLC EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A090899 001 Jun 01, 2011  
A090899 002 Jun 01, 2011  
A090899 003 Jun 01, 2011

AB VALEANT PHARMS NORTH EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A090071 001 Apr 15, 2011  
A090071 002 Apr 15, 2011  
A090071 003 Apr 15, 2011

AB WOCKHARDT EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A078865 001 Apr 14, 2011  
A078865 002 Apr 14, 2011  
A078865 003 Apr 14, 2011

AB ZYDUS PHARMS USA INC EQ 37.5MG BASE  
AB EQ 75MG BASE  
AB EQ 150MG BASE

A090174 001 Apr 14, 2011  
A090174 002 Apr 14, 2011  
A090174 003 Apr 14, 2011

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

AB ALEMBIC PHARMS LTD EQ 25MG BASE  
AB EQ 37.5MG BASE

A078932 001 Dec 14, 2010  
A078932 002 Dec 14, 2010

AB AMNEAL PHARMS EQ 50MG BASE  
AB EQ 75MG BASE  
AB EQ 100MG BASE

A078932 003 Dec 14, 2010  
A078932 004 Dec 14, 2010  
A078932 005 Dec 14, 2010

AB EQ 25MG BASE  
AB EQ 37.5MG BASE  
AB EQ 50MG BASE  
AB EQ 75MG BASE  
AB EQ 100MG BASE

A079098 001 May 11, 2010  
A079098 002 May 11, 2010  
A079098 003 May 11, 2010  
A079098 004 May 11, 2010  
A079098 005 May 11, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-427 (of 436)

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	AUROBINDO PHARMA	<u>EQ 25MG BASE</u>	<u>A090555 001</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A090555 002</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090555 003</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090555 004</u>	Apr 07, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090555 005</u>	Apr 07, 2010
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 25MG BASE</u>	<u>A078301 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078301 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078301 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078301 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078301 005</u>	Jun 13, 2008
<u>AB</u>	HERITAGE PHARMS INC	<u>EQ 25MG BASE</u>	<u>A078554 001</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078554 002</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078554 003</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078554 004</u>	Jan 09, 2009
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078554 005</u>	Jan 09, 2009
<u>AB</u>	MYLAN	<u>EQ 25MG BASE</u>	<u>A077166 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A077166 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077166 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A077166 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077166 005</u>	Jun 13, 2008
<u>AB</u>	PRINSTON INC	<u>EQ 25MG BASE</u>	<u>A090027 001</u>	Aug 04, 2010
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A090027 002</u>	Aug 04, 2010
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A090027 003</u>	Aug 04, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A090027 004</u>	Aug 04, 2010
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A090027 005</u>	Aug 04, 2010
<u>AB</u>	SUN PHARM IND'S INC	<u>EQ 25MG BASE</u>	<u>A078627 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A078627 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A078627 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A078627 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A078627 005</u>	Jun 13, 2008
<u>AB</u>	TEVA	<u>EQ 25MG BASE</u>	<u>A076690 001</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A076690 002</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A076690 003</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A076690 004</u>	Aug 03, 2006
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A076690 005</u>	Aug 03, 2006
<u>AB</u>	YAOPHARMA CO LTD	<u>EQ 25MG BASE</u>	<u>A202036 001</u>	May 28, 2015
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A202036 002</u>	May 28, 2015
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A202036 003</u>	May 28, 2015
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A202036 004</u>	May 28, 2015
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A202036 005</u>	May 28, 2015
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 25MG BASE</u>	<u>A077653 001</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 37.5MG BASE</u>	<u>A077653 002</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>A077653 003</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A077653 004</u>	Jun 13, 2008
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>A077653 005</u>	Jun 13, 2008

TABLET, EXTENDED RELEASE; ORAL

VENLAFAXINE HYDROCHLORIDE

<u>AB</u>	NOSTRUM LABS INC	<u>EQ 150MG BASE</u>	<u>A205468 002</u>	Mar 24, 2017
<u>AB</u>		<u>EQ 225MG BASE</u>	<u>A205468 003</u>	Mar 24, 2017
<u>AB</u>	+	<u>OSMOTICA PHARM</u>	<u>N022104 001</u>	May 20, 2008
<u>AB</u>	+		<u>N022104 002</u>	May 20, 2008
<u>AB</u>	!+		<u>N022104 003</u>	May 20, 2008
<u>AB</u>	+		<u>N022104 004</u>	May 20, 2008
<u>AB</u>	SUN PHARMA GLOBAL	<u>EQ 37.5MG BASE</u>	<u>A091272 001</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>A091272 002</u>	Aug 18, 2010
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>A091272 003</u>	Aug 18, 2010

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>100MG</u>	<u>A078306 001</u>	Aug 09, 2007
<u>AB</u>		<u>120MG</u>	<u>A075138 001</u>	Apr 20, 1999
<u>AB</u>		<u>180MG</u>	<u>A075138 002</u>	Apr 20, 1999
<u>AB</u>		<u>200MG</u>	<u>A078306 002</u>	Aug 09, 2007
<u>AB</u>		<u>240MG</u>	<u>A075138 003</u>	Apr 20, 1999
<u>AB</u>		<u>300MG</u>	<u>A078306 003</u>	Aug 09, 2007
	<u>VERELAN</u>			
<u>AB</u>	RECRO GAINESVILLE	<u>120MG</u>	<u>N019614 001</u>	May 29, 1990
<u>AB</u>	+	<u>180MG</u>	<u>N019614 003</u>	Jan 09, 1992
<u>AB</u>	+	<u>240MG</u>	<u>N019614 002</u>	May 29, 1990

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-428 (of 436)

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

VERELAN PM

<u>AB</u>	+	RECRO GAINESVILLE	<u>100MG</u>	<u>N020943 001</u>	Nov 25, 1998
<u>AB</u>	+		<u>200MG</u>	<u>N020943 002</u>	Nov 25, 1998
<u>AB</u>	+!		<u>300MG</u>	<u>N020943 003</u>	Nov 25, 1998

VERELAN

+!	RECRO GAINESVILLE	360MG	N019614 004	May 10, 1996
----	-------------------	-------	-------------	--------------

SOLUTION;INTRAVENOUS

VERAPAMIL HYDROCHLORIDE

<u>AP</u>		EXELA PHARMA SCS LLC	<u>5MG/2ML (2.5MG/ML)</u>	<u>N018925 001</u>	Mar 30, 1984
<u>AP</u>	!	HOSPIRA	<u>10MG/4ML (2.5MG/ML)</u>	<u>A070738 001</u>	May 06, 1987
<u>AP</u>	!		<u>5MG/2ML (2.5MG/ML)</u>	<u>A075136 001</u>	Oct 20, 1998
<u>AP</u>	!		<u>5MG/2ML (2.5MG/ML)</u>	<u>A070737 001</u>	May 06, 1987
<u>AP</u>	!		<u>10MG/4ML (2.5MG/ML)</u>	<u>A070737 002</u>	May 06, 1987

TABLET;ORAL

CALAN

<u>AB</u>	+	GD SEARLE LLC	<u>80MG</u>	<u>N018817 001</u>	Sep 10, 1984
<u>AB</u>	+!		<u>120MG</u>	<u>N018817 002</u>	Sep 10, 1984

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		HERITAGE PHARMS INC	<u>40MG</u>	<u>A071881 002</u>	Oct 14, 2015
<u>AB</u>			<u>80MG</u>	<u>A071881 003</u>	Apr 05, 1988
<u>AB</u>			<u>120MG</u>	<u>A071881 001</u>	Apr 05, 1988
<u>AB</u>		MYLAN	<u>80MG</u>	<u>A071483 002</u>	Feb 15, 1989
<u>AB</u>			<u>120MG</u>	<u>A071483 001</u>	Feb 15, 1989
<u>AB</u>		WATSON LABS	<u>40MG</u>	<u>A072924 001</u>	Jun 29, 1993
<u>AB</u>			<u>80MG</u>	<u>A070995 001</u>	Oct 01, 1986
<u>AB</u>			<u>120MG</u>	<u>A070994 001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE;ORAL

CALAN SR

<u>AB</u>	+!	PFIZER	<u>120MG</u>	<u>N019152 003</u>	Mar 06, 1991
<u>AB</u>	+!		<u>240MG</u>	<u>N019152 001</u>	Dec 16, 1986

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		APOTEX CORP	<u>120MG</u>	<u>A200878 001</u>	Apr 20, 2012
<u>AB</u>			<u>180MG</u>	<u>A200878 002</u>	Apr 20, 2012
<u>AB</u>			<u>240MG</u>	<u>A200878 003</u>	Apr 20, 2012
<u>AB</u>		CADILA PHARMS LTD	<u>180MG</u>	<u>A206173 001</u>	May 05, 2017
<u>AB</u>			<u>240MG</u>	<u>A206173 002</u>	May 05, 2017
<u>AB</u>	!	GLENMARK GENERICS	<u>120MG</u>	<u>A090700 001</u>	Aug 03, 2011
<u>AB</u>	!		<u>180MG</u>	<u>A090700 002</u>	Aug 03, 2011
<u>AB</u>			<u>240MG</u>	<u>A078906 001</u>	Sep 17, 2009
<u>AB</u>		IVAX SUB TEVA PHARMS	<u>120MG</u>	<u>A073568 002</u>	Oct 10, 1997
<u>AB</u>			<u>180MG</u>	<u>A074330 001</u>	Jan 31, 1994
<u>AB</u>			<u>240MG</u>	<u>A073568 001</u>	Jul 31, 1992
<u>AB</u>		MYLAN	<u>120MG</u>	<u>A074587 002</u>	Feb 21, 1997
<u>AB</u>			<u>180MG</u>	<u>A074587 003</u>	Sep 09, 1997
<u>AB</u>			<u>240MG</u>	<u>A074587 001</u>	Mar 23, 1996
<u>AB</u>		PAR PHARM	<u>120MG</u>	<u>A075072 001</u>	May 25, 1999
<u>AB</u>			<u>240MG</u>	<u>A075072 003</u>	May 25, 1999
<u>AB</u>		SUN PHARM IND'S INC	<u>120MG</u>	<u>A090529 001</u>	Dec 30, 2011
<u>AB</u>			<u>180MG</u>	<u>A090529 002</u>	Dec 30, 2011
<u>AB</u>			<u>240MG</u>	<u>A090529 003</u>	Dec 30, 2011

VERTEPORFIN

INJECTABLE; INJECTION

VISUDYNE

+!	VALEANT LUXEMBOURG	15MG/VIAL	N021119 001	Apr 12, 2000
----	--------------------	-----------	-------------	--------------

VIGABATRIN

FOR SOLUTION;ORAL

SABRIL

<u>AA</u>	+!	LUNDBECK PHARMS LLC	<u>500MG/PACKET</u>	<u>N022006 001</u>	Aug 21, 2009
-----------	----	---------------------	---------------------	--------------------	--------------

VIGABATRIN

<u>AA</u>		PAR PHARM INC	<u>500MG/PACKET</u>	<u>A208218 001</u>	Apr 27, 2017
-----------	--	---------------	---------------------	--------------------	--------------

TABLET;ORAL

SABRIL

+!	LUNDBECK PHARMS LLC	500MG	N020427 001	Aug 21, 2009
----	---------------------	-------	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-429 (of 436)

VILAZODONE HYDROCHLORIDE

TABLET;ORAL

VIIBRYD

+!	FOREST LABS LLC	10MG	N022567 001 Jan 21, 2011
+		20MG	N022567 002 Jan 21, 2011
+		40MG	N022567 003 Jan 21, 2011

VINBLASTINE SULFATE

INJECTABLE;INJECTION

VINBLASTINE SULFATE

!	FRESENIUS KABI USA	1MG/ML	A089515 001 Apr 29, 1987
!	WEST-WARD PHARMS	10MG/VIAL INT	A089395 001 Apr 09, 1987

VINCRISTINE SULFATE

INJECTABLE;INJECTION

**VINCRISTINE SULFATE PFS**

<b>AP</b>	!	HOSPIRA	<b>1MG/ML</b>	<b>A071484 001</b> Apr 19, 1988
<b>AP</b>		TEVA PHARMS USA	<b>1MG/ML</b>	<b>A075493 001</b> Sep 01, 1999
INJECTABLE, LIPOSOMAL;INTRAVENOUS				
MARQIBO KIT				
+!	TALON THERAP		5MG/5ML (1MG/ML)	N202497 001 Aug 09, 2012

VINORELBINE TARTRATE

INJECTABLE;INJECTION

**NAVELBINE**

<b>AP</b>	+	PIERRE FABRE	<b>EQ 10MG BASE/ML</b>	<b>N020388 001</b> Dec 23, 1994
<b>VINORELBINE TARTRATE</b>				
<b>AP</b>		ACTAVIS TOTOWA	<b>EQ 10MG BASE/ML</b>	<b>A078011 001</b> Jul 22, 2009
<b>AP</b>		DR REDDYS LABS LTD	<b>EQ 10MG BASE/ML</b>	<b>A202017 001</b> Sep 12, 2013
<b>AP</b>		FRESENIUS KABI USA	<b>EQ 10MG BASE/ML</b>	<b>A076849 001</b> Apr 18, 2005
<b>AP</b>		HOSPIRA	<b>EQ 10MG BASE/ML</b>	<b>A076827 001</b> Jun 02, 2005
<b>AP</b>		JIANGSU HANSOH PHARM	<b>EQ 10MG BASE/ML</b>	<b>A091106 001</b> Sep 26, 2012
<b>AP</b>		TEVA PHARMS USA	<b>EQ 10MG BASE/ML</b>	<b>A076028 001</b> Feb 03, 2003
<b>AP</b>		WEST-WARD PHARMS INT	<b>EQ 10MG BASE/ML</b>	<b>A075992 001</b> Jun 10, 2003
<b>AP</b>			<b>EQ 10MG BASE/ML</b>	<b>A076461 001</b> Dec 11, 2003

VISMODEGIB

CAPSULE;ORAL

ERIVEDGE

+!	GENENTECH	150MG	N203388 001 Jan 30, 2012
----	-----------	-------	--------------------------

VITAMIN A PALMITATE

INJECTABLE;INJECTION

AQUASOL A

+!	HOSPIRA	EQ 50,000 UNITS BASE/ML	N006823 001
----	---------	-------------------------	-------------

VORAPAXAR SULFATE

TABLET;ORAL

ZONTIVITY

+!	ARALEZ PHARMS	EQ 2.08MG BASE	N204886 001 May 08, 2014
----	---------------	----------------	--------------------------

VORICONAZOLE

FOR SUSPENSION;ORAL

**VFEND**

<b>AB</b>	+	PF PRISM CV	<b>200MG/5ML</b>	<b>N021630 001</b> Dec 19, 2003
<b>VORICONAZOLE</b>				
<b>AB</b>		AMNEAL PHARMS	<b>200MG/5ML</b>	<b>A205034 001</b> Apr 13, 2016
<b>AB</b>		MYLAN PHARMS INC	<b>200MG/5ML</b>	<b>A202361 001</b> May 28, 2013
<b>AB</b>		NOVEL LABS INC	<b>200MG/5ML</b>	<b>A206799 001</b> May 31, 2016

INJECTABLE;IV (INFUSION)

**VFEND**

<b>AP</b>	+	PF PRISM CV	<b>200MG/VIAL</b>	<b>N021267 001</b> May 24, 2002
<b>VORICONAZOLE</b>				
<b>AP</b>		ALVOGEN INC	<b>200MG/VIAL</b>	<b>A206398 001</b> Mar 23, 2016
<b>AP</b>		SANDOZ INC	<b>200MG/VIAL</b>	<b>A090862 001</b> May 30, 2012

POWDER;IV (INFUSION)

VORICONAZOLE

XELLIA PHARMS APS	200MG/VIAL	N208562 001 Mar 09, 2017		
TABLET;ORAL				
<b>VFEND</b>				
<b>AB</b>	+	PF PRISM CV	<b>50MG</b>	<b>N021266 001</b> May 24, 2002

<b>AB</b>	+	PF PRISM CV	<b>200MG</b>	<b>N021266 002</b> May 24, 2002
-----------	---	-------------	--------------	---------------------------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-430 (of 436)

VORICONAZOLE

TABLET;ORAL

VORICONAZOLE

<u>AB</u>	AJANTA PHARMA LTD	<u>50MG</u>	<u>A206181 001</u>	May 24, 2016
<u>AB</u>		<u>200MG</u>	<u>A206181 002</u>	May 24, 2016
<u>AB</u>	AKORN	<u>50MG</u>	<u>A207049 001</u>	Sep 07, 2016
<u>AB</u>		<u>200MG</u>	<u>A207049 002</u>	Sep 07, 2016
<u>AB</u>	APPCO PHARMA LLC	<u>50MG</u>	<u>A206762 001</u>	May 24, 2016
<u>AB</u>		<u>200MG</u>	<u>A206762 002</u>	May 24, 2016
<u>AB</u>	AUROBINDO PHARMA LTD	<u>50MG</u>	<u>A206837 001</u>	Jan 22, 2016
<u>AB</u>		<u>200MG</u>	<u>A206837 002</u>	Jan 22, 2016
<u>AB</u>	GLENMARK PHARMS LTD	<u>50MG</u>	<u>A203503 001</u>	Sep 02, 2015
<u>AB</u>		<u>200MG</u>	<u>A203503 002</u>	Sep 02, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>50MG</u>	<u>A090547 001</u>	Apr 22, 2010
<u>AB</u>		<u>200MG</u>	<u>A090547 002</u>	Apr 22, 2010
<u>AB</u>	NOVEL LABS INC	<u>50MG</u>	<u>A207371 001</u>	May 24, 2016
<u>AB</u>		<u>200MG</u>	<u>A207371 002</u>	May 24, 2016
<u>AB</u>	PRINSTON INC	<u>50MG</u>	<u>A206654 001</u>	Aug 08, 2016
<u>AB</u>		<u>200MG</u>	<u>A206654 002</u>	Aug 08, 2016
<u>AB</u>	SANDOZ INC	<u>50MG</u>	<u>A200265 001</u>	Dec 12, 2011
<u>AB</u>		<u>200MG</u>	<u>A200265 002</u>	Dec 12, 2011
<u>AB</u>	TEVA PHARMS	<u>50MG</u>	<u>A091658 001</u>	Apr 06, 2012
<u>AB</u>		<u>200MG</u>	<u>A091658 002</u>	Apr 06, 2012
<u>AB</u>	ZYDUS PHARMS USA INC	<u>50MG</u>	<u>A206747 001</u>	May 24, 2016
<u>AB</u>		<u>200MG</u>	<u>A206747 002</u>	May 24, 2016

VORINOSTAT

CAPSULE;ORAL

ZOLINZA

+! MERCK

100MG

N021991 001 Oct 06, 2006

VORTIOXETINE HYDROBROMIDE

TABLET;ORAL

TRINTELLIX

+ TAKEDA PHARMS USA

EQ 5MG BASE

N204447 001 Sep 30, 2013

+

EQ 10MG BASE

N204447 002 Sep 30, 2013

+

EQ 20MG BASE

N204447 004 Sep 30, 2013

WARFARIN SODIUM

TABLET;ORAL

COUMADIN

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>1MG</u>	<u>N009218 022</u>	Mar 01, 1990
<u>AB</u>	+		<u>2MG</u>	<u>N009218 013</u>	
<u>AB</u>	+		<u>2.5MG</u>	<u>N009218 018</u>	
<u>AB</u>	+		<u>3MG</u>	<u>N009218 025</u>	Nov 18, 1996
<u>AB</u>	+		<u>4MG</u>	<u>N009218 023</u>	Aug 24, 1993
<u>AB</u>	+		<u>5MG</u>	<u>N009218 007</u>	
<u>AB</u>	+		<u>6MG</u>	<u>N009218 026</u>	Nov 18, 1996
<u>AB</u>	+		<u>7.5MG</u>	<u>N009218 016</u>	
<u>AB</u>	+		<u>10MG</u>	<u>N009218 005</u>	

JANTOVEN

<u>AB</u>	USL PHARMA	<u>1MG</u>	<u>A040416 001</u>	Oct 02, 2003
<u>AB</u>		<u>2MG</u>	<u>A040416 002</u>	Oct 02, 2003
<u>AB</u>		<u>2.5MG</u>	<u>A040416 003</u>	Oct 02, 2003
<u>AB</u>		<u>3MG</u>	<u>A040416 004</u>	Oct 02, 2003
<u>AB</u>		<u>4MG</u>	<u>A040416 005</u>	Oct 02, 2003
<u>AB</u>		<u>5MG</u>	<u>A040416 006</u>	Oct 02, 2003
<u>AB</u>		<u>6MG</u>	<u>A040416 007</u>	Oct 02, 2003
<u>AB</u>		<u>7.5MG</u>	<u>A040416 008</u>	Oct 02, 2003
<u>AB</u>		<u>10MG</u>	<u>A040416 009</u>	Oct 02, 2003

WARFARIN SODIUM

<u>AB</u>	AMNEAL PHARMS	<u>1MG</u>	<u>A202202 001</u>	Mar 04, 2013
<u>AB</u>		<u>2MG</u>	<u>A202202 002</u>	Mar 04, 2013
<u>AB</u>		<u>2.5MG</u>	<u>A202202 003</u>	Mar 04, 2013
<u>AB</u>		<u>3MG</u>	<u>A202202 004</u>	Mar 04, 2013
<u>AB</u>		<u>4MG</u>	<u>A202202 005</u>	Mar 04, 2013
<u>AB</u>		<u>5MG</u>	<u>A202202 006</u>	Mar 04, 2013
<u>AB</u>		<u>6MG</u>	<u>A202202 007</u>	Mar 04, 2013
<u>AB</u>		<u>7.5MG</u>	<u>A202202 008</u>	Mar 04, 2013
<u>AB</u>		<u>10MG</u>	<u>A202202 009</u>	Mar 04, 2013
<u>AB</u>	BARR	<u>1MG</u>	<u>A040145 001</u>	Mar 26, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-431 (of 436)

WARFARIN SODIUM

TABLET;ORAL

WARFARIN SODIUM

<u>AB</u>	<u>2MG</u>	<u>A040145 002</u>	Mar 26, 1997
<u>AB</u>	<u>2 . 5MG</u>	<u>A040145 003</u>	Mar 26, 1997
<u>AB</u>	<u>3MG</u>	<u>A040145 008</u>	Nov 05, 1998
<u>AB</u>	<u>4MG</u>	<u>A040145 004</u>	Mar 26, 1997
<u>AB</u>	<u>5MG</u>	<u>A040145 005</u>	Mar 26, 1997
<u>AB</u>	<u>6MG</u>	<u>A040145 009</u>	Nov 05, 1998
<u>AB</u>	<u>7 . 5MG</u>	<u>A040145 006</u>	Mar 26, 1997
<u>AB</u>	<u>10MG</u>	<u>A040145 007</u>	Mar 26, 1997
<u>AB</u>	<u>1MG</u>	<u>A090935 001</u>	May 25, 2011
<u>AB</u>	<u>2MG</u>	<u>A090935 002</u>	May 25, 2011
<u>AB</u>	<u>2 . 5MG</u>	<u>A090935 003</u>	May 25, 2011
<u>AB</u>	<u>3MG</u>	<u>A090935 004</u>	May 25, 2011
<u>AB</u>	<u>4MG</u>	<u>A090935 005</u>	May 25, 2011
<u>AB</u>	<u>5MG</u>	<u>A090935 006</u>	May 25, 2011
<u>AB</u>	<u>6MG</u>	<u>A090935 007</u>	May 25, 2011
<u>AB</u>	<u>7 . 5MG</u>	<u>A090935 008</u>	May 25, 2011
<u>AB</u>	<u>10MG</u>	<u>A090935 009</u>	May 25, 2011
<u>AB</u>	<u>1MG</u>	<u>A200104 001</u>	Jun 27, 2013
<u>AB</u>	<u>2MG</u>	<u>A200104 002</u>	Jun 27, 2013
<u>AB</u>	<u>2 . 5MG</u>	<u>A200104 003</u>	Jun 27, 2013
<u>AB</u>	<u>3MG</u>	<u>A200104 004</u>	Jun 27, 2013
<u>AB</u>	<u>4MG</u>	<u>A200104 005</u>	Jun 27, 2013
<u>AB</u>	<u>5MG</u>	<u>A200104 006</u>	Jun 27, 2013
<u>AB</u>	<u>6MG</u>	<u>A200104 007</u>	Jun 27, 2013
<u>AB</u>	<u>7 . 5MG</u>	<u>A200104 008</u>	Jun 27, 2013
<u>AB</u>	<u>10MG</u>	<u>A200104 009</u>	Jun 27, 2013
<u>AB</u>	<u>1MG</u>	<u>A040616 009</u>	Jul 05, 2006
<u>AB</u>	<u>2MG</u>	<u>A040616 001</u>	Jul 05, 2006
<u>AB</u>	<u>2 . 5MG</u>	<u>A040616 002</u>	Jul 05, 2006
<u>AB</u>	<u>3MG</u>	<u>A040616 003</u>	Jul 05, 2006
<u>AB</u>	<u>4MG</u>	<u>A040616 004</u>	Jul 05, 2006
<u>AB</u>	<u>5MG</u>	<u>A040616 005</u>	Jul 05, 2006
<u>AB</u>	<u>6MG</u>	<u>A040616 006</u>	Jul 05, 2006
<u>AB</u>	<u>7 . 5MG</u>	<u>A040616 007</u>	Jul 05, 2006
<u>AB</u>	<u>10MG</u>	<u>A040616 008</u>	Jul 05, 2006
<u>AB</u>	<u>1MG</u>	<u>A040301 002</u>	Jul 15, 1999
<u>AB</u>	<u>2MG</u>	<u>A040301 003</u>	Jul 15, 1999
<u>AB</u>	<u>2 . 5MG</u>	<u>A040301 004</u>	Jul 15, 1999
<u>AB</u>	<u>3MG</u>	<u>A040301 005</u>	Jul 15, 1999
<u>AB</u>	<u>4MG</u>	<u>A040301 006</u>	Jul 15, 1999
<u>AB</u>	<u>5MG</u>	<u>A040301 007</u>	Jul 15, 1999
<u>AB</u>	<u>6MG</u>	<u>A040301 008</u>	Jul 15, 1999
<u>AB</u>	<u>7 . 5MG</u>	<u>A040301 009</u>	Jul 15, 1999
<u>AB</u>	<u>10MG</u>	<u>A040301 001</u>	Jul 15, 1999
<u>AB</u>	<u>1MG</u>	<u>A040663 001</u>	May 30, 2006
<u>AB</u>	<u>2MG</u>	<u>A040663 002</u>	May 30, 2006
<u>AB</u>	<u>2 . 5MG</u>	<u>A040663 003</u>	May 30, 2006
<u>AB</u>	<u>3MG</u>	<u>A040663 004</u>	May 30, 2006
<u>AB</u>	<u>4MG</u>	<u>A040663 005</u>	May 30, 2006
<u>AB</u>	<u>5MG</u>	<u>A040663 006</u>	May 30, 2006
<u>AB</u>	<u>6MG</u>	<u>A040663 007</u>	May 30, 2006
<u>AB</u>	<u>7 . 5MG</u>	<u>A040663 008</u>	May 30, 2006
<u>AB</u>	<u>10MG</u>	<u>A040663 009</u>	May 30, 2006

XENON XE-133

GAS;INHALATION

XENON XE 133

LANTHEUS MEDCL	10mCi/VIAL 20mCi/VIAL	N017284 001 N017284 002
MALLINKRODT NUCLEAR	10mCi/VIAL 20mCi/VIAL	N018327 001 Mar 09, 1982 N018327 002 Mar 09, 1982

ZAFIRLUKAST

TABLET;ORAL

ACCOLATE

<u>AB</u>	<u>+ PAR PHARM INC</u>	<u>10MG</u>	<u>N020547 003</u>	Sep 17, 1999
<u>AB</u>	<u>+!</u>	<u>20MG</u>	<u>N020547 001</u>	Sep 26, 1996
<u>ZAFIRLUKAST</u>				
<u>AB</u>	<u>DR REDDYS LABS LTD</u>	<u>10MG</u>	<u>A090372 001</u>	Nov 18, 2010
<u>AB</u>		<u>20MG</u>	<u>A090372 002</u>	Nov 18, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-432 (of 436)

ZALEPLON

CAPSULE;ORAL

SONATA

<u>AB</u>	+	Pfizer	<u>5MG</u>	<u>N020859 001</u>	Aug 13, 1999
<u>AB</u>	+!		<u>10MG</u>	<u>N020859 002</u>	Aug 13, 1999
<u>ZALEPLON</u>					
<u>AB</u>		AUROBINDO PHARMA	<u>5MG</u>	<u>A078829 001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A078829 002</u>	Jun 06, 2008
<u>AB</u>		CIPLA LTD	<u>5MG</u>	<u>A077505 001</u>	Jun 20, 2008
<u>AB</u>			<u>10MG</u>	<u>A077505 002</u>	Jun 20, 2008
<u>AB</u>		HIKMA PHARMS	<u>5MG</u>	<u>A078147 001</u>	Nov 25, 2008
<u>AB</u>			<u>10MG</u>	<u>A078147 002</u>	Nov 25, 2008
<u>AB</u>		ORCHID HLTHCARE	<u>5MG</u>	<u>A090374 001</u>	Sep 17, 2009
<u>AB</u>			<u>10MG</u>	<u>A090374 002</u>	Sep 17, 2009
<u>AB</u>		TEVA PHARMS	<u>5MG</u>	<u>A077239 001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A077239 002</u>	Jun 06, 2008
<u>AB</u>		UNICHEM	<u>5MG</u>	<u>A078989 001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A078989 002</u>	Jun 06, 2008
<u>AB</u>		WEST-WARD PHARMS INT	<u>5MG</u>	<u>A077237 001</u>	Jun 06, 2008
<u>AB</u>			<u>10MG</u>	<u>A077237 002</u>	Jun 06, 2008

ZANAMIVIR

POWDER;INHALATION

RELENZA

+!	GLAXOSMITHKLINE	5MG
----	-----------------	-----

N021036 001 Jul 26, 1999

ZICONOTIDE ACETATE

INJECTABLE;INTRATHECAL

PRIALT

+!	JAZZ PHARMS INTL	100MCG/1ML (100MCG/ML)
+!		500MCG/20ML (25MCG/ML)
+!		500MCG/5ML (100MCG/ML)

N021060 002 Dec 28, 2004  
 N021060 001 Dec 28, 2004  
 N021060 004 Dec 28, 2004

ZIDOVUDINE

CAPSULE;ORAL

RETROVIR

<u>AB</u>	+!	VIIV HLTHCARE	<u>100MG</u>	<u>N019655 001</u>	Mar 19, 1987
<u>ZIDOVUDINE</u>					
<u>AB</u>		AUROBINDO PHARMA LTD	<u>100MG</u>	<u>A078128 001</u>	Mar 27, 2006
<u>AB</u>		CIPLA LTD	<u>100MG</u>	<u>A078349 001</u>	May 23, 2007
INJECTABLE;INJECTION					
<u>RETROVIR</u>					
<u>AP</u>	+!	VIIV HLTHCARE	<u>10MG/ML</u>	<u>N019951 001</u>	Feb 02, 1990
<u>ZIDOVUDINE</u>					
<u>AP</u>		LUITPOLD SYRUP;ORAL	<u>10MG/ML</u>	<u>A091457 001</u>	May 06, 2010
<u>RETROVIR</u>					
<u>AA</u>	+!	VIIV HLTHCARE	<u>50MG/5ML</u>	<u>N019910 001</u>	Sep 28, 1989
<u>ZIDOVUDINE</u>					
<u>AA</u>		AUROBINDO	<u>50MG/5ML</u>	<u>A077268 001</u>	Sep 19, 2005
<u>AA</u>		CIPLA LTD	<u>50MG/5ML</u>	<u>A077981 001</u>	Jun 26, 2008
TABLET;ORAL					
<u>ZIDOVUDINE</u>					
<u>AB</u>		AUROBINDO	<u>300MG</u>	<u>A077267 001</u>	Sep 19, 2005
<u>AB</u>		CIPLA	<u>300MG</u>	<u>A090561 001</u>	Oct 27, 2010
<u>AB</u>	!	HETERO LABS LTD III	<u>300MG</u>	<u>A090092 001</u>	Apr 25, 2008
<u>AB</u>		MYLAN PHARMS INC	<u>300MG</u>	<u>A078922 001</u>	Feb 14, 2008
<u>AB</u>		WEST-WARD PHARMS INT	<u>300MG</u>	<u>A076844 001</u>	Sep 19, 2005

ZILEUTON

TABLET;ORAL

ZYFLO

+!	CHIESI USA INC	600MG	N020471 003	Dec 09, 1996
TABLET, EXTENDED RELEASE;ORAL				

ZILEUTON

<u>AB</u>		RISING PHARMS INC	<u>600MG</u>	<u>A204929 001</u>	Mar 17, 2017
<u>ZYFLO CR</u>					
<u>AB</u>	+!	CHIESI USA INC	<u>600MG</u>	<u>N022052 001</u>	May 30, 2007

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-433 (of 436)

ZINC ACETATE

CAPSULE; ORAL

GALZIN

+ ! TEVA  
+ !

EQ 25MG ZINC  
EQ 50MG ZINC

N020458 001 Jan 28, 1997  
N020458 002 Jan 28, 1997

ZINC CHLORIDE

INJECTABLE; INJECTION

ZINC CHLORIDE IN PLASTIC CONTAINER

+ ! HOSPIRA

EQ 1MG ZINC/ML

N018959 001 Jun 26, 1986

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON

AB + ! PFIZER

EQ 20MG BASE

N020825 001 Feb 05, 2001

AB +

EQ 40MG BASE

N020825 002 Feb 05, 2001

AB +

EQ 60MG BASE

N020825 003 Feb 05, 2001

AB +

EQ 80MG BASE

N020825 004 Feb 05, 2001

ZIPRASIDONE HYDROCHLORIDE

AB APOTEX INC

EQ 20MG BASE

A077561 001 Mar 02, 2012

AB +

EQ 40MG BASE

A077561 002 Mar 02, 2012

AB +

EQ 60MG BASE

A077561 003 Mar 02, 2012

AB +

EQ 80MG BASE

A077561 004 Mar 02, 2012

AB AUROBINDO PHARMA LTD

EQ 20MG BASE

A204117 001 Dec 27, 2016

AB +

EQ 40MG BASE

A204117 002 Dec 27, 2016

AB +

EQ 60MG BASE

A204117 003 Dec 27, 2016

AB +

EQ 80MG BASE

A204117 004 Dec 27, 2016

AB DR REDDYS LABS INC

EQ 20MG BASE

A077565 001 Mar 02, 2012

AB +

EQ 40MG BASE

A077565 002 Mar 02, 2012

AB +

EQ 60MG BASE

A077565 003 Mar 02, 2012

AB +

EQ 80MG BASE

A077565 004 Mar 02, 2012

AB LUPIN PHARMS

EQ 20MG BASE

A077560 001 Mar 02, 2012

AB +

EQ 40MG BASE

A077560 002 Mar 02, 2012

AB +

EQ 60MG BASE

A077560 003 Mar 02, 2012

AB +

EQ 80MG BASE

A077560 004 Mar 02, 2012

AB MACLEODS PHARMS LTD

EQ 20MG BASE

A204375 001 Feb 17, 2017

AB +

EQ 40MG BASE

A204375 002 Feb 17, 2017

AB +

EQ 60MG BASE

A204375 003 Feb 17, 2017

AB +

EQ 80MG BASE

A204375 004 Feb 17, 2017

AB MYLAN PHARMS INC

EQ 20MG BASE

A202395 001 Oct 10, 2013

AB +

EQ 40MG BASE

A202395 002 Oct 10, 2013

AB +

EQ 60MG BASE

A202395 003 Oct 10, 2013

AB +

EQ 80MG BASE

A202395 004 Oct 10, 2013

AB SANDOZ INC

EQ 20MG BASE

A077562 001 Jun 01, 2012

AB +

EQ 40MG BASE

A077562 002 Jun 01, 2012

AB +

EQ 60MG BASE

A077562 003 Jun 01, 2012

AB +

EQ 80MG BASE

A077562 004 Jun 01, 2012

AB WOCKHARDT LTD

EQ 20MG BASE

A090348 001 Sep 05, 2012

AB +

EQ 40MG BASE

A090348 002 Sep 05, 2012

AB +

EQ 60MG BASE

A090348 003 Sep 05, 2012

AB +

EQ 80MG BASE

A090348 004 Sep 05, 2012

AB ZYDUS PHARMS USA INC

EQ 20MG BASE

A208988 001 Aug 22, 2017

AB +

EQ 40MG BASE

A208988 002 Aug 22, 2017

AB +

EQ 60MG BASE

A208988 003 Aug 22, 2017

AB +

EQ 80MG BASE

A208988 004 Aug 22, 2017

ZIPRASIDONE MESYLATE

INJECTABLE; INTRAMUSCULAR

GEODON

+ ! PFIZER

EQ 20MG BASE/ML

N020919 001 Jun 21, 2002

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

RECLAST

AP + ! NOVARTIS

EQ 5MG BASE/100ML

N021817 001 Apr 16, 2007

ZOLEDRONIC ACID

ACCORD HLTHCARE

EQ 4MG BASE/5ML

A205279 001 Nov 28, 2016

ACS DOBFAR INFO SA

EQ 4MG BASE/100ML

N203231 001 Aug 02, 2013

ACTAVIS INC

EQ 5MG BASE/100ML

A202828 001 Sep 23, 2013

AKORN

EQ 4MG BASE/5ML

A202472 001 Mar 04, 2013

AKORN INC

EQ 5MG BASE/100ML

A200918 001 Aug 21, 2014

APOTEX INC

EQ 4MG BASE/5ML

A202548 001 May 22, 2014

APOTEX INC

EQ 5MG BASE/100ML

A204367 001 Dec 24, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-434 (of 436)

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

ZOLEDRONIC ACID

<u>AP</u>	AUROBINDO PHARMA LTD	<u>EQ 4MG BASE/5ML</u>	<u>A207751 001</u>	Sep 26, 2016
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A209125 001</u>	Dec 08, 2017
<u>AP</u>	BPI LABS LLC	<u>EQ 4MG BASE/5ML</u>	<u>A207341 001</u>	Dec 29, 2017
<u>AP</u>	BRECKENRIDGE PHARM	<u>EQ 4MG BASE/5ML</u>	<u>A091170 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 4MG BASE/5ML</u>	<u>A202571 001</u>	May 07, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202163 001</u>	Aug 05, 2013
<u>AP</u>	CIPLA LTD	<u>EQ 4MG BASE/100ML</u>	<u>A210174 001</u>	Oct 27, 2017
<u>AP</u>	DR REDDYS LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A091186 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A091363 001</u>	Mar 29, 2013
<u>AP</u>	EMCURE PHARMS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A201783 001</u>	Mar 12, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A201801 001</u>	Mar 29, 2013
<u>AP</u>	FRESENIUS KABI USA	<u>EQ 4MG BASE/5ML</u>	<u>A091516 001</u>	Apr 23, 2015
<u>AP</u>	GLAND PHARMA LTD	<u>EQ 4MG BASE/5ML</u>	<u>A202930 001</u>	Aug 05, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A204217 001</u>	Aug 18, 2016
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 4MG BASE/5ML</u>	<u>A202182 001</u>	Jun 03, 2013
<u>AP</u>	HOSPIRA INC	<u>EQ 4MG BASE/5ML</u>	<u>A090621 001</u>	Mar 19, 2015
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A202837 001</u>	Apr 05, 2013
<u>AP</u>	MYLAN LABS LTD	<u>EQ 4MG BASE/5ML</u>	<u>A202650 001</u>	Mar 04, 2013
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A203841 001</u>	Feb 14, 2017
<u>AP</u>		<u>EQ 5MG BASE/100ML</u>	<u>A205254 001</u>	Oct 27, 2017
<u>AP</u>	SAGENT PHARMS	<u>EQ 4MG BASE/5ML</u>	<u>A091493 001</u>	Nov 24, 2014
<u>AP</u>	! SUN PHARMA GLOBAL	<u>EQ 4MG BASE/VIAL</u>	<u>A090018 001</u>	Mar 04, 2013
<u>AP</u>	USV NORTH AMERICA	<u>EQ 4MG BASE/5ML</u>	<u>A202923 001</u>	Sep 04, 2014
<u>ZOMETA</u>				
<u>AP</u>	+! NOVARTIS	<u>EQ 4MG BASE/5ML</u>	<u>N021223 002</u>	Mar 07, 2003
<u>AP</u>	+!	<u>EQ 4MG BASE/100ML</u>	<u>N021223 003</u>	Jun 17, 2011

SOLUTION; IV (INFUSION)

ZOLEDRONIC ACID

HOSPIRA INC

EQ 4MG BASE/100ML (EQ 0.04MG BASE/ML)

N204016 001 Dec 28, 2015

ZOLMITRIPTAN

SPRAY; NASAL

ZOMIG

+ ASTRAZENECA

2.5MG/SPRAY

N021450 003 Sep 16, 2013

++!

5MG/SPRAY

N021450 004 Sep 30, 2003

TABLET; ORAL

ZOLMITRIPTAN

<u>AB</u>	AJANTA PHARMA LTD	<u>2.5MG</u>	<u>A204041 001</u>	May 20, 2016
<u>AB</u>		<u>5MG</u>	<u>A204041 002</u>	May 20, 2016
<u>AB</u>	ALEMBIC PHARMS LTD	<u>2.5MG</u>	<u>A204232 001</u>	Sep 30, 2015
<u>AB</u>		<u>5MG</u>	<u>A204232 002</u>	Sep 30, 2015
<u>AB</u>	APOTEX INC	<u>2.5MG</u>	<u>A202078 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A202078 002</u>	May 14, 2013
<u>AB</u>	APPCO PHARMA LLC	<u>2.5MG</u>	<u>A206973 001</u>	Jun 30, 2017
<u>AB</u>		<u>5MG</u>	<u>A206973 002</u>	Jun 30, 2017
<u>AB</u>	AUROBINDO PHARMA LTD	<u>2.5MG</u>	<u>A207021 001</u>	May 11, 2016
<u>AB</u>		<u>5MG</u>	<u>A207021 002</u>	May 11, 2016
<u>AB</u>	GLENMARK GENERICS	<u>2.5MG</u>	<u>A201779 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A201779 002</u>	May 14, 2013
<u>AB</u>	INVAGEN PHARMS	<u>2.5MG</u>	<u>A204284 001</u>	Apr 09, 2014
<u>AB</u>		<u>5MG</u>	<u>A204284 002</u>	Apr 09, 2014
<u>AB</u>	JUBILANT GENERICS	<u>2.5MG</u>	<u>A202279 001</u>	Nov 20, 2014
<u>AB</u>		<u>5MG</u>	<u>A202279 002</u>	Nov 20, 2014
<u>AB</u>	MACLEODS PHARMS LTD	<u>2.5MG</u>	<u>A203772 001</u>	Sep 30, 2015
<u>AB</u>		<u>5MG</u>	<u>A203772 002</u>	Sep 30, 2015
<u>AB</u>	MYLAN PHARMS INC	<u>2.5MG</u>	<u>A203186 001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A203186 002</u>	May 14, 2013
<u>AB</u>	PLD ACQUISITIONS LLC	<u>2.5MG</u>	<u>A207867 001</u>	Feb 27, 2017
<u>AB</u>		<u>5MG</u>	<u>A207867 002</u>	Feb 27, 2017
<u>AB</u>	TEVA PHARMS USA	<u>2.5MG</u>	<u>A090861 001</u>	Mar 04, 2014
<u>AB</u>		<u>5MG</u>	<u>A090861 002</u>	Mar 04, 2014

ZOMIG

<u>AB</u>	+ IPR	<u>2.5MG</u>	<u>N020768 001</u>	Nov 25, 1997
<u>AB</u>	++!	<u>5MG</u>	<u>N020768 002</u>	Nov 25, 1997
TABLET, ORALLY DISINTEGRATING; ORAL				
<u>AB</u>	<u>ZOLMITRIPTAN</u>	<u>2.5MG</u>	<u>A205074 001</u>	Dec 01, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-435 (of 436)

ZOLMITRIPTAN

TABLET, ORALLY DISINTEGRATING;ORAL

ZOLMITRIPTAN

<u>AB</u>		<u>5MG</u>	<u>A205074</u> <u>002</u>	Dec 01, 2016
<u>AB</u>	APOTEX INC	<u>2 . 5MG</u>	<u>A202476</u> <u>001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A202476</u> <u>002</u>	May 14, 2013
<u>AB</u>	GLENMARK GENERICS	<u>2 . 5MG</u>	<u>A202560</u> <u>001</u>	May 14, 2013
<u>AB</u>		<u>5MG</u>	<u>A202560</u> <u>002</u>	May 14, 2013
<u>AB</u>	JUBILANT GENERICS	<u>2 . 5MG</u>	<u>A202956</u> <u>001</u>	Sep 17, 2015
<u>AB</u>		<u>5MG</u>	<u>A202956</u> <u>002</u>	Sep 17, 2015
<u>AB</u>	MACLEODS PHARMS LTD	<u>2 . 5MG</u>	<u>A204336</u> <u>001</u>	Oct 22, 2015
<u>AB</u>		<u>5MG</u>	<u>A204336</u> <u>002</u>	Oct 22, 2015
<u>AB</u>	ZYDUS PHARMS USA INC	<u>2 . 5MG</u>	<u>A202890</u> <u>001</u>	May 15, 2013
<u>AB</u>		<u>5MG</u>	<u>A202890</u> <u>002</u>	May 15, 2013

ZOMIG-ZMT

<u>AB</u> + ASTRAZENECA	<u>2 . 5MG</u>	<u>N021231</u> <u>001</u>	Feb 13, 2001
<u>AB</u> +!	<u>5MG</u>	<u>N021231</u> <u>002</u>	Sep 17, 2001

ZOLPIDEM TARTRATE

SPRAY, METERED;ORAL

ZOLPIMIST			
+!	MAGNA PHARMS	5MG/SPRAY	
TABLET;ORAL			N022196 001 Dec 19, 2008

AMBIEN

<u>AB</u> + SANOFI AVENTIS US	<u>5MG</u>	<u>N019908</u> <u>001</u>	Dec 16, 1992
<u>AB</u> +!	<u>10MG</u>	<u>N019908</u> <u>002</u>	Dec 16, 1992

ZOLPIDEM TARTRATE

<u>AB</u>	ACME LABS	<u>5MG</u>	<u>A077214</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077214</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	APOTEX INC	<u>5MG</u>	<u>A077884</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077884</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	AUROBINDO PHARMA	<u>5MG</u>	<u>A078413</u> <u>001</u>	May 04, 2007
<u>AB</u>		<u>10MG</u>	<u>A078413</u> <u>002</u>	May 04, 2007
<u>AB</u>	CIPLA LTD	<u>5MG</u>	<u>A077388</u> <u>001</u>	Jul 30, 2012
<u>AB</u>		<u>10MG</u>	<u>A077388</u> <u>002</u>	Jul 30, 2012
<u>AB</u>	INVAGEN PHARMS	<u>5MG</u>	<u>A078184</u> <u>001</u>	Sep 07, 2007
<u>AB</u>		<u>10MG</u>	<u>A078184</u> <u>002</u>	Sep 07, 2007
<u>AB</u>	MYLAN	<u>5MG</u>	<u>A076578</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A076578</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	SANDOZ INC	<u>5MG</u>	<u>A077322</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077322</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	SUN PHARM IND'S INC	<u>5MG</u>	<u>A077359</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077359</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	SUN PHARM IND'S LTD	<u>5MG</u>	<u>A078055</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A078055</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	TEVA	<u>5MG</u>	<u>A076410</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A076410</u> <u>002</u>	Apr 23, 2007
<u>AB</u>	TORRENT PHARMS	<u>5MG</u>	<u>A077903</u> <u>001</u>	Aug 17, 2007
<u>AB</u>		<u>10MG</u>	<u>A077903</u> <u>002</u>	Aug 17, 2007
<u>AB</u>	VINTAGE	<u>5MG</u>	<u>A078616</u> <u>001</u>	Nov 21, 2008
<u>AB</u>		<u>10MG</u>	<u>A078616</u> <u>002</u>	Nov 21, 2008
<u>AB</u>	WOCKHARDT	<u>5MG</u>	<u>A078426</u> <u>001</u>	May 15, 2007
<u>AB</u>		<u>10MG</u>	<u>A078426</u> <u>002</u>	May 15, 2007
<u>AB</u>	YUNG SHIN PHARM	<u>5MG</u>	<u>A077990</u> <u>001</u>	Apr 23, 2007
<u>AB</u>		<u>10MG</u>	<u>A077990</u> <u>002</u>	Apr 23, 2007

TABLET;SUBLINGUAL

EDLUAR

<u>AB</u> + MYLAN SPECIALITY LP	<u>5MG</u>	<u>N021997</u> <u>001</u>	Mar 13, 2009
<u>AB</u> +!	<u>10MG</u>	<u>N021997</u> <u>002</u>	Mar 13, 2009

INTERMEZZO

<u>AB</u> + PURDUE PHARMA	<u>1 . 75MG</u>	<u>N022328</u> <u>001</u>	Nov 23, 2011
<u>AB</u> +!	<u>3 . 5MG</u>	<u>N022328</u> <u>002</u>	Nov 23, 2011

ZOLPIDEM TARTRATE

<u>AB</u>	DR REDDYS LABS INC	<u>1 . 75MG</u>	<u>A204503</u> <u>001</u>	Nov 18, 2016
<u>AB</u>		<u>3 . 5MG</u>	<u>A204503</u> <u>002</u>	Nov 18, 2016
<u>AB</u>	MYLAN PHARMS INC	<u>5MG</u>	<u>A202657</u> <u>001</u>	Aug 08, 2016
<u>AB</u>		<u>10MG</u>	<u>A202657</u> <u>002</u>	Aug 08, 2016
<u>AB</u>	NOVEL LABS INC	<u>1 . 75MG</u>	<u>A204299</u> <u>001</u>	Jun 03, 2015
<u>AB</u>		<u>3 . 5MG</u>	<u>A204299</u> <u>002</u>	Jun 03, 2015
<u>AB</u>	PAR FORM	<u>5MG</u>	<u>A201509</u> <u>001</u>	Aug 01, 2016
<u>AB</u>		<u>10MG</u>	<u>A201509</u> <u>002</u>	Aug 01, 2016
<u>AB</u>	PAR PHARM INC	<u>1 . 75MG</u>	<u>A204229</u> <u>001</u>	Sep 11, 2017

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**PRESCRIPTION DRUG PRODUCT LIST**

3-436 (of 436)

ZOLPIDEM TARTRATE

TABLET; SUBLINGUAL

ZOLPIDEM TARTRATE

<u>AB</u>		<u>3.5MG</u>	<u>A204229 002</u>	Sep 11, 2017
	TABLET, EXTENDED RELEASE; ORAL			
<u>AB</u>	<u>AMBIEN CR</u>			
<u>AB</u>	+ SANOFI AVENTIS US	<u>6.25MG</u>	<u>N021774 002</u>	Sep 02, 2005
<u>AB</u>	+!	<u>12.5MG</u>	<u>N021774 001</u>	Sep 02, 2005
	<u>ZOLPIDEM TARTRATE</u>			
<u>AB</u>	ACTAVIS ELIZABETH	<u>6.25MG</u>	<u>A078179 002</u>	Oct 13, 2010
<u>AB</u>		<u>12.5MG</u>	<u>A078179 001</u>	Jun 06, 2011
<u>AB</u>	ACTAVIS LABS FL INC	<u>6.25MG</u>	<u>A090153 001</u>	Mar 25, 2013
<u>AB</u>		<u>12.5MG</u>	<u>A090153 002</u>	Mar 25, 2013
<u>AB</u>	ANCHEM PHARMS	<u>6.25MG</u>	<u>A078148 002</u>	Apr 14, 2011
<u>AB</u>		<u>12.5MG</u>	<u>A078148 001</u>	Dec 03, 2010
<u>AB</u>	APOTEX INC	<u>6.25MG</u>	<u>A200266 001</u>	Sep 10, 2013
<u>AB</u>		<u>12.5MG</u>	<u>A200266 002</u>	Sep 10, 2013
<u>AB</u>	LUPIN LTD	<u>6.25MG</u>	<u>A078970 001</u>	Sep 11, 2013
<u>AB</u>		<u>12.5MG</u>	<u>A078970 002</u>	Sep 11, 2013
<u>AB</u>	SANDOZ	<u>6.25MG</u>	<u>A090107 001</u>	Jul 01, 2011
<u>AB</u>		<u>12.5MG</u>	<u>A090107 002</u>	Jul 01, 2011
<u>AB</u>	SUN PHARMA GLOBAL	<u>6.25MG</u>	<u>A204170 001</u>	Jan 24, 2017
<u>AB</u>		<u>12.5MG</u>	<u>A204170 002</u>	Jan 24, 2017

ZONISAMIDE

CAPSULE; ORAL

<u>AB</u>	<u>ZONEGRAN</u>			
<u>AB</u>	+ SUNOVION PHARMS INC	<u>25MG</u>	<u>N020789 003</u>	Aug 22, 2003
<u>AB</u>	+!	<u>50MG</u>	<u>N020789 002</u>	Aug 22, 2003
<u>AB</u>		<u>100MG</u>	<u>N020789 001</u>	Mar 27, 2000

<u>AB</u>	<u>ZONISAMIDE</u>			
<u>AB</u>	APOTEX INC	<u>25MG</u>	<u>A077642 001</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>A077642 002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>A077642 003</u>	Dec 22, 2005
<u>AB</u>	BIONPHARMA INC	<u>25MG</u>	<u>A077813 001</u>	Aug 16, 2006
<u>AB</u>		<u>50MG</u>	<u>A077813 002</u>	Aug 16, 2006
<u>AB</u>		<u>100MG</u>	<u>A077813 003</u>	Aug 16, 2006
<u>AB</u>	GLENMARK GENERICS	<u>25MG</u>	<u>A077651 001</u>	Jan 30, 2006
<u>AB</u>		<u>50MG</u>	<u>A077651 002</u>	Jan 30, 2006
<u>AB</u>		<u>100MG</u>	<u>A077651 003</u>	Jan 30, 2006
<u>AB</u>	INVAGEN PHARMS	<u>25MG</u>	<u>A077869 001</u>	May 31, 2006
<u>AB</u>		<u>50MG</u>	<u>A077869 002</u>	May 31, 2006
<u>AB</u>		<u>100MG</u>	<u>A077869 003</u>	May 31, 2006
<u>AB</u>	MYLAN	<u>25MG</u>	<u>A077637 001</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>A077637 002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>A077637 003</u>	Dec 22, 2005
<u>AB</u>	SUN PHARM INDUS (IN)	<u>25MG</u>	<u>A077634 001</u>	Mar 17, 2006
<u>AB</u>		<u>50MG</u>	<u>A077634 002</u>	Mar 17, 2006
<u>AB</u>		<u>100MG</u>	<u>A077634 003</u>	Mar 17, 2006
<u>AB</u>	WOCKHARDT	<u>25MG</u>	<u>A077636 003</u>	Jul 27, 2006
<u>AB</u>		<u>50MG</u>	<u>A077636 002</u>	Jul 27, 2006
<u>AB</u>		<u>100MG</u>	<u>A077636 001</u>	Dec 22, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>A077625 001</u>	Oct 16, 2006
<u>AB</u>		<u>50MG</u>	<u>A077625 002</u>	Oct 16, 2006
<u>AB</u>		<u>100MG</u>	<u>A077625 003</u>	Oct 16, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-1 (of 20)

**ACETAMINOPHEN**

SUPPOSITORY;RECTAL

ACEPHEN

G AND W LABS	120MG 325MG 325MG 650MG 650MG	N018060 001 A072344 001 Mar 27, 1992 N018060 003 Dec 18, 1986 A072237 001 Mar 27, 1992 N018060 002
--------------	---	--

ACETAMINOPHEN

PERRIGO NEW YORK	120MG 650MG	A070607 001 Apr 06, 1987 A070608 001 Dec 01, 1986
+ TARO PHARMS NORTH	120MG 325MG	N018337 003 Sep 12, 1983 N018337 002
+!	650MG	N018337 001
INFANTS' FEVERALL		

+ TARO PHARMS NORTH 80MG

N018337 004 Aug 26, 1992

NEOPAP

POLYMEDICA	120MG	N016401 001
------------	-------	-------------

TABLET, EXTENDED RELEASE;ORAL

ACETAMINOPHEN

AUROBINDO PHARMA LTD	650MG	A207229 001 Nov 09, 2016
OHM LABS	650MG	A076200 001 Mar 19, 2002
PERRIGO	650MG	A075077 001 Feb 25, 2000
SUN PHARM INDS LTD	650MG	A078569 001 Dec 14, 2011

TYLENOL

+! J AND J CONSUMER INC	650MG	N019872 001 Jun 08, 1994
+!	650MG	N019872 002 Jan 11, 2001

**ACETAMINOPHEN; ASPIRIN; CAFFEINE**

TABLET;ORAL

ACETAMINOPHEN, ASPIRIN AND CAFFEINE PERRIGO	250MG;250MG;65MG	A075794 001 Nov 26, 2001
EXCEDRIN (MIGRAINE) +!	GLAXOSMITHKLINE CONS	N020802 001 Jan 14, 1998

**ADAPALENE**

GEL;TOPICAL

DIFFERIN

+! GALDERMA LABS LP	0.1%	N020380 002 Jul 08, 2016
---------------------	------	--------------------------

**ALCOHOL; CHLORHEXIDINE GLUCONATE**

SOLUTION;TOPICAL

AVAGARD +!	3M	61%;1%	N021074 001 Jun 07, 2001
------------	----	--------	--------------------------

**ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE**

TABLET, CHEWABLE;ORAL

GAVISCON		
+ SANOFI AVENTIS US	80MG;20MG	N018685 001 Dec 09, 1983
+!	160MG;40MG	N018685 002 Dec 09, 1983

**ASPIRIN**

CAPSULE;ORAL

ASPIRIN +!	PLX PHARMA	325MG	N203697 001 Jan 14, 2013
------------	------------	-------	--------------------------

**AVOBENZONE; ECAMSULE; OCTOCRYLENE**

CREAM;TOPICAL

ANTHELIOS SX +!	LOREAL USA	2%;2%;10%	N021502 001 Jul 21, 2006
CAPITAL SOLEIL 15 +!	LOREAL USA	2%;3%;10%	N021501 001 Oct 02, 2006

**AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE**

CREAM;TOPICAL

ANTHELIOS 20 +!	LOREAL USA	2%;2%;10%;2%	N021471 001 Oct 05, 2006
ANTHELIOS 40 +!	LOREAL USA	2%;3%;10%;5%	N022009 001 Mar 31, 2008 N022009 002 Oct 29, 2009

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-2 (of 20)

BENTOQUATAM

LOTION;TOPICAL		
IVY BLOCK		
+! STAND HOMEOPATH	5%	N020532 001 Aug 26, 1996

BRIMONIDINE TARTRATE

SOLUTION/DROPS;OPHTHALMIC		
LUMIFY		
+! BAUSCH AND LOMB INC	0.025%	N208144 001 Dec 22, 2017

BUDESONIDE

SPRAY, METERED;NASAL		
BUDESONIDE		
APOTEX INC	0.032MG/SPRAY	A078949 002 Nov 20, 2015
RHINOCORT ALLERGY		
+! ASTRAZENECA PHARMS	0.032MG/SPRAY	N020746 003 Mar 23, 2015

BUTENAFINE HYDROCHLORIDE

CREAM;TOPICAL		
BUTENAFINE HYDROCHLORIDE		
TARO	1%	A205181 001 Nov 16, 2017
LOTRIMIN ULTRA		
+! BAYER HEALTHCARE LLC	1%	N021307 001 Dec 07, 2001

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE;ORAL		
FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE		
PERRIGO R AND D	800MG;10MG;165MG	A077355 001 Feb 06, 2008
	800MG;10MG;165MG	A204782 001 Aug 29, 2016
PEPCID COMPLETE		
+! J AND J CONSUMER INC	800MG;10MG;165MG	N020958 001 Oct 16, 2000

CETIRIZINE HYDROCHLORIDE

CAPSULE;ORAL		
CETIRIZINE HYDROCHLORIDE ALLERGY		
APOTEX INC	10MG	A207235 001 Aug 12, 2016
+ BIONPHARMA INC	5MG	N022429 001 Jul 23, 2009
+!	10MG	N022429 004 Jul 23, 2009
STRIDES PHARMA	10MG	A205291 001 Jul 21, 2017
CETIRIZINE HYDROCHLORIDE HIVES RELIEF		
+ BIONPHARMA INC	5MG	N022429 003 Jul 23, 2009
+!	10MG	N022429 002 Jul 23, 2009

SYRUP;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY		
ALLIED PHARMA INC	5MG/5ML	A091327 001 Oct 17, 2011
AMNEAL PHARMS	5MG/5ML	A090765 002 Oct 07, 2009
AUROBINDO PHARMA	5MG/5ML	A090750 002 Feb 02, 2010
BIO PHARM INC	5MG/5ML	A090474 002 Mar 30, 2009
PERRIGO R AND D	5MG/5ML	A204226 001 Sep 09, 2013
	5MG/5ML	A090254 002 Apr 09, 2008
SILARX	5MG/5ML	A091130 001 Apr 22, 2011
TARO	5MG/5ML	A090182 002 Apr 22, 2008
	5MG/5ML	A201546 001 May 20, 2011
TRIS PHARMA INC	5MG/5ML	A090572 001 Nov 16, 2012

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

ALLIED PHARMA INC	5MG/5ML	A091327 002 Oct 17, 2011
AMNEAL PHARMS	5MG/5ML	A090765 001 Oct 07, 2009
AUROBINDO PHARMA	5MG/5ML	A090750 001 Feb 02, 2010
BIO PHARM INC	5MG/5ML	A090474 001 Mar 30, 2009
PERRIGO R AND D	5MG/5ML	A090254 001 Apr 09, 2008
SILARX	5MG/5ML	A091130 002 Apr 22, 2011
TARO	5MG/5ML	A090182 001 Apr 22, 2008
	5MG/5ML	A201546 002 May 20, 2011
TRIS PHARMA INC	5MG/5ML	A090572 002 Nov 16, 2012

CHILDREN'S ZYRTEC ALLERGY

+! J AND J CONSUMER INC	5MG/5ML	N022155 002 Nov 16, 2007
-------------------------	---------	--------------------------

CHILDREN'S ZYRTEC HIVES RELIEF

+! J AND J CONSUMER INC	5MG/5ML	N022155 001 Nov 16, 2007
-------------------------	---------	--------------------------

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS NY	5MG	A078780 001 Jan 21, 2010
	10MG	A078780 004 Jan 21, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-3 (of 20)

**CETIRIZINE HYDROCHLORIDE**

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

APOTEX INC	5MG	A078317 001	Dec 27, 2007
	10MG	A078317 002	Dec 27, 2007
AUROBINDO PHARMA LTD	5MG	A090760 001	Aug 05, 2015
	10MG	A090760 003	Aug 05, 2015
CIPLA LTD	5MG	A077318 001	Jul 25, 2013
	10MG	A077318 002	Jul 25, 2013
CONTRACT PHARMACAL	5MG	A076047 001	Dec 27, 2007
	10MG	A076047 002	Dec 27, 2007
DR REDDYS LABS LTD	5MG	A078343 004	Jan 15, 2008
	10MG	A078343 003	Jan 15, 2008
GRANULES INDIA LTD	10MG	A209274 001	Dec 22, 2017
IPCA LABS LTD	5MG	A202277 002	Mar 11, 2014
	10MG	A202277 004	Mar 11, 2014
JUBILANT CADISTA	5MG	A078933 001	Jun 15, 2010
	10MG	A078933 002	Jun 15, 2010
MYLAN	5MG	A076677 001	Dec 27, 2007
	10MG	A076677 002	Dec 27, 2007
ORCHID HLTHCARE	5MG	A078862 001	Feb 19, 2009
	10MG	A078862 002	Feb 19, 2009
PERRIGO R AND D	5MG	A078336 001	Dec 27, 2007
	10MG	A078336 002	Dec 27, 2007
SANDOZ	5MG	A077946 001	Dec 27, 2007
	10MG	A077946 002	Dec 27, 2007
SUN PHARM INDs INC	5MG	A077499 001	Dec 27, 2007
	10MG	A077499 002	Dec 27, 2007
SUN PHARM INDs LTD	5MG	A077498 001	Dec 27, 2007
	10MG	A077498 002	Dec 27, 2007
TARO	5MG	A078072 001	Jul 22, 2009
	5MG	A078072 003	Jul 22, 2009
TORRENT PHARMS LLC	5MG	A079191 001	Apr 15, 2010
	10MG	A079191 004	Apr 15, 2010
UNICHEM	5MG	A078680 003	Jun 26, 2009
	10MG	A078680 004	Jun 26, 2009
UNIQUE PHARM LABS	5MG	A077829 001	Aug 26, 2009
	10MG	A077829 004	Aug 26, 2009
WOCKHARDT	5MG	A078427 003	Dec 28, 2007
	10MG	A078427 004	Dec 28, 2007

CETIRIZINE HYDROCHLORIDE HIVES

DR REDDYS LABS LTD	5MG	A078343 001	Jan 15, 2008
	10MG	A078343 002	Jan 15, 2008
IPCA LABS LTD	5MG	A202277 001	Mar 11, 2014
	10MG	A202277 003	Mar 11, 2014
JUBILANT CADISTA	5MG	A078933 003	Jun 15, 2010
	10MG	A078933 004	Jun 15, 2010
MYLAN	5MG	A076677 004	Dec 27, 2007
	10MG	A076677 003	Dec 27, 2007
ORCHID HLTHCARE	5MG	A078862 003	Feb 19, 2009
	10MG	A078862 004	Feb 19, 2009
PERRIGO R AND D	5MG	A078336 003	Dec 27, 2007
	10MG	A078336 004	Dec 27, 2007
SUN PHARM INDs INC	5MG	A077499 003	Dec 27, 2007
	10MG	A077499 004	Dec 27, 2007
SUN PHARM INDs LTD	5MG	A077498 003	Dec 27, 2007
	10MG	A077498 004	Dec 27, 2007
UNICHEM	5MG	A078680 001	Jun 26, 2009
	10MG	A078680 002	Jun 26, 2009
UNIQUE PHARM LABS	5MG	A077829 003	Aug 26, 2009
	10MG	A077829 002	Aug 26, 2009

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS NY	5MG	A078780 003	Jan 21, 2010
	10MG	A078780 002	Jan 21, 2010
AUROBINDO PHARMA LTD	5MG	A090760 002	Aug 05, 2015
	10MG	A090760 004	Aug 05, 2015
TARO	10MG	A078072 002	Jul 22, 2009
	10MG	A078072 004	Jul 22, 2009
TORRENT PHARMS LLC	5MG	A079191 003	Apr 15, 2010
	10MG	A079191 002	Apr 15, 2010

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
OTC DRUG PRODUCT LIST

4-4 (of 20)

CETIRIZINE HYDROCHLORIDE

TABLET;ORAL

ZYRTEC ALLERGY

+ J AND J CONSUMER INC	5MG	N019835 003	Nov 16, 2007
+!	10MG	N019835 004	Nov 16, 2007
ZYRTEC HIVES RELIEF			
+ J AND J CONSUMER INC	5MG	N019835 005	Nov 16, 2007
+!	10MG	N019835 006	Nov 16, 2007

TABLET, CHEWABLE;ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

JUBILANT GENERICS	5MG	A091116 001	Feb 19, 2015
	10MG	A091116 002	Feb 19, 2015
NOVEL LABS INC	5MG	A206793 001	Mar 08, 2016
	10MG	A206793 002	Mar 08, 2016
SANDOZ	5MG	A078692 001	Feb 14, 2008
!	10MG	A078692 002	Feb 14, 2008
SUN PHARMA GLOBAL	5MG	A090142 001	Aug 30, 2011
	10MG	A090142 002	Aug 30, 2011

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

JUBILANT GENERICS	5MG	A091116 003	Feb 19, 2015
	10MG	A091116 004	Feb 19, 2015
SUN PHARMA GLOBAL	5MG	A090142 003	Aug 30, 2011
	10MG	A090142 004	Aug 30, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

PERRIGO R AND D	10MG	A205490 001	Sep 02, 2015
ZYRTEC ALLERGY			

+! J AND J CONSUMER INC	10MG	N022578 001	Sep 03, 2010
-------------------------	------	-------------	--------------

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

IVAX SUB TEVA	5MG;120MG	A077170 001	Feb 25, 2008
PHARMS			
SANDOZ	5MG;120MG	A077991 001	Mar 05, 2008
SUN PHARM INDs LTD	5MG;120MG	A090922 001	Sep 28, 2012

ZYRTEC-D 12 HOUR

+! J AND J CONSUMER INC	5MG;120MG	N021150 002	Nov 09, 2007
-------------------------	-----------	-------------	--------------

CHLORHEXIDINE GLUCONATE

AEROSOL, METERED;TOPICAL

EXIDINE

+! XTTRIUM	4%	N019127 001	Dec 24, 1984
------------	----	-------------	--------------

CLOTH;TOPICAL

CHLORHEXIDINE GLUCONATE

+! SAGE PRODS	2%	N021669 001	Apr 25, 2005
---------------	----	-------------	--------------

SOLUTION;TOPICAL

BRIAN CARE

SOAPCO	4%	A071419 001	Dec 17, 1987
--------	----	-------------	--------------

CHG SCRUB

ECOLAB	4%	N019258 002	Jul 22, 1986
--------	----	-------------	--------------

CIDA-STAT

ECCOLAB	2%	N019258 001	Jul 22, 1986
---------	----	-------------	--------------

DYNA-HEX

BAJAJ MEDICAL LLC	0.75%	N020111 001	Sep 11, 1997
-------------------	-------	-------------	--------------

EXIDINE

+! XTTRIUM	2%	N019422 001	Dec 17, 1985
------------	----	-------------	--------------

	4%	N019125 001	Dec 24, 1984
--	----	-------------	--------------

HIBICLENS

+! MOLNLYCKE HLTH	4%	N017768 001	
-------------------	----	-------------	--

HIBISTAT

+! MOLNLYCKE HLTH	0.5%	N018300 001	
-------------------	------	-------------	--

SPONGE;TOPICAL

BIOSCRUB

GRIFFEN	4%	N019822 001	Mar 31, 1989
---------	----	-------------	--------------

CHLORHEXIDINE GLUCONATE

! BECTON DICKINSON	4%	A072525 001	Oct 24, 1989
--------------------	----	-------------	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-5 (of 20)

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SPONGE;TOPICAL

CHLORAPREP ONE-STEP

+!	BECTON DICKINSON CO	2%;70% (3ML)
+!		2%;70% (10.5ML)
+!		2%;70% (26ML)

N020832	001	Jul 14, 2000
N020832	004	Aug 20, 2003
N020832	006	Nov 21, 2006

CHLORAPREP ONE-STEP FREPP

+!	BECTON DICKINSON CO	2%;70% (1.5ML)
----	---------------------	----------------

N020832	003	Apr 26, 2002
---------	-----	--------------

CHLORAPREP WITH TINT

+!	BECTON DICKINSON CO	2%;70% (26ML)
+!		2%;70% (10.5ML)
+!		2%;70% (3ML)

N020832	002	May 03, 2005
N020832	005	Apr 03, 2006
N020832	007	Oct 10, 2006

SWAB;TOPICAL

CHLORAPREP ONE-STEP SEPP

+!	BECTON DICKINSON CO	2%;70% (0.67ML)
----	---------------------	-----------------

N021555	001	Oct 07, 2002
---------	-----	--------------

CHLORAPREP SINGLE SWABSTICK

+!	BECTON DICKINSON CO	2%;70% (1.75ML)
----	---------------------	-----------------

N021555	002	May 10, 2005
---------	-----	--------------

CHLORAPREP TRIPLE SWABSTICK

+!	BECTON DICKINSON CO	2%;70% (5.25ML)
----	---------------------	-----------------

N021555	003	Jun 10, 2009
---------	-----	--------------

PREVANTICS MAXI SWABSTICK

+!	PROF DSPLS	3.15%;70% (5.1ML)
----	------------	-------------------

N021524	003	Jun 03, 2005
---------	-----	--------------

PREVANTICS SWAB

+!	PROF DSPLS	3.15%;70% (1ML)
----	------------	-----------------

N021524	001	Jun 03, 2005
---------	-----	--------------

PREVANTICS SWABSTICK

+!	PROF DSPLS	3.15%;70% (1.6ML)
----	------------	-------------------

N021524	002	Jun 03, 2005
---------	-----	--------------

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE;ORAL

CHLOR-TRIMETON

+!	BAYER HEALTHCARE LLC	12MG
----	----------------------	------

N007638	002	
---------	-----	--

CHLORPHENIRAMINE MALEATE

AVANTHI INC	12MG
-------------	------

A040829	001	May 13, 2009
---------	-----	--------------

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET;ORAL

ADVIL ALLERGY AND CONGESTION RELIEF

+!	PFIZER	4MG;200MG;10MG
----	--------	----------------

N022113	001	Dec 21, 2011
---------	-----	--------------

ADVIL MULTI-SYMPOTM COLD & FLU

+!	PFIZER	4MG;200MG;10MG
----	--------	----------------

N022113	002	Apr 28, 2017
---------	-----	--------------

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION;ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+!	PFIZER	1MG/5ML;100MG/5ML;15MG/5ML
----	--------	----------------------------

N021587	001	Feb 24, 2004
---------	-----	--------------

TABLET;ORAL

ADVIL ALLERGY SINUS

+!	PFIZER	2MG;200MG;30MG
----	--------	----------------

N021441	001	Dec 19, 2002
---------	-----	--------------

CIMETIDINE

TABLET;ORAL

CIMETIDINE

APOTEX	100MG
--------	-------

A074948	001	Jun 19, 1998
---------	-----	--------------

200MG
-------

A074948	002	Jul 26, 2002
---------	-----	--------------

IVAX SUB TEVA PHARMS	200MG
----------------------	-------

A075345	001	Jun 16, 1999
---------	-----	--------------

L PERRIGO CO	200MG
--------------	-------

A075285	001	Oct 29, 1998
---------	-----	--------------

TAGAMET HB

+!	MEDTECH PRODUCTS	200MG
----	------------------	-------

N020238	002	Aug 21, 1996
---------	-----	--------------

CLEMASTINE FUMARATE

TABLET;ORAL

CLEMASTINE FUMARATE

L PERRIGO CO	1.34MG
--------------	--------

A074512	001	Nov 22, 1995
---------	-----	--------------

SANDOZ	1.34MG
--------	--------

A073458	001	Oct 31, 1993
---------	-----	--------------

TAVIST-1

+!	GLAXOSMITHKLINE CONS	1.34MG
----	----------------------	--------

N020925	001	Aug 21, 1992
---------	-----	--------------

CLOTRIMAZOLE

CREAM;VAGINAL

CLOTRIMAZOLE

!	ACTAVIS MID	1%
---	-------------	----

A074165	001	Jul 16, 1993
---------	-----	--------------

ATLANTIC	
----------	--

TARO	
------	--

	1%
--	----

A072641	001	Dec 04, 1995
---------	-----	--------------

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
OTC DRUG PRODUCT LIST

4-6 (of 20)

CLOTRIMAZOLE

CREAM;VAGINAL			
MYCELEX-7			
BAYER HEALTHCARE	1%		N018230 002 Dec 26, 1991
LLC			
TRIVAGIZOLE 3			
TARO	2%		N021143 001 Apr 12, 2000
CREAM, TABLET;TOPICAL, VAGINAL			
MYCELEX-7 COMBINATION PACK			
BAYER HEALTHCARE	1%,100MG		N020389 002 Jun 23, 1994
LLC			
TABLET;VAGINAL			
MYCELEX-7			
BAYER HEALTHCARE	100MG		N018182 002 Dec 26, 1991
LLC			

CROMOLYN SODIUM

SPRAY, METERED;NASAL			
CROMOLYN SODIUM			
! BAUSCH AND LOMB	5.2MG/SPRAY		A075702 001 Jul 03, 2001
PERRIGO	5.2MG/SPRAY		A075427 001 Dec 12, 2001

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL			
DEXBROMPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE SULFATE			
! AVANTHI INC	6MG;120MG		A078648 001 Feb 27, 2013

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE;ORAL			
GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE			
ACTAVIS LABS FL	30MG;600MG		A091070 001 Aug 31, 2015
	60MG;1.2GM		A091070 002 Aug 31, 2015
AUROBINDO PHARMA LTD	30MG;600MG		A206941 001 Mar 17, 2017
	60MG;1.2GM		A206941 002 Mar 17, 2017
MUCINEX DM			
+ RECKITT BENCKISER	30MG;600MG		N021620 002 Apr 29, 2004
+!	60MG;1.2GM		N021620 001 Apr 29, 2004

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL			
DELSYM			
+! RECKITT BENCKISER	EQ 30MG HBR/5ML		N018658 001 Oct 08, 1982
DEXTROMETHORPHAN POLISTIREX			
AMNEAL PHARMS LLC	EQ 30MG HBR/5ML		A203133 001 Jul 28, 2017
TRIS PHARMA INC	EQ 30MG HBR/5ML		A091135 001 May 25, 2012

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET;ORAL			
ADVIL PM			
+! PFIZER	38MG;200MG		N021394 001 Dec 21, 2005
IBUPROFEN AND DIPHENHYDRAMINE CITRATE			
DR REDDYS LABS LTD	38MG;200MG		A090619 001 Jul 08, 2009
PERRIGO R AND D	38MG;200MG		A079113 001 Dec 22, 2008

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE;ORAL			
ADVIL PM			
+! PFIZER	25MG;EQ 200MG FREE ACID AND POTASSIUM SALT		N021393 001 Dec 21, 2005
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE			
BIONPHARMA INC	25MG;EQ 200MG FREE ACID AND POTASSIUM SALT		A090397 001 Nov 22, 2010
STRIDES PHARMA	25MG;EQ 200MG FREE ACID AND POTASSIUM SALT		A200888 001 Mar 05, 2012

DIPHENHYDRAMINE HYDROCHLORIDE; NAPROXEN SODIUM

TABLET;ORAL			
ALEVE PM			
+! BAYER HLTHCARE	25MG;220MG		N205352 001 Jan 17, 2014

DOCOSANOL

CREAM;TOPICAL			
ABREVA			
+! GLAXOSMITHKLINE	10%		N020941 001 Jul 25, 2000

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-7 (of 20)

DOXYLAMINE SUCCINATE

TABLET;ORAL

DOXYLAMINE SUCCINATE

LNK	25MG	A040564 001 Aug 27, 2004
PERRIGO	25MG	A040167 001 Sep 18, 1996
UNISOM		

+! CHATTEM

25MG

N018066 001

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE MAGNESIUM

AUROBINDO PHARMA LTD	EQ 20MG BASE	A209339 001 Oct 16, 2017
PERRIGO R AND D	EQ 20MG BASE	A207193 001 Aug 18, 2017
NEXIUM 24HR		
+! ASTRAZENECA LP	EQ 20MG BASE	N204655 001 Mar 28, 2014
TABLET, DELAYED RELEASE;ORAL		
NEXIUM 24HR		
+! ASTRAZENECA LP	EQ 20MG BASE	N207920 001 Nov 23, 2015

FAMOTIDINE

TABLET;ORAL

FAMOTIDINE

AUROBINDO PHARMA LTD	10MG	A206531 001 Apr 26, 2016
	20MG	A206531 002 Apr 26, 2016
DR REDDYS LABS LTD	10MG	A075758 001 Aug 17, 2001
	20MG	A077367 001 Sep 25, 2006
IVAX SUB TEVA PHARMS	10MG	A075512 001 Jul 26, 2001
MYLAN	10MG	A075674 001 Dec 21, 2001
PERRIGO	10MG	A075400 001 Mar 18, 2005
PERRIGO R AND D	20MG	A077351 001 Sep 25, 2006
SUN PHARM INDs LTD	10MG	A090283 001 Nov 17, 2009
	20MG	A090283 002 Nov 17, 2009
TEVA	10MG	A075312 001 May 31, 2001
WOCKHARDT	10MG	A077146 001 Mar 07, 2005
	20MG	A090837 001 Aug 04, 2010
PEPCID AC		
+ J AND J CONSUMER INC	10MG	N020325 001 Apr 28, 1995
+!	20MG	N020325 002 Sep 23, 2003
PEPCID AC		
J AND J CONSUMER INC	10MG	N020902 001 Aug 05, 1999
TABLET, CHEWABLE;ORAL		
FAMOTIDINE		
PERRIGO	10MG	A075715 001 Aug 22, 2003
PEPCID AC		
+! J AND J CONSUMER INC	20MG	N020801 002 Dec 17, 2007

FEXOFENADINE HYDROCHLORIDE

SUSPENSION;ORAL

CHILDREN'S ALLEGRA ALLERGY

+! SANOFI AVENTIS US	30MG/5ML	N201373 001 Jan 24, 2011
----------------------	----------	--------------------------

CHILDREN'S ALLEGRA HIVES

+! SANOFI AVENTIS US	30MG/5ML	N201373 002 Jan 24, 2011
----------------------	----------	--------------------------

CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY

ACTAVIS MID ATLANTIC	30MG/5ML	A203330 001 Nov 18, 2014
----------------------	----------	--------------------------

TARO PHARM

30MG/5ML

A208123 001 Nov 09, 2017

CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES

ACTAVIS MID ATLANTIC

30MG/5ML

A203330 002 Nov 18, 2014

TARO PHARM

30MG/5ML

A208123 002 Nov 09, 2017

TABLET;ORAL

ALLEGRA ALLERGY

+ SANOFI AVENTIS US

60MG

N020872 007 Jan 24, 2011

+!

180MG

N020872 010 Jan 24, 2011

ALLEGRA HIVES

+ SANOFI AVENTIS US

60MG

N020872 008 Jan 24, 2011

+!

180MG

N020872 009 Jan 24, 2011

CHILDREN'S ALLEGRA ALLERGY

+ SANOFI AVENTIS US

30MG

N020872 005 Jan 24, 2011

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-8 (of 20)

**FEXOFENADINE HYDROCHLORIDE**

TABLET; ORAL

CHILDREN'S ALLEGRA HIVES					
+ SANOFI AVENTIS US	30MG		N020872	006	Jan 24, 2011
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY					
AUROLIFE PHARMA LLC	30MG		A202039	001	Nov 19, 2014
DR REDDYS LABS LTD	30MG		A076502	004	Apr 12, 2011
HETERO LABS LTD V	30MG		A204097	001	Aug 19, 2016
MYLAN	30MG		A077081	004	Jul 21, 2011
SUN PHARM INDUS	30MG		A091567	002	Feb 06, 2012
TEVA	30MG		A076447	004	Apr 13, 2011
WOCKHARDT LTD	30MG		A079112	002	Feb 08, 2012
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES					
DR REDDYS LABS LTD	30MG		A076502	005	Apr 12, 2011
MYLAN	30MG		A077081	005	Jul 21, 2011
SUN PHARM INDUS	30MG		A091567	001	Feb 06, 2012
TEVA	30MG		A076447	005	Apr 13, 2011
WOCKHARDT LTD	30MG		A079112	001	Feb 08, 2012
FEXOFENADINE HYDROCHLORIDE ALLERGY					
AUROLIFE PHARMA LLC	60MG		A202039	002	Nov 19, 2014
	180MG		A202039	003	Nov 19, 2014
DR REDDYS LABS LTD	60MG		A076502	006	Apr 12, 2011
	180MG		A076502	008	Apr 12, 2011
HETERO LABS LTD V	60MG		A204097	002	Aug 19, 2016
	180MG		A204097	003	Aug 19, 2016
MYLAN	60MG		A077081	006	Jul 21, 2011
	180MG		A077081	008	Jul 21, 2011
SCIEGEN PHARMS INC	60MG		A204507	002	Sep 16, 2015
	180MG		A204507	003	Sep 16, 2015
SUN PHARM INDUS	60MG		A091567	004	Feb 06, 2012
	180MG		A091567	006	Feb 06, 2012
TEVA	60MG		A076447	006	Apr 13, 2011
	180MG		A076447	008	Apr 13, 2011
WOCKHARDT LTD	60MG		A079112	004	Feb 08, 2012
	180MG		A079112	006	Feb 08, 2012
FEXOFENADINE HYDROCHLORIDE HIVES					
DR REDDYS LABS LTD	60MG		A076502	007	Apr 12, 2011
	180MG		A076502	009	Apr 12, 2011
MYLAN	60MG		A077081	007	Jul 21, 2011
	180MG		A077081	009	Jul 21, 2011
SCIEGEN PHARMS INC	60MG		A204507	004	Sep 16, 2015
	180MG		A204507	005	Sep 16, 2015
SUN PHARM INDUS	60MG		A091567	003	Feb 06, 2012
	180MG		A091567	005	Feb 06, 2012
TEVA	60MG		A076447	007	Apr 13, 2011
	180MG		A076447	009	Apr 13, 2011
WOCKHARDT LTD	60MG		A079112	003	Feb 08, 2012
	180MG		A079112	005	Feb 08, 2012
TABLET, ORALLY DISINTEGRATING; ORAL					
CHILDREN'S ALLEGRA ALLERGY					
+! SANOFI AVENTIS US	30MG		N021909	002	Jan 24, 2011
CHILDREN'S ALLEGRA HIVES					
+! SANOFI AVENTIS US	30MG		N021909	003	Jan 24, 2011
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY					
DR REDDYS LABS LTD	30MG		A202978	001	Jan 18, 2013
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES					
DR REDDYS LABS LTD	30MG		A202978	002	Jan 18, 2013

**FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHENDRINE HYDROCHLORIDE**

TABLET, EXTENDED RELEASE; ORAL					
ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION					
+! SANOFI AVENTIS US	60MG;120MG		N020786	002	Jan 24, 2011
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION					
+! SANOFI AVENTIS US	180MG;240MG		N021704	002	Jan 24, 2011
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHENDRINE HYDROCHLORIDE					
AUROBINDO PHARMA LTD	60MG;120MG		A209116	001	Oct 30, 2017
DR REDDYS LABS LTD	60MG;120MG		A076667	001	Nov 18, 2014
	180MG;240MG		A079043	002	Jun 22, 2011
SUN PHARMA GLOBAL	60MG;120MG		A090818	001	Jan 29, 2015

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-9 (of 20)

**FLUTICASONE FUROATE**

SPRAY, METERED;NASAL		
FLONASE SENSI-MIST ALLERGY RELIEF		
+! GLAXOSMITHKLINE	0.0275MG/SPRAY	
CONS		N022051 002 Aug 02, 2016

**FLUTICASONE PROPIONATE**

SPRAY, METERED;NASAL		
FLONASE ALLERGY RELIEF		
+! GLAXOSMITHKLINE	0.05MG/SPRAY	
CONS		N205434 001 Jul 23, 2014
FLUTICASONE PROPIONATE		
APOTEX INC	0.05MG/SPRAY	A208150 001 Feb 29, 2016
WEST-WARD PHARMS INT	0.05MG/SPRAY	A207957 001 May 26, 2016

**GUAIFENESIN**

TABLET, EXTENDED RELEASE;ORAL		
GUAIFENESIN		
ACTAVIS LABS FL	1.2GM	A091009 002 Sep 03, 2015
GUARDIAN PHARMS	600MG	A209215 001 Sep 06, 2017
	1.2GM	A209215 002 Sep 06, 2017
PERRIGO R AND D	600MG	A078912 001 Nov 23, 2011
MUCINEX		
+ RECKITT BENCKISER	600MG	N021282 001 Jul 12, 2002
+!	1.2GM	N021282 002 Dec 18, 2002

**GUAIFENESIN: PSEUDOEPHEDRINE HYDROCHLORIDE**

TABLET, EXTENDED RELEASE;ORAL		
GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE		
ACTAVIS LABS FL	600MG;60MG	A091071 001 May 27, 2015
	1.2GM;120MG	A091071 002 May 27, 2015
DR REDDYS LABS LTD	600MG;60MG	A208369 001 Dec 29, 2017
	1.2GM;120MG	A208369 002 Dec 29, 2017
MUCINEX D		
+ RECKITT BENCKISER	600MG;60MG	N021585 001 Jun 22, 2004
+!	1.2GM;120MG	N021585 002 Jun 22, 2004

**IBUPROFEN**

CAPSULE;ORAL		
ADVIL LIQUI-GELS		
+! PFIZER	EQ 200MG FREE ACID AND POTASSIUM SALT	N020402 001 Apr 20, 1995
ADVIL MIGRAINE LIQUI-GELS		
+! PFIZER	EQ 200MG FREE ACID AND POTASSIUM SALT	N020402 002 Mar 16, 2000
IBUPROFEN		
AMNEAL PHARMS	EQ 200MG FREE ACID AND POTASSIUM SALT	A202300 001 Dec 23, 2011
ASCENT PHARMS INC	EQ 200MG FREE ACID AND POTASSIUM SALT	A206999 001 Dec 21, 2017
BIONPHARMA INC	EQ 200MG FREE ACID AND POTASSIUM SALT	A078682 001 Mar 24, 2009
HUMANWELL PURACAP	EQ 200MG FREE ACID AND POTASSIUM SALT	A206568 001 Jun 21, 2016
MARKSANS PHARMA	EQ 200MG FREE ACID AND POTASSIUM SALT	A079205 001 Jun 26, 2009
P AND L DEV LLC	EQ 200MG FREE ACID AND POTASSIUM SALT	A077338 001 Jul 10, 2009
SOFGEN PHARMS	EQ 200MG FREE ACID AND POTASSIUM SALT	A203599 001 Sep 07, 2016
MIDOL LIQUID GELS		
+! BIONPHARMA INC	200MG	N021472 001 Oct 18, 2002
SUSPENSION;ORAL		
CHILDREN'S ADVIL		
PFIZER	100MG/5ML	N020589 001 Jun 27, 1996
CHILDREN'S ADVIL-FLAVORED		
PFIZER	100MG/5ML	N020589 002 Nov 07, 1997
CHILDREN'S ELIXSURE		
MOBERG PHARMA NORTH	100MG/5ML	N021604 001 Jan 07, 2004
CHILDREN'S IBUPROFEN		
PERRIGO	100MG/5ML	A074937 001 Dec 22, 1998
CHILDREN'S MOTRIN		
+! J AND J CONSUMER INC	100MG/5ML	N020516 001 Jun 16, 1995
IBUPROFEN		
ACTAVIS MID	100MG/5ML	A074916 001 Apr 30, 1999
ATLANTIC		
ARISE PHARMS	100MG/5ML	A200457 001 Aug 18, 2011
TARO	100MG/5ML	A209207 001 Jun 27, 2017
SUSPENSION/DROPS;ORAL		
CHILDREN'S MOTRIN		
+! J AND J CONSUMER INC	40MG/ML	N020603 001 Jun 10, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-10 (of 20)

IBUPROFEN

SUSPENSION/DROPS;ORAL

IBUPROFEN

L PERRIGO CO	40MG/ML	A075217 001 Dec 16, 1998
TRIS PHARMA INC	40MG/ML	A079058 001 Aug 31, 2009

PEDIATRIC ADVIL

+! PFIZER	100MG/2.5ML	N020812 001 Jan 30, 1998
-----------	-------------	--------------------------

TABLET;ORAL

ADVIL

PFIZER	200MG	N018989 001 May 18, 1984
--------	-------	--------------------------

IBU-TAB 200

ALRA	200MG	A071057 001 Aug 11, 1988
------	-------	--------------------------

IBUPROFEN

AIPING PHARM INC	200MG	A207095 001 May 05, 2017
------------------	-------	--------------------------

AMNEAL PHARMS	200MG	A079233 001 Mar 18, 2014
---------------	-------	--------------------------

AMNEAL PHARMS NY	200MG	A071333 001 Feb 17, 1987
------------------	-------	--------------------------

	200MG	A072199 001 May 23, 1988
--	-------	--------------------------

AUROBINDO PHARMA LTD	200MG	A208865 001 Nov 08, 2017
----------------------	-------	--------------------------

AVEMA PHARMA	200MG	A076460 001 Nov 26, 2003
--------------	-------	--------------------------

CONTRACT PHARMACAL	200MG	A071732 001 Sep 10, 1987
--------------------	-------	--------------------------

	200MG	A072299 001 Jul 01, 1988
--	-------	--------------------------

DR REDDYS LA	200MG	A075661 001 Dec 12, 2001
--------------	-------	--------------------------

DR REDDYS LABS INC	100MG	A076117 001 Nov 20, 2001
--------------------	-------	--------------------------

GRANULES INDIA	200MG	A079174 001 Dec 10, 2010
----------------	-------	--------------------------

GRANULES INDIA LTD	200MG	A202312 001 Oct 07, 2016
--------------------	-------	--------------------------

LNK	100MG	A076741 001 Jun 17, 2004
-----	-------	--------------------------

	200MG	A075010 001 Mar 01, 1999
--	-------	--------------------------

	200MG	A075139 001 Mar 01, 1999
--	-------	--------------------------

MARKSANS PHARMA	200MG	A091237 001 Feb 08, 2011
-----------------	-------	--------------------------

	200MG	A091239 001 Feb 01, 2011
--	-------	--------------------------

MCNEIL	200MG	A073019 001 Mar 30, 1994
--------	-------	--------------------------

MERRO PHARM	200MG	A070985 001 Oct 02, 1987
-------------	-------	--------------------------

OHM	200MG	A071163 001 Jul 15, 1986
-----	-------	--------------------------

PAR PHARM	200MG	A070481 001 Sep 24, 1986
-----------	-------	--------------------------

PERRIGO	200MG	A072096 001 Dec 08, 1987
---------	-------	--------------------------

	200MG	A075995 001 Mar 14, 2002
--	-------	--------------------------

PERRIGO R AND D	200MG	A077349 001 Jun 21, 2005
-----------------	-------	--------------------------

STRIDES PHARMA	200MG	A079129 001 Mar 28, 2011
----------------	-------	--------------------------

	200MG	A091355 001 Apr 04, 2011
--	-------	--------------------------

	200MG	A207052 001 May 30, 2017
--	-------	--------------------------

VINTAGE PHARMS	200MG	A071229 001 Apr 01, 1987
----------------	-------	--------------------------

	200MG	A071639 001 Feb 02, 1988
--	-------	--------------------------

IBUPROHMG

OHM LABS	200MG	A071214 001 Dec 01, 1986
----------	-------	--------------------------

JUNIOR STRENGTH ADVIL

PFIZER	100MG	N020267 002 Dec 13, 1996
--------	-------	--------------------------

JUNIOR STRENGTH IBUPROFEN

L PERRIGO CO	100MG	A075367 001 Apr 22, 1999
--------------	-------	--------------------------

JUNIOR STRENGTH MOTRIN

J AND J CONSUMER INC	100MG	N020602 001 Jun 10, 1996
----------------------	-------	--------------------------

MOTRIN IB

+! J AND J CONSUMER INC	200MG	N019012 003 Dec 17, 1990
-------------------------	-------	--------------------------

PROFEN

CONTRACT PHARMACAL	200MG	A071265 001 Oct 15, 1986
--------------------	-------	--------------------------

TAB-PROFEN

PERRIGO	200MG	A072095 001 Dec 08, 1987
---------	-------	--------------------------

TABLET, CHEWABLE;ORAL

CHILDREN'S ADVIL

PFIZER	50MG	N020944 001 Dec 18, 1998
--------	------	--------------------------

CHILDREN'S MOTRIN

+! J AND J CONSUMER INC	50MG	N020601 001 Nov 15, 1996
-------------------------	------	--------------------------

IBUPROFEN

PERRIGO	50MG	A076359 001 Jan 16, 2004
---------	------	--------------------------

	100MG	A076359 002 Jan 16, 2004
--	-------	--------------------------

JUNIOR STRENGTH ADVIL

PFIZER	100MG	N020944 002 Dec 18, 1998
--------	-------	--------------------------

JUNIOR STRENGTH MOTRIN

+! J AND J CONSUMER INC	100MG	N020601 003 Nov 15, 1996
-------------------------	-------	--------------------------

	100MG	
--	-------	--

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-11 (of 20)

IBUPROFEN SODIUM

TABLET;ORAL

ADVIL

+!	Pfizer Cons Hlthcare	EQ 200MG BASE
	IBUPROFEN SODIUM	
	Perrigo R AND D	EQ 200MG BASE

N201803 001	Jun 12, 2012
A206581 001	Aug 03, 2015

IBUPROFEN; PHENYLEPHRINE HYDROCHLORIDE

TABLET;ORAL

ADVIL CONGESTION RELIEF

+!	Pfizer	200MG;10MG
	IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE	
	Perrigo R AND D	200MG;10MG

N022565 001	May 27, 2010
A203200 001	Jul 03, 2014

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE;ORAL

ADVIL COLD AND SINUS

+!	Pfizer	EQ 200MG FREE ACID AND POTASSIUM SALT;30MG
	IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE	
	Aurobindo Pharma Ltd	EQ 200MG FREE ACID AND POTASSIUM SALT;30MG

N021374 001	May 30, 2002
A209235 001	Dec 01, 2017

SUSPENSION;ORAL

CHILDREN'S ADVIL COLD

Pfizer	100MG/5ML;15MG/5ML
--------	--------------------

N021373 001	Apr 18, 2002
-------------	--------------

CHILDREN'S MOTRIN COLD

+! J AND J CONSUMER INC	100MG/5ML;15MG/5ML
----------------------------	--------------------

N021128 001	Aug 01, 2000
-------------	--------------

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

Perrigo	100MG/5ML;15MG/5ML
---------	--------------------

A076478 001	Nov 05, 2003
-------------	--------------

TABLET;ORAL

ADVIL COLD AND SINUS

+! Pfizer	200MG;30MG
	IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE
DR REDDYS LABS LTD	200MG;30MG
	IBUPROHM COLD AND SINUS
OHM LABS	200MG;30MG
	SINE-AID IB
J AND J CONSUMER INC	200MG;30MG

N019771 001	Sep 19, 1989
A077628 001	Aug 14, 2006
A074567 001	Apr 17, 2001
N019899 001	Dec 31, 1992

INSULIN RECOMBINANT HUMAN

INJECTABLE;INJECTION

HUMULIN R PEN

+! Lilly	100 UNITS/ML
NOVOLIN R	
+! NOVO NORDISK INC	100 UNITS/ML

N018780 005	Aug 06, 1998
N019938 001	Jun 25, 1991

INSULIN RECOMBINANT HUMAN

INJECTABLE;INJECTION

HUMULIN R

+! Lilly	100 UNITS/ML
----------	--------------

N018780 001	Oct 28, 1982
-------------	--------------

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE;INJECTION

HUMULIN 70/30

+! Lilly	30 UNITS/ML;70 UNITS/ML
HUMULIN 70/30 PEN	
+! Lilly	30 UNITS/ML;70 UNITS/ML
NOVOLIN 70/30	
+! NOVO NORDISK INC	30 UNITS/ML;70 UNITS/ML

N019717 001	Apr 25, 1989
N019717 002	Aug 06, 1998
N019991 001	Jun 25, 1991

INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE;INJECTION

HUMULIN N

+! Lilly	100 UNITS/ML
NOVOLIN N	
+! NOVO NORDISK INC	100 UNITS/ML

N018781 001	Oct 28, 1982
N019959 001	Jul 01, 1991

IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE;TOPICAL

DURAPREP

+! 3M	EQ 0.7% IODINE;74% (6ML)
+!	EQ 0.7% IODINE;74% (26ML)

N021586 001	Sep 29, 2006
N021586 002	Sep 29, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-12 (of 20)

**KETOCONAZOLE**

SHAMPOO;TOPICAL	
NIZORAL A-D	
+! JOHNSON AND JOHNSON 1%	N020310 001 Oct 10, 1997

**KETOTIFEN FUMARATE**

SOLUTION/DROPS;OPHTHALMIC		
ALAWAY		
+! BAUSCH AND LOMB	EQ 0.025% BASE	N021996 001 Dec 01, 2006
+	EQ 0.035% BASE	N021996 002 Feb 11, 2015
KETOTIFEN FUMARATE		
AKORN	EQ 0.025% BASE	A077958 001 Jul 26, 2007
! ALCON PHARMS LTD	EQ 0.025% BASE	A077200 001 Sep 02, 2008

**LANSOPRAZOLE**

CAPSULE, DELAYED REL PELLETS;ORAL		
LANSOPRAZOLE		
DR REDDYS LABS LTD	15MG	A202194 001 May 18, 2012
KREMERS URBAN	15MG	A207157 001 Sep 29, 2017
PHARMS		
MYLAN PHARMS INC	15MG	A203187 001 Jun 01, 2016
NATCO PHARMA LTD	15MG	A203306 001 Jan 13, 2016
PERRIGO R AND D	15MG	A202319 001 May 18, 2012
WOCKHARDT LTD	15MG	A202727 001 May 18, 2012
PREVACID 24 HR		
+! GLAXOSMITHKLINE CONS	15MG	N022327 001 May 18, 2009
TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL		
LANSOPRAZOLE		
DEXCEL PHARMA	15MG	N208025 001 Jun 07, 2016

**LEVOCECIRIZINE DIHYDROCHLORIDE**

SOLUTION;ORAL		
XYZAL ALLERGY 24HR		
+ SANOFI-AVENTIS US	2.5MG/5ML (0.5MG/ML)	N209090 001 Jan 31, 2017
TABLET;ORAL		
XYZAL ALLERGY 24HR		
+ SANOFI-AVENTIS US	5MG	N209089 001 Jan 31, 2017

**LEVONORGESTREL**

TABLET;ORAL		
ATHENTIA NEXT		
AUROBINDO PHARMA LTD	1.5MG	A206867 001 Dec 08, 2015
FALLBACK SOLO		
LUPIN LTD	1.5MG	A201446 001 Jun 19, 2014
HER STYLE		
NOVAST LABS LTD	1.5MG	A207976 001 Mar 11, 2016
LEVONORGESTREL		
FDN CONSUMER	1.5MG	A200670 001 Jul 12, 2012
GLENMARK PHARMS LTD	1.5MG	A207044 001 Mar 25, 2016
LOTUS PHARM CO LTD	0.75MG	A202684 001 Sep 02, 2016
MYLAN LABS LTD	0.75MG	A202740 001 Sep 02, 2016
	1.5MG	A202739 001 Oct 31, 2014
NOVEL LABS INC	1.5MG	A202508 001 Feb 22, 2013
OC PHARMA	1.5MG	A202380 001 May 29, 2015
! PERRIGO R AND D	0.75MG	A090740 001 Dec 30, 2010
	1.5MG	A202334 001 Aug 20, 2014
RECKITT BENCKISER	1.5MG	A202246 001 Jun 05, 2015
OPCICON ONE-STEP		
SUN PHARM INDs LTD	1.5MG	A202635 001 Sep 11, 2014
PLAN B ONE-STEP		
+! FDN CONSUMER	1.5MG	N021998 001 Jul 10, 2009

**LOPERAMIDE HYDROCHLORIDE**

CAPSULE;ORAL		
LOPERAMIDE HYDROCHLORIDE		
+ BIONPHARMA INC	1MG	N021855 001 Aug 04, 2005
+!	2MG	N021855 002 Aug 04, 2005
SOLUTION;ORAL		
IMODIUM A-D		
+! J AND J CONSUMER INC	1MG/5ML	N019487 001 Mar 01, 1988
LOPERAMIDE HYDROCHLORIDE		
ALLIED PHARMA INC	1MG/5ML	A073079 001 Apr 30, 1992
HI TECH PHARMA	1MG/5ML	A074352 001 Nov 17, 1995

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-13 (of 20)

LOPERAMIDE HYDROCHLORIDE

SOLUTION;ORAL			
LOPERAMIDE HYDROCHLORIDE			
PERRIGO	1MG/5ML	A073243	001 Jan 21, 1992
WOCKHARDT BIO AG	1MG/5ML	A074730	001 Aug 28, 1997
SUSPENSION;ORAL			
IMODIUM A-D			
+! J AND J CONSUMER INC	1MG/7.5ML	N019487	002 Jul 08, 2004
LOPERAMIDE HYDROCHLORIDE			
PERRIGO R AND D	1MG/7.5ML	A091292	001 May 20, 2011
TABLET;ORAL			
IMODIUM A-D			
+! J AND J CONSUMER INC	2MG	N019860	001 Nov 22, 1989
LOPERAMIDE HYDROCHLORIDE			
AUROBINDO PHARMA LTD	2MG	A206548	001 Dec 15, 2015
L PERRIGO CO	2MG	A075232	001 Jan 06, 2000
LNK	2MG	A076497	001 Jun 10, 2003
OHM LABS	2MG	A074091	001 Dec 10, 1992

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET;ORAL			
IMODIUM MULTI-SYMPTOM RELIEF			
+! J AND J CONSUMER INC	2MG;125MG	N021140	001 Nov 30, 2000
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE			
SUN PHARM INDs LTD	2MG;125MG	A077500	001 Sep 06, 2006
TABLET, CHEWABLE;ORAL			
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE			
! PERRIGO	2MG;125MG	A076029	001 Aug 30, 2002

LORATADINE

CAPSULE;ORAL			
CLARITIN			
+! BAYER HEALTHCARE LLC	10MG	N021952	001 Jun 16, 2008
LORATADINE			
MARKSANS PHARMA	10MG	A206214	001 Sep 23, 2016
SUSPENSION;ORAL			
LORATADINE			
+! TARO	1MG/ML	N021734	001 Oct 04, 2005
SYRUP;ORAL			
CLARITIN			
+! BAYER HEALTHCARE LLC	1MG/ML	N020641	002 Nov 27, 2002
LORATADINE			
PERRIGO	1MG/ML	A075728	001 Aug 20, 2004
SILARX	1MG/ML	A077421	001 Jun 29, 2006
TARO	1MG/ML	A076805	001 Aug 20, 2004
TARO PHARM	1MG/ML	A201865	001 Jul 31, 2015
TEVA	1MG/ML	A075505	001 Nov 07, 2003
WOCKHARDT BIO AG	1MG/ML	A075815	001 Aug 20, 2004
TABLET;ORAL			
CLARITIN			
+! BAYER HEALTHCARE LLC	10MG	N019658	002 Nov 27, 2002
CLARITIN HIVES RELIEF			
+! BAYER HEALTHCARE LLC	10MG	N019658	003 Nov 19, 2003
LORATADINE			
APOTEX INC	10MG	A076471	001 Feb 14, 2006
MYLAN	10MG	A075790	001 Nov 07, 2008
	10MG	A076154	001 Aug 20, 2003
	10MG	A078447	001 Aug 12, 2011
PERRIGO	10MG	A076301	001 Jun 25, 2004
SANDOZ	10MG	A075209	001 Jan 21, 2003
SUN PHARM INDs LTD	10MG	A076134	001 Aug 18, 2003
TABLET, CHEWABLE;ORAL			
CHILDREN'S CLARITIN			
+! BAYER HEALTHCARE LLC	5MG	N021891	001 Aug 23, 2006

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-14 (of 20)

LORATADINE

TABLET, ORALLY DISINTEGRATING;ORAL			
ALAVERT			
Pfizer	10MG	N021375	001 Dec 19, 2002
CLARITIN HIVES RELIEF REDITAB			
+! BAYER HEALTHCARE LLC	10MG	N020704	003 Nov 19, 2003
CLARITIN REDITABS			
+! BAYER HEALTHCARE LLC	5MG	N021993	001 Dec 12, 2006
+!	10MG	N020704	002 Nov 27, 2002
LORATADINE			
ACTAVIS LABS FL INC	10MG	A075990	001 Nov 03, 2003
PERRIGO PHARMA INTL	10MG	A076011	001 Sep 29, 2003
Pfizer	10MG	A075822	001 Feb 10, 2003
LORATADINE REDIDOSE			
SUN PHARM INDs LTD	10MG	A077153	001 Apr 11, 2007

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL			
CLARITIN-D			
+! BAYER HEALTHCARE LLC	5MG;120MG	N019670	002 Nov 27, 2002
CLARITIN-D 24 HOUR			
+! BAYER HEALTHCARE LLC	10MG;240MG	N020470	002 Nov 27, 2002
LORATADINE AND PSEUDOEPHEDRINE SULFATE			
ACTAVIS LABS FL INC	10MG;240MG	A075706	001 Feb 21, 2003
PERRIGO PHARMA INTL	5MG;120MG	A076050	001 Jan 30, 2003
	10MG;240MG	A075989	001 Mar 04, 2004
SUN PHARM INDs LTD	10MG;240MG	A076557	001 Sep 22, 2004

MENTHOL; METHYL SALICYLATE

PATCH;TOPICAL			
SALONPAS			
+! HISAMITSU PHARM CO	3%;10%	N022029	001 Feb 20, 2008
+	3%;10%	N022029	002 Nov 05, 2012

MICONAZOLE NITRATE

CREAM;TOPICAL, VAGINAL			
MICONAZOLE 3 COMBINATION PACK			
PERRIGO	2%,4%	A076357	001 Mar 30, 2004
MONISTAT 3 COMBINATION PACK			
+ MEDTECH PRODUCTS	2%,4%	N021261	003 Jun 17, 2003
MONISTAT 3 COMBINATION PACK (PREFILLED)			
+! MEDTECH PRODUCTS	2%,4%	N021261	001 Feb 02, 2001
CREAM;VAGINAL			
MICONAZOLE 3			
TARO	4%	A076773	001 Mar 02, 2005
MICONAZOLE 7			
ACTAVIS MID ATLANTIC	2%	A074164	001 Mar 29, 1996
MICONAZOLE NITRATE			
G AND W LABS INC	2%	A074366	001 Feb 22, 1996
PERRIGO	2%	A074760	001 May 15, 1997
PERRIGO R AND D	4%	A091366	001 Jan 15, 2010
TARO	2%	A074444	001 Jan 13, 1997
MONISTAT 3			
+! MEDTECH PRODUCTS	4%	N020827	001 Mar 30, 1998
MONISTAT 7			
+! MEDTECH PRODUCTS	2%	N017450	002 Feb 15, 1991
CREAM, SUPPOSITORY;TOPICAL, VAGINAL			
M-ZOLE 3 COMBINATION PACK			
ACTAVIS MID ATLANTIC	2%,200MG	A074926	001 Apr 16, 1999
MICONAZOLE 7 COMBINATION PACK			
G AND W LABS	2%,100MG	A076585	001 Mar 26, 2004
MICONAZOLE NITRATE			
PERRIGO R AND D	2%,1.2GM	A079114	001 Jun 02, 2010
MICONAZOLE NITRATE COMBINATION PACK			
PERRIGO	2%,200MG	A075329	001 Apr 20, 1999
MONISTAT 1 COMBINATION PACK			
+! MEDTECH PRODUCTS	2%,1.2GM	N021308	001 Jun 29, 2001
MONISTAT 3 COMBINATION PACK			
+! MEDTECH PRODUCTS	2%,200MG	N020670	002 Apr 16, 1996

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-15 (of 20)

MICONAZOLE NITRATE

CREAM, SUPPOSITORY;TOPICAL, VAGINAL		
MONISTAT 7 COMBINATION PACK		
+! MEDTECH PRODUCTS	2%,100MG	N020288 002 Apr 26, 1993
SUPPOSITORY;VAGINAL		
MICONAZOLE NITRATE		
ACTAVIS PHARMA	100MG	A073507 001 Nov 19, 1993
G AND W LABS	100MG	A074414 001 Apr 30, 1997
! PERRIGO	100MG	A074395 001 Mar 20, 1997
MONISTAT 7		
+! MEDTECH PRODUCTS	100MG	N018520 002 Feb 15, 1991

MINOXIDIL

AEROSOL, FOAM;TOPICAL		
MEN'S ROGAINE		
+! JOHNSON AND JOHNSON	5%	N021812 001 Jan 20, 2006
MINOXIDIL		
PERRIGO ISRAEL	5%	A091344 001 Apr 28, 2011
MINOXIDIL (FOR MEN)		
WATSON LABS INC	5%	A208092 001 Feb 17, 2017
MINOXIDIL (FOR WOMEN)		
WATSON LABS INC	5%	A208092 002 Jul 27, 2017
WOMEN'S ROGAINE		
+! JOHNSON AND JOHNSON	5%	N021812 002 Feb 28, 2014
SOLUTION;TOPICAL		
MINOXIDIL (FOR MEN)		
ACTAVIS MID	2%	A074588 001 Apr 05, 1996
ATLANTIC		
HI TECH PHARMA	2%	A074731 001 Dec 24, 1996
L PERRIGO CO	2%	A075357 001 Jul 30, 1999
WOCKHARDT BIO AG	2%	A074767 001 Feb 28, 1997
MINOXIDIL (FOR WOMEN)		
HI TECH PHARMA	2%	A074731 002 May 11, 2005
L PERRIGO CO	2%	A075357 002 Jul 30, 1999
MINOXIDIL EXTRA STRENGTH (FOR MEN)		
ACTAVIS MID	5%	A075518 001 Nov 17, 2000
ATLANTIC		
AVACOR PRODS	5%	A075619 001 Nov 17, 2000
PERRIGO	5%	A075598 001 Jun 13, 2001
PERRIGO NEW YORK	5%	A075737 001 Mar 15, 2002
WOCKHARDT BIO AG	5%	A075438 001 Feb 27, 2003
ROGAINE (FOR MEN)		
+! JOHNSON AND JOHNSON	2%	N019501 002 Feb 09, 1996
ROGAINE (FOR WOMEN)		
+! JOHNSON AND JOHNSON	2%	N019501 003 Feb 09, 1996
ROGAINE EXTRA STRENGTH (FOR MEN)		
+! JOHNSON AND JOHNSON	5%	N020834 001 Nov 14, 1997
ATHEROXIDIL		
EI INC	2%	A078176 001 Nov 09, 2007
	5%	A076239 001 Aug 24, 2004

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC		
NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE		
AKORN INC	0.025%;0.3%	A202795 001 Jan 24, 2013
ALTAIRE PHARMS INC	0.02675%;0.315%	A078208 001 Sep 27, 2010
NAPHCON-A		
+! ALCON	0.025%;0.3%	N020226 001 Jun 08, 1994
OPCON-A		
+! BAUSCH AND LOMB	0.02675%;0.315%	N020065 001 Jun 08, 1994
VISINE-A		
+! JOHNSON AND JOHNSON	0.025%;0.3%	N020485 001 Jan 31, 1996

NAPROXEN SODIUM

CAPSULE;ORAL		
NAPROXEN SODIUM		
+! BIONPHARMA INC	EQ 200MG BASE	N021920 001 Feb 17, 2006
TABLET;ORAL		
ALEVE		
+! BAYER	EQ 200MG BASE	N020204 002 Jan 11, 1994
NAPROXEN SODIUM		
AMNEAL PHARMS NY	EQ 200MG BASE	A079096 001 Dec 16, 2008
AUROBINDO PHARMA LTD	EQ 200MG BASE	A205497 001 Mar 18, 2016

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-16 (of 20)

NAPROXEN SODIUM

TABLET;ORAL

NAPROXEN SODIUM

CONTRACT PHARMACAL	EQ 200MG BASE	A074635 001	Jan 13, 1997
DR REDDYS LABS INC	EQ 200MG BASE	A075168 001	Jul 28, 1998
GRANULES INDIA	EQ 200MG BASE	A091353 001	Sep 20, 2011
LNK INTL INC	EQ 200MG BASE	A204872 001	Jan 23, 2017
MARKSANS PHARMA	EQ 200MG BASE	A090545 001	Mar 16, 2011
PERRIGO	EQ 200MG BASE	A074661 001	Jan 13, 1997
SUN PHARM INDNS LTD	EQ 200MG BASE	A091183 001	May 20, 2011

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

ALEVE-D SINUS & COLD

+! BAYER	200MG;120MG	N021076 001	Nov 29, 1999
NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE			
DR REDDYS LABS INC	EQ 220MG BASE;120MG	A077381 001	Sep 27, 2006

PERRIGO EQ 200MG BASE;120MG

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

HABITROL

+	DR REDDYS LABS SA	7MG/24HR	N020076 004	Nov 12, 1999
+		14MG/24HR	N020076 005	Nov 12, 1999
+!		21MG/24HR	N020076 006	Nov 12, 1999

NICODERM CQ

+	SANOFI AVENTIS US	7MG/24HR	N020165 006	Aug 02, 1996
+		14MG/24HR	N020165 005	Aug 02, 1996
+!		21MG/24HR	N020165 004	Aug 02, 1996

NICOTINE

AVEVA	7MG/24HR	A074612 002	Jul 28, 2003
	14MG/24HR	A074612 003	Oct 20, 1997
	21MG/24HR	A074612 001	Oct 20, 1997

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICORETTE

+	GLAXOSMITHKLINE	EQ 2MG BASE	N018612 002	Feb 09, 1996
+		EQ 2MG BASE	N018612 004	Sep 25, 2000
+!		EQ 4MG BASE	N020066 002	Feb 09, 1996
+		EQ 4MG BASE	N020066 004	Sep 25, 2000

NICORETTE (MINT)

+	GLAXOSMITHKLINE	EQ 2MG BASE	N018612 003	Dec 23, 1998
+		EQ 4MG BASE	N020066 003	Dec 23, 1998

NICOTINE POLACRILEX

ACTAVIS LABS NY INC	EQ 2MG BASE	A074507 001	Mar 15, 1999
	EQ 2MG BASE	A076569 001	Jul 29, 2004
	EQ 2MG BASE	A078699 001	Dec 29, 2008
	EQ 2MG BASE	A079216 001	Jul 08, 2009
	EQ 2MG BASE	A204794 001	May 10, 2016
	EQ 4MG BASE	A074707 001	Mar 19, 1999
	EQ 4MG BASE	A076568 002	Jul 29, 2004
	EQ 4MG BASE	A078697 001	Dec 29, 2008
	EQ 4MG BASE	A079038 001	Jul 08, 2009
	EQ 4MG BASE	A079219 001	Jul 08, 2009
	EQ 4MG BASE	A204833 001	Feb 26, 2016

L PERRIGO CO

	EQ 2MG BASE	A076775 001	Sep 16, 2004
	EQ 2MG BASE	A076776 001	Sep 16, 2004
	EQ 2MG BASE	A076777 001	Sep 16, 2004
	EQ 4MG BASE	A076778 001	Sep 16, 2004
	EQ 4MG BASE	A076779 001	Sep 16, 2004

	EQ 4MG BASE	A076789 001	Sep 16, 2004
	EQ 4MG BASE	A078325 001	Oct 30, 2006
	EQ 4MG BASE	A078547 001	May 24, 2007
	EQ 4MG BASE	A078967 001	Apr 23, 2008
	EQ 4MG BASE	A091349 001	Jul 20, 2011

	EQ 2MG BASE	A206394 001	Dec 15, 2016
	EQ 4MG BASE	A078326 001	Oct 30, 2006
	EQ 4MG BASE	A078546 001	May 24, 2007
	EQ 4MG BASE	A078968 001	Apr 23, 2008
	EQ 4MG BASE	A091354 001	Jul 20, 2011

	EQ 4MG BASE	A206393 001	Dec 15, 2016
	EQ 2MG BASE	A079044 001	Jul 08, 2009




38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-17 (of 20)

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL			
THRIVE			
NOVARTIS	EQ 2MG BASE	A077658 001	Jun 19, 2007
	EQ 4MG BASE	A077656 001	Jun 19, 2007
TROCHE/LOZENGE;ORAL			
COMMIT			
+ GLAXOSMITHKLINE	EQ 2MG BASE	N021330 001	Oct 31, 2002
CONS			
+!	EQ 4MG BASE	N021330 002	Oct 31, 2002
NICORETTE			
+ GLAXOSMITHKLINE	EQ 2MG BASE	N022360 001	May 18, 2009
CONS			
+!	EQ 4MG BASE	N022360 002	May 18, 2009
NICOTINE POLACRILEX			
PERRIGO R AND D	EQ 2MG BASE	A077007 001	Jan 31, 2006
	EQ 2MG BASE	A090711 001	Jul 10, 2009
	EQ 2MG BASE	A090821 001	Jul 10, 2009
	EQ 2MG BASE	A203690 001	Oct 09, 2012
	EQ 4MG BASE	A077007 002	Jan 31, 2006
	EQ 4MG BASE	A090711 002	Jul 10, 2009
	EQ 4MG BASE	A090821 002	Jul 10, 2009
	EQ 4MG BASE	A203690 002	Oct 09, 2012

NIZATIDINE

TABLET;ORAL			
AXID AR			
+! PFIZER	75MG	N020555 001	May 09, 1996

NONOXYNOL-9

SPONGE;VAGINAL			
TODAY			
+! MAYER LABS INC	1GM	N018683 001	Apr 01, 1983

OMEPRAZOLE

TABLET, DELAYED RELEASE;ORAL			
OMEPRAZOLE			
+! DEXCEL PHARMA	20MG	N022032 001	Dec 04, 2007
TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL			
OMEPRAZOLE			
+ DEXCEL PHARMA	20MG	N209400 001	Jul 05, 2017

OMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED RELEASE;ORAL			
OMEPRAZOLE MAGNESIUM			
! DR REDDYS LABS LTD	EQ 20MG BASE	A078878 001	Jun 05, 2009
TABLET, DELAYED RELEASE;ORAL			
OMEPRAZOLE MAGNESIUM			
PERRIGO R AND D	EQ 20MG BASE	A204152 001	Jul 30, 2015
PRILOSEC OTC			
+! ASTRazeneca PHARMS	EQ 20MG BASE	N021229 001	Jun 20, 2003

OMEPRAZOLE; SODIUM BICARBONATE

CAPSULE;ORAL			
OMEPRAZOLE AND SODIUM BICARBONATE			
ACTAVIS ELIZABETH	20MG;1.1GM	A204137 001	Jul 15, 2016
AUROLIFE PHARMA LLC	20MG;1.1GM	A204923 001	Nov 07, 2016
PAR PHARM	20MG;1.1GM	A201946 001	Jul 15, 2016
PERRIGO R AND D	20MG;1.1GM	A201361 001	Jul 15, 2016
ZEGERID OTC			
+! BAYER HEALTHCARE	20MG;1.1GM	N022281 001	Dec 01, 2009
LLC			
FOR SUSPENSION;ORAL			
ZEGERID OTC			
+! BAYER HEALTHCARE	20MG/PACKET;1.68GM/PACKET	N022283 001	Jun 17, 2013
LLC			

ORLISTAT

CAPSULE;ORAL			
ALLI			
+! GLAXOSMITHKLINE	60MG	N021887 001	Feb 07, 2007
CONS			

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-18 (of 20)

OXYBUTYNIN

FILM, EXTENDED RELEASE;TRANSDERMAL	
OXYTROL FOR WOMEN	
+! ALLERGAN SALES LLC	3.9MG/24HR
	N202211 001 Jan 25, 2013

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC	
VISINE L.R.	
+! JOHNSON AND JOHNSON	0.025%
	N019407 001 Mar 31, 1989

PERMETHRIN

LOTION;TOPICAL	
NIX	
+! MEDTECH PRODUCTS	1%
PERMETHRIN	
ACTAVIS MID	1%
ATLANTIC	
PERRIGO NEW YORK	1%
	A075014 001 Mar 28, 2000
	A076090 001 Dec 20, 2001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION;ORAL	
GLYCOLAX	
KREMERS URBAN PHARMS	17GM/PACKET
	17GM/SCOOPFUL
MIRALAX	
+! BAYER HEALTHCARE LLC	17GM/SCOOPFUL
	N022015 001 Oct 06, 2006
POLYETHYLENE GLYCOL 3350	
AILEX PHARMS LLC	17GM/SCOOPFUL
ANI PHARMS INC	17GM/SCOOPFUL
MYLAN	17GM/PACKET
	17GM/SCOOPFUL
NEXGEN PHARMA	17GM/SCOOPFUL
NOVEL LABS INC	17GM/SCOOPFUL
NUVO PHARM INC	17GM/SCOOPFUL
PAR PHARM	17GM/SCOOPFUL
PERRIGO R AND D	17GM/PACKET
	A079214 001 Jan 31, 2013
STRIDES PHARMA	17GM/SCOOPFUL
	A090685 001 Oct 06, 2009
	A090685 002 Oct 06, 2009
	A203928 001 Aug 24, 2016
	A203928 002 Aug 24, 2016
	A090600 001 Oct 06, 2009
	A090600 002 Oct 06, 2009
	A202071 001 Dec 28, 2012
	A202850 001 Dec 15, 2015
	A078915 001 Oct 06, 2009
	A078915 002 Oct 06, 2009
	A090812 001 Oct 07, 2009
	A091077 001 Oct 06, 2009
	A206105 001 Oct 28, 2016
	A079214 001 Jan 31, 2013
	A090685 001 Oct 06, 2009
	A090685 002 Oct 06, 2009
	A203928 001 Aug 24, 2016
	A203928 002 Aug 24, 2016

POTASSIUM IODIDE

SOLUTION;ORAL	
POTASSIUM IODIDE	
MISSION PHARMACAL CO	65MG/ML
	A206211 001 Mar 24, 2016
THYROSCHILD	
! ARCO PHARMS LLC	65MG/ML
	A077218 001 Jan 12, 2005
TABLET;ORAL	
IOSAT	
+ ANBEX	65MG
+!	130MG
THYROSAFE	
! RECIP	65MG
	A076350 001 Sep 10, 2002

POVIDONE-IODINE

SOLUTION;TOPICAL	
POVIDONE IODINE	
+! ALLEGIANCE HLTHCARE	1%
	N019522 001 Mar 31, 1989
SPONGE;TOPICAL	
E-Z SCRUB 201	
+! BECTON DICKINSON	20%
	N019240 001 Nov 29, 1985
E-Z SCRUB 241	
+! BECTON DICKINSON	10%
	N019476 001 Jan 07, 1987

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL	
PSEUDOEPHEDRINE HYDROCHLORIDE	
AUROBINDO PHARMA LTD	120MG
L PERRIGO CO	120MG
SUN PHARM INDS LTD	120MG
SUDAFED 12 HOUR	
! MCNEIL CONS	120MG
	A209008 001 Jun 09, 2017
	A075153 001 Feb 26, 1999
	A077442 001 Sep 28, 2005
	A073585 001 Oct 31, 1991

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-19 (of 20)

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL	SUDAFED 24 HOUR	+! J AND J CONSUMER INC	240MG	N020021 002 Dec 15, 1992
--------------------------------	-----------------	-------------------------	-------	--------------------------

PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL	AFRINOL	+! SCHERING PLOUGH	120MG	N018191 001
--------------------------------	---------	--------------------	-------	-------------

PURIFIED WATER

SOLUTION; OPHTHALMIC	PUR-WASH	+! NIAGARA PHARMS	98.3%	N022305 001 Sep 01, 2011
----------------------	----------	-------------------	-------	--------------------------

RANITIDINE HYDROCHLORIDE

TABLET; ORAL	RANITIDINE HYDROCHLORIDE	APOTEX INC	EQ 75MG BASE	A075167 001 May 04, 2000
			EQ 150MG BASE	A200172 001 May 31, 2012
		AUROBINDO PHARMA LTD	EQ 75MG BASE	A207579 001 Nov 13, 2017
			EQ 150MG BASE	A207578 001 Nov 13, 2017
		DR REDDYS LABS LTD	EQ 75MG BASE	A075294 001 Mar 28, 2000
			EQ 150MG BASE	A078192 001 Aug 31, 2007
		IVAX SUB TEVA PHARMS	EQ 75MG BASE	A075296 001 Jan 14, 2000
		MYLAN	EQ 75MG BASE	A075497 001 Jan 14, 2000
		PERRIGO	EQ 75MG BASE	A076195 001 Aug 30, 2002
		PERRIGO R AND D	EQ 150MG BASE	A091429 001 May 11, 2011
			EQ 150MG BASE	A091429 002 May 11, 2011
		STRIDES PHARMA	EQ 75MG BASE	A201745 001 Feb 29, 2012
			EQ 150MG BASE	A200536 001 Jun 28, 2011
		WOCKHARDT	EQ 75MG BASE	A076760 001 Feb 24, 2006
ZANTAC 150				
+! SANOFI US			EQ 150MG BASE	N021698 001 Aug 31, 2004
+			EQ 150MG BASE	N021698 002 Mar 13, 2007
ZANTAC 75				
+	SANOFI US		EQ 75MG BASE	N020520 001 Dec 19, 1995

SODIUM CHLORIDE

AEROSOL, METERED; INHALATION	BRONCHO SALINE	+! BLAIREX	0.9%	N019912 001 Sep 03, 1992
------------------------------	----------------	------------	------	--------------------------

SODIUM FLUORIDE; TRICLOSAN

PASTE; DENTAL	COLGATE TOTAL	+! COLGATE PALMOLIVE	0.24%; 0.3%	N020231 001 Jul 11, 1997
---------------	---------------	----------------------	-------------	--------------------------

TERBINAFINE

GEL; TOPICAL	LAMISIL AT	+! NOVARTIS	1%	N021958 001 Jul 24, 2006
--------------	------------	-------------	----	--------------------------

TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL	LAMISIL	+! GLAXOSMITHKLINE	1%	N020980 001 Mar 09, 1999
		TERBINAFINE HYDROCHLORIDE		
		TARO	1%	A077511 001 Jul 02, 2007
SOLUTION; TOPICAL				
	LAMISIL AT	+! NOVARTIS	1%	N021124 001 Mar 17, 2000
SPRAY; TOPICAL				
	LAMISIL AT	+! NOVARTIS	1%	N021124 002 Mar 17, 2000

TIOCONAZOLE

OINTMENT; VAGINAL	TIOCONAZOLE	PERRIGO	6.5%	A075915 001 Nov 21, 2001
		VAGISTAT-1		
		+! COMBE	6.5%	N020676 001 Feb 11, 1997

38TH EDITION - 2018 - APPROVED DRUG PRODUCT LIST  
**OTC DRUG PRODUCT LIST**

4-20 (of 20)

TRIAMCINOLONE ACETONIDE

SPRAY, METERED;NASAL		
NASACORT ALLERGY 24 HOUR		
+! SANOFI AVENTIS US	0.055MG/SPRAY	N020468 002 Oct 11, 2013
TRIAMCINOLONE ACETONIDE		
PERRIGO ISRAEL	0.055MG/SPRAY	A078104 002 Nov 14, 2014

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT  
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT 4% SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

HAEMONETICS  
MANUFACTURING INC

N760305

Jun 30, 1978

ANTICOAGULANT CITRATE DEXTROSE SOLUTION (ACD)

INJECTABLE; INJECTION

CITRA LABS LLC

N020037

Aug 26, 2003

ACD-A SOLUTION

A010228

Feb 25, 2002

TERUMO BCT INC

N000922

Aug 29, 2002

ADSOL WITH ACD-A

A980728

Feb 06, 2002

FENWAL INC

N001214

May 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION FORMULA A

HAEMONETICS CORP

A710497

Nov 06, 1987

AS3 SOLUTION/ACD-A

TERUMO BCT INC

NONE

HAEMONETICS

MANUFACTURING INC

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N160918

Mar 17, 1978

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP (ACD-A)

INJECTABLE; INJECTION

NONE

ARTERIOCYTE MEDICAL  
SYSTEMS, INC

N160767

May 11, 2012

ANTICOAGULANT CITRATE PHOSPHATE 2X DEXTROSE SOLUTION (CP2D)

INJECTABLE; INJECTION

CITRATE PHOSPHATE DOUBLE DEXTROSE/ADDITIVE SOLUTION 3

HAEMONETICS CORP

N000127

Jan 18, 2002

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION

INJECTABLE; INJECTION

NONE

TERUMO MEDICAL CORP

N820528

Nov 03, 1982

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION (CPDA)

INJECTABLE; INJECTION

CPDA-1 BLOOD-PACK UNIT (PL 146 PLASTIC) 250, 450, 500 ML BLOOD PACK UNITS

FENWAL INC

N770420

May 12, 1978

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION USP

INJECTABLE; INJECTION

BLOOD PACK UNIT CPDA-1 IN PLASTIC CONTAINER

FENWAL INC

N940404

Jul 28, 1994

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT  
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS  
MANUFACTURING INC

N800077

Nov 06, 1980

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION

INJECTABLE; INJECTION

ADSOL IN PLASTIC CONTAINER  
FENWAL INC

N900223

Dec 27, 1991

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

INJECTABLE; INJECTION

CPD ANTICOAGULANT IN PL 2209 PLASTIC CONTAINER

FENWAL INC

N900224

Dec 27, 1991

MACOPRODUCTIONS SAS CPD/AS-1: MACOPHARMA LEUCOFLEX MTL1 LEUKOREDUCTION SYSTEM FOR BLOOD  
COMPONENTS KNOWN AS MTL1-WB

MACOPRODUCTIONS SAS

N040083

Nov 21, 2005

NONE

TERUMO BCT INC

A070025

Jan 06, 2008

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC

N170401

Dec 06, 1977

HAEMONETICS

N811012

Jun 28, 1983

MANUFACTURING INC

N800222

Aug 23, 1982

TERUMO MEDICAL CORP

N781211

Jun 10, 1981

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-1:DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

ADSOL RED BLOOD CELL PRESERVATIVE SOLUTION

FENWAL INC

N811104

May 16, 1983

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-5:DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

OPTISOL RED BLOOD CELL PRESERVATIVE SOLUTION

TERUMO MEDICAL CORP

N880217

Oct 07, 1988

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:AS-3: CITRIC ACID USP; MONOBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;  
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-3 NUTRICEL ADDITIVE SYSTEM

HAEMONETICS            0.042GM/100ML; 0.276GM/100ML;  
MANUFACTURING INC    0.410GM/100ML; 0.30GM/100ML;  
                          1.10GM/100ML; 0.588GM/100ML

N820915

Oct 19, 1984

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT  
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:  
AS-2: CITRIC ACID USP; DIBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;  
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-2 NUTRICEL ADDITIVE SYSTEM

MEDSEP CORP            0.042GM/100ML;0.285GM/100ML;  
                          0.718GM/100ML;0.017GM/100ML;  
                          0.396GM/100ML;0.588GM/100ML

N820915

Sep 22, 1983

ANTICOAGULANT SODIUM CITRATE 4% SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS  
CORPORATION

N980123

Mar 03, 2000

ANTICOAGULANT SODIUM CITRATE SOLUTION

INJECTABLE; INJECTION

TRICITRASOL

CYTOSOL  
LABORATORIES INC

N010409

Jul 10, 2003

ANTICOAGULANT SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

FENWAL INC  
TERUMO MEDICAL CORP

N770923

Jan 20, 1978

N781214

Feb 08, 1980

CORD BLOOD STERILE COLLECTION BAG, ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION  
(CPD)

STERILE CORD BLOOD COLLECTION UNIT

NONE

MACOPHARMA

N125552

Dec 21, 2016

DEXTRAN 1 IN SODIUM CHLORIDE 0.6%

INJECTABLE; INJECTION

PROMIT

MEDA AB

N830715

Oct 30, 1984

DEXTRAN 40, 10% IN DEXTROSE 5%

INJECTABLE; INJECTION

LMD IN GLASS BOTTLE

HOSPIRA INC            10GM/100ML;5GM/100ML

A720563

Oct 30, 1992

DEXTRAN 40, 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

LMD IN PLASTIC CONTAINER

HOSPIRA INC            10GM/100ML;0.9GM/100ML

A720562

Oct 30, 1992

HETASTARCH 6% IN LACTATED ELECTROLYTE INJECTION

INJECTABLE; INJECTION

HEXTEND

BIOTIME INC            6GM/100ML

N200952

Mar 31, 1999

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT  
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

**HETASTARCH 6% IN SODIUM CHLORIDE 0.9%**

INJECTABLE; INJECTION

6% HETASTARCH IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
HOSPIRA INC	6GM/100ML;0.9GM/100ML	A740193	Jan 30, 1995
HESSPAN IN PLASTIC CONTAINER			
B BRAUN MEDICAL INC	6GM/100ML;0.9GM/100ML	N890105	Apr 04, 1991
NONE			
TEVA PARENTERAL MEDICINES INC	6GM/100ML;0.9GM/100ML	A740592	Nov 12, 1998

**HYDROXYETHYL STARCH 130/0.4 IN 6% SODIUM CHLORIDE 0.9%**

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE INFUSED DIRECTLY TO THE PATIENT.

NONE			
B. BRAUN MEDICAL VOLUVEN		A110013	Jan 09, 2015
FRESENIUS KABI DEUTSCHLAND GMBH	6GM/100ML;0.9GM/100ML	N070012	Dec 27, 2007

**ISOPLATE SOLUTION IN THE 500 ML EXCEL CONTAINER**

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE INFUSED DIRECTLY TO THE PATIENT.

ISOPLATE SOLUTION HAEMONETICS CORP		N90067	Mar 05, 2013
------------------------------------	--	--------	--------------

**LEUKOCYTE REDUCTION FILTRATION SYSTEM FOR WHOLE BLOOD WITH CPD ANTICOAGULANT AND SOLX ADDITIVE**

INJECTABLE; INJECTION

LEUKOSEP HWB-600-XL HAEMONETICS CORP		N110059	Apr 25, 2013
--------------------------------------	--	---------	--------------

**RED BLOOD CELL PROCESSING SOLUTION**

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE INFUSED DIRECTLY TO THE PATIENT.

REJUVESOL CITRA LABS LLC		N950522	Feb 26, 1997
--------------------------	--	---------	--------------

**SODIUM CHLORIDE; SODIUM ACETATE; SODIUM CITRATE DIHYDRATE; SODIUM PHOSPHATE, DIABASIC ANHYDROUS; SODIUM PHOSPHATE MONOBASIC, MONOHYDRATE**

STORAGE/PROCESSING SOLUTION ONLY; SHOULD NEVER BE INFUSED DIRECTLY TO THE PATIENT.

INTERSOL SOLUTION FENWAL INC.	2.26G/500ML; 2.21G/500ML; 1.59G/500ML; 1.53G/500ML; 0.465G/500ML	N080041	Dec 09, 2009
-------------------------------	---	---------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-1(of 375)

\*\* See List Footnote

ABARELIX

INJECTABLE; INTRAMUSCULAR  
 PLENAXIS  
 SPECIALITY EUROPEAN 100MG/VIAL

N021320 001 Nov 25, 2003

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL  
 CAMPRAL  
 + FOREST LABS 333MG \*\*

N021431 001 Jul 29, 2004

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL  
 ACEBUTOLOL HYDROCHLORIDE  
 WATSON LABS EQ 200MG BASE  
 EQ 400MG BASE

A074007 001 Oct 18, 1995  
 A074007 002 Oct 18, 1995ACETAMINOPHEN

INJECTABLE; INJECTION  
 INJECTAPAP  
 ORTHO MCNEIL PHARM 100MG/ML  
 SUPPOSITORY; RECTAL  
 ACEPHEN  
 G AND W LABS 120MG  
 ACETAMINOPHEN  
 ABLE 120MG  
 325MG  
 650MG  
 ACINO PRODS 120MG  
 650MG  
 TYLENOL  
 J AND J CONSUMER INC 120MG  
 650MG

N017785 001 Mar 07, 1986  
 A072218 001 Mar 27, 1992  
 A073106 001 Feb 27, 1995  
 A073107 001 Feb 27, 1995  
 A073108 001 Feb 27, 1995  
 A071010 001 May 12, 1987  
 A071011 001 May 12, 1987  
 N017756 002  
 N017756 001

TABLET, EXTENDED RELEASE; ORAL  
 ACETAMINOPHEN  
 SUN PHARM INDs LTD 650MG

A090205 001 Nov 18, 2009

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL  
 ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE  
 MIKART 150MG;180MG;15MG  
 150MG;180MG;30MG  
 150MG;180MG;60MG  
 CODEINE, ASPIRIN, APAP FORMULA NO. 2  
 SCHERER LABS 150MG;180MG;15MG  
 CODEINE, ASPIRIN, APAP FORMULA NO. 3  
 SCHERER LABS 150MG;180MG;30MG  
 CODEINE, ASPIRIN, APAP FORMULA NO. 4  
 SCHERER LABS 150MG;180MG;60MG

A081095 001 Oct 26, 1990  
 A081096 001 Oct 26, 1990  
 A081097 001 Oct 26, 1990  
 A085640 001  
 A085639 001  
 A085638 001ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL  
 BANCAP  
 FOREST PHARMS 325MG;50MG  
 BUCET  
 MALLINCKRODT 650MG;50MG  
 PHRENILIN FORTE  
 VALEANT 650MG;50MG  
 TENCON  
 MALLINCKRODT 650MG;50MG  
 TRIAPRIN  
 DUNHALL 325MG;50MG

A088889 001 Jan 16, 1986  
 A088991 001 Jun 28, 1985  
 A088831 001 Jun 19, 1985  
 A089405 001 May 15, 1990  
 A089268 001 Jul 02, 1987  
 A089568 001 Oct 05, 1988  
 A087550 001 Oct 19, 1984

TABLET; ORAL  
 BUTALBITAL AND ACETAMINOPHEN  
 HALSEY 325MG;50MG  
 WATSON LABS 325MG;50MG  
 BUTAPAP  
 MIKART 650MG;50MG  
 PHRENILIN  
 VALEANT 325MG;50MG \*\*  
 SEDAPAP  
 MAYRAND 650MG;50MG

A089988 001 Oct 26, 1992  
 A087811 001 Jun 19, 1985  
 A088944 001 Oct 17, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-2(of 375)

\*\* See List Footnote

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE;ORAL

ANOQUAN

SHIRE	325MG;50MG;40MG	A087628 001 Oct 01, 1986
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE		
GRAHAM DM	325MG;50MG;40MG	A088743 001 Jul 18, 1985
	325MG;50MG;40MG	A088765 001 Mar 27, 1985
	325MG;50MG;40MG	A089067 001 Apr 19, 1985
HIKMA PHARMS	500MG;50MG;40MG	A040261 001 Oct 28, 1998
MALLINCKRODT	325MG;50MG;40MG	A088758 001 Mar 27, 1985
BUTALIBITAL, ACETAMINOPHEN AND CAFFEINE		
GILBERT LABS	325MG;50MG;40MG **	A088825 001 Dec 05, 1984
ESGIC-PLUS		
MIKART	500MG;50MG;40MG	A040085 001 Mar 28, 1996
FEMCET		
MALLINCKRODT	325MG;50MG;40MG	A089102 001 Jun 19, 1985
MEDIGESIC PLUS		
US CHEM	325MG;50MG;40MG	A089115 001 Jan 14, 1986
TRIAD		
MALLINCKRODT	325MG;50MG;40MG	A089023 001 Jun 19, 1985
TABLET;ORAL		
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE		
ABLE	325MG;50MG;40MG	A040390 001 Jul 23, 2001
	500MG;50MG;40MG	A040394 001 Jul 23, 2001
GILBERT LABS	325MG;50MG;40MG	A087629 001 Nov 13, 1984
HIKMA PHARMS	500MG;50MG;40MG	A040336 001 Aug 18, 1999
MIKART	750MG;50MG;40MG	A040496 001 Dec 23, 2003
MIRROR PHARMS LLC	500MG;50MG;40MG	A040883 001 Dec 23, 2008
NOVAST LABS LTD	325MG;50MG;40MG	A040864 001 Dec 01, 2008
SUN PHARM INDUSTRIES	325MG;50MG;40MG	A040601 001 Jul 29, 2005
VINTAGE PHARMS	500MG;50MG;40MG	A040513 001 Aug 25, 2003
WATSON LABS	325MG;50MG;40MG	A089536 001 Feb 16, 1988
	500MG;50MG;40MG	A040267 001 Jul 30, 1998
ESGIC		
FOREST PHARMS	325MG;50MG;40MG	A089660 001 Dec 23, 1988
ESGIC-PLUS		
MIKART	500MG;50MG;40MG	A089451 001 May 23, 1988

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE		
ABLE	325MG;50MG;40MG;30MG	A076528 001 Aug 21, 2003
HIKMA INTL PHARMS	325MG;50MG;40MG;30MG	A075618 001 Mar 23, 2001
PHRENILIN WITH CAFFEINE AND CODEINE		
VALEANT	325MG;50MG;40MG;30MG	A074911 001 Aug 22, 2001

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE;ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE		
MIKART	356.4MG;30MG;16MG	A040109 001 Aug 26, 1997
WRASER PHARMS LLC	356.4MG;30MG;16MG	A040688 001 Apr 03, 2007
DHC PLUS		
PHARM RES ASSOC	356.4MG;30MG;16MG	A088584 001 Mar 04, 1986
SYNALGOS-DC-A		
LEITNER PHARMS	356.4MG;30MG;16MG	A089166 001 May 14, 1986

TABLET;ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE		
BOCA PHARMA LLC	712.8MG;60MG;32MG	A040701 001 Apr 03, 2007
MIKART	712.8MG;60MG;32MG	A040316 001 Apr 28, 1999
WEST-WARD PHARM CORP	712.8MG;60MG;32MG	A040637 001 Sep 22, 2006

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET;ORAL

TAVIST ALLERGY/SINUS/HEADACHE

NOVARTIS 500MG;EQ 0.25MG BASE;30MG

N021082 001 Mar 01, 2001

**DISCONTINUED DRUG PRODUCT LIST**

6-3(of 375)

\*\* See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

## CAPSULE;ORAL

## ACETAMINOPHEN AND CODEINE PHOSPHATE

TEVA	300MG;15MG	A088537 001 Jun 04, 1984
	300MG;30MG	A088324 001 Dec 29, 1983
	300MG;60MG	A088599 001 Jun 01, 1984

## PHENAPHEN W/ CODEINE NO. 2

ROBINS AH	325MG;15MG	A084444 001
-----------	------------	-------------

## PHENAPHEN W/ CODEINE NO. 3

ROBINS AH	325MG;30MG	A084445 001
-----------	------------	-------------

## PHENAPHEN W/ CODEINE NO. 4

ROBINS AH	325MG;60MG	A084446 001
-----------	------------	-------------

## PROVAL #3

SOLVAY	325MG;30MG	A085685 001
--------	------------	-------------

## TYLENOL W/ CODEINE NO. 3

ORTHO MCNEIL PHARM	300MG;30MG	A087422 001
--------------------	------------	-------------

## TYLENOL W/ CODEINE NO. 4

ORTHO MCNEIL PHARM	300MG;60MG	A087421 001
--------------------	------------	-------------

## SOLUTION;ORAL

## ACETAMINOPHEN AND CODEINE PHOSPHATE

ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A085861 001
ALLIED PHARMA INC	120MG/5ML;12MG/5ML	A086366 001
DAVA PHARMS INC	120MG/5ML;12MG/5ML	A040098 001 Sep 20, 1996

## TYLENOL W/ CODEINE

ORTHO MCNEIL PHARM	120MG/5ML;12MG/5ML	A085057 001
--------------------	--------------------	-------------

## SUSPENSION;ORAL

## CAPITAL AND CODEINE

ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	A085883 001
VALEANT PHARMS LLC	120MG/5ML;12MG/5ML	A086024 001

## TABLET;ORAL

## ACETAMINOPHEN AND CODEINE PHOSPHATE

ABLE	300MG;30MG	A040452 001 Aug 01, 2002
	300MG;60MG	A040459 001 Aug 01, 2002

AM THERAP	300MG;15MG	A089478 001 Mar 03, 1987
	300MG;15MG	A089481 001 Mar 03, 1987
	300MG;30MG	A089479 001 Mar 03, 1987
	300MG;30MG	A089482 001 Mar 03, 1987

	300MG;60MG	A089480 001 Mar 03, 1987
	300MG;60MG	A089483 001 Mar 03, 1987

ANDA REPOSITORY	300MG;15MG	A089673 002 Feb 10, 1988
	300MG;30MG	A089673 003 Feb 10, 1988
	300MG;60MG	A089673 001 Feb 10, 1988

DURAMED PHARMS BARR	300MG;15MG	A040223 001 Nov 18, 1997
	300MG;15MG	A088353 001 Feb 06, 1984
	300MG;30MG	A040223 002 Nov 18, 1997

	300MG;30MG	A088354 001 Feb 06, 1984
	300MG;60MG	A040223 003 Nov 18, 1997
	300MG;60MG	A088355 001 Feb 06, 1984

EVERYLIFE	325MG;30MG	A085217 001
HALSEY	300MG;15MG	A083871 001
	300MG;30MG	A083872 001

	300MG;60MG	A086549 001
KV PHARM	300MG;30MG	A085288 001
	300MG;60MG	A085365 001

	325MG;15MG	A085364 001
	325MG;45MG **	A085363 001

LEDERLE	300MG;30MG	A087141 001
MIKART	300MG;30MG	A089238 001 Feb 25, 1986
	300MG;60MG	A089244 001 Feb 25, 1986

	650MG;30MG	A089231 001 Mar 03, 1986
	650MG;60MG	A089363 001 Sep 09, 1991

MUTUAL PHARM	300MG;15MG	A085795 001
	300MG;30MG	A085794 001
	300MG;60MG	A087653 001 Apr 13, 1982

PURACAP PHARM	300MG;30MG	A087762 001 Dec 10, 1982
PUREPAC PHARM	300MG;30MG	A086681 001
	300MG;30MG	A089080 001 Jul 17, 1986
	300MG;60MG	A086683 001

ROXANE	300MG;15MG	A084659 001
--------	------------	-------------

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-4(of 375)

\*\* See List Footnote

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET;ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

	300MG;30MG	A084656 001
	300MG;60MG	A084667 001
	500MG;15MG	A089511 001 Apr 25, 1989
	500MG;30MG	A089512 001 Apr 25, 1989
	500MG;60MG	A089513 001 Apr 25, 1989
SANDOZ	300MG;15MG	A087433 001
	300MG;30MG	A081250 001 Jul 16, 1992
	300MG;30MG	A085291 002
	300MG;30MG	A085917 001
	300MG;60MG	A081249 001 Jul 16, 1992
	300MG;60MG	A085964 001
	300MG;60MG	A087423 001
SUPERPHARM	300MG;15MG	A089183 001 Oct 18, 1985
	300MG;30MG	A089184 001 Oct 18, 1985
	300MG;30MG	A089253 001 May 19, 1986
	300MG;60MG	A089185 001 Oct 18, 1985
	300MG;60MG	A089254 001 May 19, 1986
USL PHARMA	300MG;30MG	A087919 001 Jun 22, 1982
	300MG;60MG	A087920 001 Jun 22, 1982
VALEANT PHARM INTL	300MG;30MG	A085896 001
VITARINE	300MG;30MG	A085676 001
WARNER CHILCOTT	300MG;15MG	A085992 001
	300MG;30MG	A085218 002
	300MG;60MG	A087306 001
WATSON LABS	300MG;15MG	A087277 001 May 26, 1982
	300MG;15MG	A089997 001 Dec 28, 1994
	300MG;30MG	A087276 001 May 26, 1982
	300MG;30MG	A089998 001 Dec 28, 1994
	300MG;60MG	A087275 001 May 26, 1982
	300MG;60MG	A089999 001 Dec 28, 1994
WATSON LABS FLORIDA	300MG;15MG	A040443 001 Jan 22, 2003
	300MG;30MG	A040443 002 Jan 22, 2003
	300MG;60MG	A040443 003 Jan 22, 2003
WHITEWORTH TOWN PLSN	300MG;30MG	A084360 001
	300MG;60MG	A085607 001
CAPITAL AND CODEINE		
CARNRICK	325MG;30MG	A083643 001
CODRIX		
WATSON LABS FLORIDA	500MG;15MG	A040447 001 Feb 26, 2003
	500MG;30MG	A040441 001 Mar 27, 2003
	500MG;60MG	A040488 001 Mar 28, 2003
EMPRACET W/ CODEINE PHOSPHATE #3		
GLAXOSMITHKLINE	300MG;30MG	A083951 001
EMPRACET W/ CODEINE PHOSPHATE #4		
GLAXOSMITHKLINE	300MG;60MG	A083951 002
PAPA-DEINE #3		
VANGARD	300MG;30MG	A088037 001 Mar 20, 1984
PAPA-DEINE #4		
VANGARD	300MG;60MG	A088715 001 Mar 20, 1984
PHENAPHEN-650 W/ CODEINE		
ROBINS AH	650MG;30MG	A085856 001
TYLENOL W/ CODEINE		
ORTHO MCNEIL PHARM	325MG; 7.5MG **	A085056 001
	325MG;15MG **	A085056 002
	325MG;30MG **	A085056 003
	325MG;60MG **	A085056 004
TYLENOL W/ CODEINE NO. 1		
JANSSEN PHARMS	300MG;7.5MG	A085055 001
TYLENOL W/ CODEINE NO. 2		
JANSSEN PHARMS	300MG;15MG	A085055 002

**DISCONTINUED DRUG PRODUCT LIST**

6-5(of 375)

\*\* See List Footnote

ACETAMINOPHEN; DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

DRIXORAL PLUS

SCHERING PLOUGH 500MG;3MG;60MG

N019453 001 May 22, 1987

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE

CENT PHARMS 500MG;5MG

A088898 001 Mar 27, 1985

ALLAY

IVAX PHARMS 500MG;5MG

A089907 001 Jan 13, 1989

BANCAP HC

FOREST PHARMS 500MG;5MG

A087961 001 Mar 17, 1983

CO-GESIC

CENT PHARMS 500MG;5MG

A089360 001 Mar 02, 1988

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MALLINCKRODT 500MG;5MG

A088956 001 Jul 19, 1985

500MG;5MG

A089006 001 Aug 09, 1985

MIKART 500MG;5MG

A081067 001 Nov 30, 1989

500MG;5MG

A081068 001 Nov 30, 1989

500MG;5MG

A081069 001 Nov 30, 1989

500MG;5MG

A081070 001 Nov 30, 1989

500MG;5MG

A089008 001 Feb 21, 1986

LORCET-HD

MALLINCKRODT 500MG;5MG

A087336 001 Jul 08, 1982

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

MALLINCKRODT 500MG/15ML;7.5MG/15ML

A040418 001 Jun 27, 2001

MALLINCKRODT INC 500MG/15ML;10MG/15ML

A040508 001 Aug 29, 2003

MIKART 500MG/15ML;5MG/15ML

A081226 001 Oct 27, 1992

500MG/15ML;5MG/15ML

A089557 001 Apr 29, 1992

500MG/15ML;7.5MG/15ML

A081051 001 Aug 28, 1992

NESHER PHARMS 500MG/15ML;7.5MG/15ML

A040366 001 Jan 23, 2002

PHARM ASSOC 500MG/15ML;7.5MG/15ML

A040182 001 Mar 13, 1998

VINTAGE PHARMS 500MG/15ML;7.5MG/15ML

A040520 001 Oct 30, 2003

ZYFREL

CYPRESS PHARM INC 325MG/15ML;7.5MG/15ML

A090468 001 Apr 14, 2016

TABLET; ORAL

ANEXSIA

MALLINCKRODT 500MG;5MG

A089160 001 Apr 23, 1987

750MG;10MG

A040468 001 Oct 31, 2002

ANEXSIA 7.5/650

MALLINCKRODT 650MG;7.5MG

A089725 001 Sep 30, 1987

CO-GESIC

UCB INC 500MG;5MG

A087757 001 May 03, 1982

DURADYNE DHC

FOREST PHARMS 500MG;5MG

A087809 001 Mar 17, 1983

HY-PHEN

ASCHER 500MG;5MG

A087677 001 May 03, 1982

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

ABLE 325MG;5MG

A040478 001 Nov 08, 2002

325MG;7.5MG

A040464 001 Oct 23, 2002

325MG;10MG

A040464 002 Oct 23, 2002

500MG;5MG

A040477 001 Nov 06, 2002

500MG;7.5MG

A040490 001 May 21, 2003

500MG;10MG

A040473 001 Nov 06, 2002

650MG;7.5MG

A040474 001 Jan 02, 2003

650MG;10MG

A040476 001 Oct 23, 2002

750MG;7.5MG

A040469 001 Oct 25, 2002

AMNEAL PHARMS NY 500MG;5MG

A040729 001 Aug 25, 2006

500MG;7.5MG

A040748 001 Aug 25, 2006

500MG;10MG

A040813 001 Feb 23, 2007

650MG;7.5MG

A040754 001 Aug 25, 2006

650MG;10MG

A040757 001 Aug 25, 2006

750MG;7.5MG

A040769 001 Aug 28, 2006

APIL 500MG;10MG

A040148 002 Feb 14, 1997

BARR 500MG;2.5MG

A040307 001 Jul 26, 2000

500MG;5MG

A040308 001 Jul 26, 2000

500MG;5MG

A088577 001 Dec 21, 1984

500MG;7.5MG

A040307 002 Jul 26, 2000

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-6(of 375)

\*\* See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	500MG;10MG	A040309 001	Jul 26, 2000
	650MG;7.5MG	A040307 003	Jul 26, 2000
	650MG;10MG	A040307 004	Jul 26, 2000
	750MG;7.5MG	A040308 002	Jul 26, 2000
CARACO	500MG;5MG	A090265 001	Dec 23, 2008
	500MG;7.5MG	A090265 002	Dec 23, 2008
	500MG;10MG	A090265 003	Dec 23, 2008
	650MG;7.5MG	A090380 001	Dec 23, 2008
	650MG;10MG	A090380 002	Dec 23, 2008
	660MG;10MG	A090380 003	Dec 23, 2008
	750MG;7.5MG	A090380 004	Dec 23, 2008
HALSEY	500MG;5MG	A089554 001	Jun 12, 1987
IVAX PHARMS	500MG;5MG	A089696 001	Apr 21, 1988
MALLINCKRODT	500MG;5MG	A040084 002	Jun 01, 1995
	500MG;7.5MG	A040201 001	Feb 27, 1998
	500MG;10MG	A040201 002	Feb 27, 1998
	650MG;10MG	A040084 004	Oct 16, 1996
	660MG;10MG	A040084 003	Jul 29, 1996
	750MG;7.5MG	A040084 001	Jun 01, 1995
MIKART	500MG;2.5MG	A089698 001	Aug 25, 1989
	500MG;5MG	A089271 001	Jul 16, 1986
	500MG;5MG	A089697 001	Jan 28, 1992
	500MG;7.5MG	A089699 001	Aug 25, 1989
	650MG;5MG	A040849 001	Jun 09, 2010
	650MG;7.5MG	A089689 001	Jun 29, 1988
	650MG;10MG	A081223 001	May 29, 1992
MUTUAL PHARM	500MG;5MG	A040236 001	Sep 25, 1997
	650MG;7.5MG	A040240 002	Nov 26, 1997
	650MG;10MG	A040240 001	Nov 26, 1997
	750MG;7.5MG	A040236 002	Sep 25, 1997
RANBAXY	500MG;5MG	A040825 001	Aug 16, 2007
	500MG;10MG	A040824 001	Aug 16, 2007
RANBAXY LABS LTD	750MG;7.5MG	A040822 001	Aug 16, 2007
SANDOZ	500MG;5MG	A040149 001	Jan 27, 1997
	750MG;7.5MG	A040149 002	Jan 27, 1997
SUN PHARM INDs LTD	325MG;10MG	A040826 001	Aug 16, 2007
UCB INC	500MG;10MG	A040210 001	Aug 13, 1997
	650MG;7.5MG	A040134 001	Nov 21, 1996
USL PHARMA	500MG;5MG	A089290 001	May 29, 1987
	500MG;5MG	A089291 001	May 29, 1987
VINTAGE PHARMS	500MG;2.5MG	A040144 002	Apr 25, 1997
	500MG;5MG	A089831 001	Sep 07, 1988
	500MG;5MG	A089971 001	Dec 02, 1988
	500MG;7.5MG	A040144 001	Feb 22, 1996
	500MG;10MG	A040356 001	May 31, 2000
	650MG;7.5MG	A040155 001	Apr 14, 1997
	650MG;10MG	A040143 001	Feb 22, 1996
	660MG;10MG	A040358 001	May 31, 2000
	750MG;7.5MG	A040157 001	Apr 12, 1996
VINTAGE PHARMS LLC	500MG;5MG	A040281 001	Sep 30, 1998
	500MG;7.5MG	A040280 001	Sep 30, 1998
	650MG;7.5MG	A040280 002	Sep 30, 1998
	650MG;10MG	A040280 003	Sep 30, 1998
	750MG;7.5MG	A040281 002	Sep 30, 1998
WATSON LABS	325MG;7.5MG	A040248 001	Apr 28, 2000
	325MG;10MG	A040248 002	Apr 28, 2000
	500MG;2.5MG	A040123 003	Mar 04, 1996
	500MG;2.5MG	A081079 001	Aug 30, 1991
	500MG;5MG	A040122 001	Mar 04, 1996
	500MG;7.5MG	A089883 001	Dec 01, 1988
	500MG;7.5MG	A040123 004	Mar 04, 1996
	500MG;7.5MG	A081080 001	Aug 30, 1991
	650MG;7.5MG	A040094 001	Sep 29, 1995
	650MG;7.5MG	A040123 001	Mar 04, 1996
	650MG;10MG	A040094 002	Sep 29, 1995
	650MG;10MG	A040123 002	Mar 04, 1996

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-7(of 375)

\*\* See List Footnote

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET;ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

	660MG;10MG	A040094 003	Aug 08, 2000
	750MG;7.5MG	A040122 002	Mar 04, 1996
	750MG;7.5MG	A081083 001	Aug 30, 1991
	750MG;10MG	A040094 004	Mar 22, 1999
WATSON LABS FLORIDA	500MG;5MG	A040493 001	May 28, 2003
	660MG;10MG	A040495 001	May 28, 2003
	750MG;7.5MG	A040494 001	May 28, 2003
LORTAB			
UCB INC	500MG;5MG	A087722 001	Jul 09, 1982
	500MG;10MG	A040100 001	Jan 26, 1996
NORCET			
ABANA	500MG;5MG	A088871 001	May 15, 1986
TYCOLET			
ORTHO MCNEIL PHARM	500MG;5MG	A089385 001	Aug 27, 1986
VICODIN			
ABBOTT	500MG;5MG	A085667 001	
ABBVIE	500MG;5MG	A088058 001	Jan 07, 1983
VICODIN ES			
ABBVIE	750MG;7.5MG	A089736 001	Dec 09, 1988
VICODIN HP			
ABBVIE	660MG;10MG	A040117 001	Sep 23, 1996
ZYDONE			
VINTAGE PHARMS LLC	400MG;5MG	A040288 001	Nov 27, 1998
	400MG;7.5MG	A040288 002	Nov 27, 1998
	400MG;10MG	A040288 003	Nov 27, 1998

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE;ORAL

OXYCODONE AND ACETAMINOPHEN

ACTAVIS ELIZABETH	500MG;5MG	A040199 001	Dec 30, 1998
BARR	500MG;5MG	A040304 001	Oct 02, 2000
DURAMED PHARMS BARR	500MG;5MG	A040289 001	Mar 16, 1999
HALSEY	500MG;5MG	A089994 001	May 04, 1989
MALLINCKRODT	500MG;5MG	A040257 001	Aug 04, 1998
MUTUAL PHARM	500MG;5MG	A040219 001	Jan 22, 1998
VINTAGE PHARMS	500MG;5MG	A040106 001	Jul 30, 1996
VINTAGE PHARMS LLC	500MG;5MG	A040303 001	Dec 30, 1999
WATSON LABS	500MG;5MG	A040234 001	Oct 30, 1997

ROXILOX

ROXANE 500MG;5MG

A040061 001 Jul 03, 1995

TYLOX

JANSSEN PHARMS 500MG;5MG

A088790 001 Dec 12, 1984

TYLOX-325

ORTHO MCNEIL PHARM 325MG;5MG

A088246 001 Nov 08, 1984

SOLUTION;ORAL

ROXICET

WEST-WARD PHARMS INT 325MG/5ML;5MG/5ML

A089351 001 Dec 03, 1986

TABLET;ORAL

OXYCODONE 2.5/APAP 500

BRISTOL MYERS SQUIBB 500MG;2.5MG

A085910 001

OXYCODONE 5/APAP 500

BRISTOL MYERS SQUIBB 500MG;5MG

A085911 001

OXYCODONE AND ACETAMINOPHEN

ACTAVIS ELIZABETH 325MG;5MG

A040203 001 Mar 15, 1999

325MG;7.5MG

A040800 001 Apr 03, 2012

325MG;10MG

A040800 002 Apr 03, 2012

AMNEAL PHARMS NY 500MG;7.5MG

A040789 001 Nov 27, 2007

650MG;10MG

A040789 002 Nov 27, 2007

BARR 325MG;5MG

A087406 001

DURAMED PHARMS BARR 325MG;5MG

A040272 001 Jun 30, 1998

MALLINCKRODT 500MG;7.5MG

A040550 001 Jun 30, 2004

650MG;10MG

A040550 002 Jun 30, 2004

MAYNE PHARMA INC 500MG;7.5MG

A090177 005 Oct 20, 2008

650MG;10MG

A090177 006 Oct 20, 2008

MIKART 400MG;2.5MG

A040679 001 May 16, 2006

400MG;5MG

A040687 001 Apr 27, 2006

400MG;7.5MG

A040698 001 Apr 27, 2006

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-8(of 375)

\*\* See List Footnote

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

OXYCODONE AND ACETAMINOPHEN

	400MG;10MG	A040692 001	Apr 27, 2006
	500MG;10MG	A040676 001	Apr 19, 2006
WATSON LABS	500MG;7.5MG	A040371 001	Dec 29, 2000
	650MG;10MG	A040371 002	Dec 29, 2000
PERCOCET			
VINTAGE PHARMS LLC	325MG;5MG	A085106 002	
	500MG;7.5MG	A040341 001	Jul 26, 1999
	650MG;10MG	A040341 002	Jul 26, 1999
ROXICET 5/500			
ROXANE	500MG;5MG	A089775 001	Jan 12, 1989
TABLET, EXTENDED RELEASE;ORAL			
XARTEMIS XR			
+ MALLINCKRODT INC	325MG;7.5MG	N204031 001	Mar 11, 2014

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

CAPSULE;ORAL

TYLOX

ORTHO MCNEIL PHARM 500MG;4.5MG;0.38MG

A085375 001

ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET;ORAL

ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE

GAVIS PHARMS	650MG;EQ 25MG BASE	A076202 001	Aug 02, 2002
WATSON LABS	650MG;EQ 25MG BASE	A074699 001	Mar 24, 2000
TALACEN			

SANOFI AVENTIS US 650MG;EQ 25MG BASE

N018458 001 Sep 23, 1982

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET;ORAL

DARVOCET

AAIPHARMA LLC 325MG;32.5MG

N016844 001

DOLENE AP-65

LEDERLE 650MG;65MG

A085100 001

PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN

MYLAN 325MG;32MG

A083689 001

650MG;65MG

A083978 001

SANDOZ 650MG;65MG

A089959 001 Jul 18, 1989

VINTAGE PHARMS 650MG;65MG

A040507 001 Jul 30, 2003

WATSON LABS 650MG;65MG

A040139 001 Dec 16, 1996

WYGESIC

CARACO 650MG;65MG

A084999 001

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET;ORAL

DARVOCET A500

XANODYNE PHARM 500MG;100MG

A076429 001 Sep 10, 2003

DARVOCET-N 100

XANODYNE PHARM 650MG;100MG

N017122 002

DARVOCET-N 50

XANODYNE PHARM 325MG;50MG

N017122 001

PROPACET 100

TEVA 650MG;100MG

A070107 001 Jun 12, 1985

PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

ABLE 650MG;100MG

A075838 001 Jul 11, 2001

ACTAVIS ELIZABETH 650MG;100MG

A070910 001 Jan 02, 1987

CORNERSTONE 325MG;100MG

A076743 001 May 07, 2004

500MG;100MG

A076750 001 Jun 28, 2004

HALSEY 325MG;50MG

A072105 001 May 13, 1988

650MG;100MG

A072106 001 May 13, 1988

IVAX SUB TEVA PHARMS 650MG;100MG

A070146 001 Aug 02, 1985

MALLINCKRODT 650MG;100MG

A075738 001 Feb 02, 2001

MIRROR PHARMS 650MG;100MG

A0777821 001 Feb 11, 2008

MUTUAL PHARM 325MG;50MG

A070115 001 Jun 12, 1985

650MG;100MG

A070116 001 Jun 12, 1985

650MG;100MG

A070615 001 Mar 21, 1986

650MG;100MG

A070771 001 Mar 21, 1986

650MG;100MG

A070775 001 Mar 21, 1986

MYLAN 650MG;100MG

A072195 001 Feb 16, 1988

**DISCONTINUED DRUG PRODUCT LIST**

6-9(of 375)

\*\* See List Footnote

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET;ORAL

PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

MYLAN PHARMS INC	650MG;100MG	A070145 001	Jun 12, 1985
SANDOZ	650MG;100MG	A070443 001	Jan 23, 1986
SUPERPHARM	650MG;100MG	A071319 001	Jan 06, 1987
TEVA	650MG;100MG	A070732 001	Jan 03, 1986
	650MG;100MG	A074119 001	Dec 19, 1994
VINTAGE PHARMS	325MG;50MG	A074843 002	Feb 15, 2001
	650MG;100MG	A074843 001	Feb 12, 1997
WATSON LABS	325MG;50MG	A070398 001	Dec 18, 1986
	650MG;100MG	A070399 001	Dec 18, 1986
WATSON LABS FLORIDA	500MG;100MG	A077196 001	Jun 28, 2005
	650MG;100MG	A076609 001	Nov 16, 2004
WOCKHARDT LTD	325MG;50MG	A077677 001	Mar 16, 2007
	650MG;100MG	A077677 002	Mar 16, 2007

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET;ORAL

TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

CSPC OUYI PHARM CO	325MG;37.5MG	A076914 001	Jul 26, 2006
--------------------	--------------	-------------	--------------

ACETAZOLAMIDE

TABLET;ORAL

ACETAZOLAMIDE

ALRA	250MG	A083320 001	
ASCOT	250MG	A087686 001	Oct 20, 1982
SUN PHARM INDUSTRIES	250MG	A089753 001	Jun 22, 1988
VANGARD	250MG	A087654 001	Feb 05, 1982
WATSON LABS	250MG	A084498 002	
	250MG	A088882 001	Oct 22, 1985
DIAMOX			
+ TEVA BRANDED PHARM	125MG **	N008943 001	
+	250MG **	N008943 002	

ACETAZOLAMIDE SODIUM

INJECTABLE;INJECTION

ACETAZOLAMIDE SODIUM

HOSPIRA	EQ 500MG BASE/VIAL	A040108 001	Oct 30, 1995
DIAMOX			

+ TEVA WOMENS	EQ 500MG BASE/VIAL **	N009388 001	Dec 05, 1990
---------------	-----------------------	-------------	--------------

ACETIC ACID, GLACIAL

SOLUTION/DROPS;OTIC

ACETASOL

ACTAVIS MID ATLANTIC	2%	A087146 001	
ACETIC ACID			
KV PHARM	2%	A085493 001	
ORLEX			

WARNER CHILCOTT	2%	A086845 001	
-----------------	----	-------------	--

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS;OTIC

ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE

BAUSCH AND LOMB	2%;0.79%	A040063 001	Feb 25, 1994
BOROFAIR			
PHARMAFAIR	2%;0.79%	A088606 001	Aug 21, 1985
DOMEBORO			

BAYER PHARMS	2%;0.79%	A084476 001	
--------------	----------	-------------	--

ACETIC ACID, GLACIAL; DESONIDE

SOLUTION/DROPS;OTIC

TRIDESILON

BAYER PHARMS	2%;0.05%	N017914 001	
--------------	----------	-------------	--

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

ACETIC ACID W/ HYDROCORTISONE

KV PHARM	2%;1%	A085492 001	
HYDROCORTISONE AND ACETIC ACID			
BAUSCH AND LOMB	2%;1%	A040097 001	Oct 31, 1994

WOCKHARDT	2%;1%	A040168 001	Aug 30, 1996
-----------	-------	-------------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-10(of 375)

\*\* See List Footnote

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS;OTIC

ORLEX HC

WARNER CHILCOTT 2%;1%

A086844 001

ACETIC ACID, GLACIAL; HYDROCORTISONE; NEOMYCIN SULFATE

SUSPENSION/DROPS;OTIC

NEO-CORT-DOME

BAYER PHARMS 2%;1%;EQ 0.35% BASE

N050238 001

ACETOHEXAMIDE

TABLET;ORAL

ACETOHEXAMIDE

ANI PHARMS INC	250MG	A070869 001	Feb 09, 1987
	500MG	A070870 001	Feb 09, 1987
USL PHARMA	250MG	A070753 001	Nov 03, 1986
	500MG	A070754 001	Nov 03, 1986
WATSON LABS TEVA	250MG	A071893 001	Nov 25, 1987
	500MG	A071894 001	Nov 25, 1987
DYMELOR			
LILLY	250MG	N013378 002	
	500MG	N013378 001	

ACETOPHENAZINE MALEATE

TABLET;ORAL

TINDAL

SCHERING 20MG

N012254 002

ACETRIZOATE SODIUM

SOLUTION;INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM 53%

N009008 001

ACETYLCHOLINE CHLORIDE

FOR SOLUTION;OPHTHALMIC

MIOCHOL

NOVARTIS 20MG/VIAL

N016211 001

ACETYLCYSTEINE

SOLUTION;INHALATION, ORAL

ACETYLCYSTEINE

HOSPIRA	10%	A071364 001	May 01, 1989
	20%	A071365 001	May 01, 1989
ROXANE	10%	A072323 001	Apr 30, 1992
	10%	A072621 001	Sep 30, 1992
	20%	A072324 001	Apr 30, 1992
	20%	A072622 001	Sep 30, 1992
MUCOMYST			
+ APOTHECON	10% **	N013601 002	
+	20% **	N013601 001	
MUCOSIL-10			
DEY	10%	A070575 001	Oct 14, 1986
MUCOSIL-20			
DEY	20%	A070576 001	Oct 14, 1986

ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION;INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON 10%;0.05%

N017366 001

ACETYLDIGITOXIN

TABLET;ORAL

ACYLANID

NOVARTIS 0.1MG

N009436 001

ACITRETN

CAPSULE;ORAL

ACITRETN

MYLAN PHARMS INC 17.5MG  
22.5MGA203707 001 Sep 10, 2015  
A203707 002 Sep 10, 2015

**DISCONTINUED DRUG PRODUCT LIST**

6-11(of 375)

\*\* See List Footnote

ACRISORCIN

CREAM;TOPICAL

AKRINOL

SCHERING

2MG/GM

N012470 001

ACYCLOVIR

CAPSULE;ORAL

ACYCLOVIR

ACTAVIS ELIZABETH	200MG	A074906 001	Aug 26, 1997
CHARTWELL MOLECULES	200MG	A074872 001	Apr 22, 1997
HERITAGE PHARMS INC	200MG	A074889 001	Oct 31, 1997
IVAX SUB TEVA PHARMS	200MG	A074674 001	Apr 22, 1997
LEK PHARM	200MG	A074750 001	Apr 22, 1997
MYLAN	200MG	A074727 001	Apr 22, 1997
	200MG	A074977 001	Apr 13, 1998
RANBAXY	200MG	A074975 001	Sep 30, 1998
ROXANE	200MG	A074570 002	Apr 22, 1997
TEVA	200MG	A074828 001	Apr 22, 1997
TEVA PHARMS	200MG	A074914 001	Nov 26, 1997
WATSON LABS	200MG	A075101 001	Apr 15, 1998

TABLET;ORAL

ACYCLOVIR

ACTAVIS ELIZABETH	400MG	A074870 001	Jun 05, 1997
	800MG	A074870 002	Jun 05, 1997
CHARTWELL MOLECULES	400MG	A074834 001	Apr 24, 1997
	800MG	A074834 002	Apr 24, 1997
IVAX SUB TEVA PHARMS	400MG	A074836 001	Apr 22, 1997
	800MG	A074836 002	Apr 22, 1997
LEK PHARM	400MG	A074658 001	Apr 22, 1997
	800MG	A074658 002	Apr 22, 1997
MYLAN	400MG	A074976 001	Apr 13, 1998
	800MG	A074976 002	Apr 13, 1998
SUN PHARM INDS LTD	400MG	A074980 001	Sep 30, 1998
	800MG	A074980 002	Sep 30, 1998
TEVA	200MG **	A074556 001	Apr 22, 1997
TEVA PHARMS	400MG	A075021 001	Mar 18, 1998
	800MG	A075021 002	Mar 18, 1998

ACYCLOVIR SODIUM

INJECTABLE;INJECTION

ACYCLOVIR

ABBVIE	EQ 50MG BASE/ML	A075114 001	Jul 26, 1999
ACYCLOVIR IN SODIUM CHLORIDE	0.9% PRESERVATIVE FREE		
EUROHLTH INTL SARL	EQ 500MG BASE/VIAL	A074885 001	Dec 19, 1997
	EQ 1GM BASE/VIAL	A074885 002	Dec 19, 1997
ACYCLOVIR SODIUM			
APOTHECON	EQ 500MG BASE/VIAL	A074897 001	Feb 27, 1998
	EQ 1GM BASE/VIAL	A074897 002	Feb 27, 1998
ATHENEX INC	EQ 500MG BASE/VIAL	A074596 002	Apr 22, 1997
	EQ 1GM BASE/VIAL	A074596 001	Apr 22, 1997
EUROHLTH INTL SARL	EQ 500MG BASE/VIAL	A074913 001	Oct 15, 1997
	EQ 1GM BASE/VIAL	A074913 002	Oct 15, 1997
HOSPIRA	EQ 25MG BASE/ML	A074720 001	Apr 22, 1997
	EQ 50MG BASE/ML	A075065 001	Feb 25, 1999
	EQ 500MG BASE/VIAL	A074663 001	Apr 22, 1997
	EQ 500MG BASE/VIAL	A074758 001	Apr 22, 1997
	EQ 1GM BASE/VIAL	A074663 002	Apr 22, 1997
	EQ 1GM BASE/VIAL	A074758 002	Apr 22, 1997
MYLAN LABS LTD	EQ 500MG BASE/VIAL	A203927 001	Mar 29, 2017
	EQ 1GM BASE/VIAL	A203927 002	Mar 29, 2017
TEVA PARENTERAL	EQ 50MG BASE/ML	A075627 001	Mar 28, 2001
	EQ 500MG BASE/VIAL	A074969 001	Aug 26, 1997
	EQ 1GM BASE/VIAL	A074969 002	Aug 26, 1997
ZOVIRAX			
+ GLAXOSMITHKLINE	EQ 250MG BASE/VIAL **	N018603 003	Aug 30, 1983
+	EQ 500MG BASE/VIAL **	N018603 001	Oct 22, 1982
+	EQ 1GM BASE/VIAL **	N018603 002	Jun 29, 1989

**DISCONTINUED DRUG PRODUCT LIST**

6-12(of 375)

\*\* See List Footnote

ADAPALENE

SOLUTION;TOPICAL

DIFERIN

+ GALDERMA LABS LP 0.1% \*\*

N020338 001 May 31, 1996

ADENOSINE

INJECTABLE;INJECTION

ADENOSINE

TEVA PHARMS USA 3MG/ML

3MG/ML

WEST-WARD PHARMS INT 3MG/ML

WOCKHARDT 3MG/ML

A076564 001 Jun 16, 2004

A078676 001 Jul 31, 2008

A076501 001 Jun 16, 2004

A090220 001 Jul 20, 2009

SOLUTION;IV (INFUSION)

ADENOSCAN

+ ASTELLAS 60MG/20ML (3MG/ML) \*\*

N020059 001 May 18, 1995

+ 90MG/30ML (3MG/ML) \*\*

N020059 002 May 18, 1995

ALATROFLOXACIN MESYLATE

INJECTABLE;INJECTION

TROVAN PRESERVATIVE FREE

PFIZER EQ 200MG BASE/VIAL

N020760 001 Dec 18, 1997

EQ 300MG BASE/VIAL

N020760 002 Dec 18, 1997

ALBENDAZOLE

TABLET, CHEWABLE;ORAL

ALBENZA

AMEDRA PHARMS LLC 200MG

N207844 001 Jun 11, 2015

ALBUMIN CHROMATED CR-51 SERUM

INJECTABLE;INJECTION

CHROMALBIN

ISO TEX 100uCi/VIAL

N017835 001

250uCi/VIAL

N017835 002

500uCi/VIAL

N017835 003

ALBUMIN IODINATED I-125 SERUM

INJECTABLE;INJECTION

RADIO-IODINATED (I 125) SERUM ALBUMIN (HUMAN)

BAYER PHARMS 2.5uCi/AMP

N017846 001

RADIOIODINATED SERUM ALBUMIN (HUMAN) IHSA I 125

MALLINCKRODT 6.67uCi/ML

N017844 003

10uCi/ML

N017844 001

100uCi/ML

N017844 002

ALBUMIN IODINATED I-131 SERUM

INJECTABLE;INJECTION

MEGATOPE

ISO TEX 2mCi/VIAL

N017837 003

5uCi/AMP

N017837 004

20uCi/AMP

N017837 005

ALBUTEROL

AEROSOL, METERED;INHALATION

ALBUTEROL

ARMSTRONG PHARMS 0.09MG/INH

A072273 001 Aug 14, 1996

GENPHARM 0.09MG/INH

A073045 001 Aug 19, 1997

IVAX SUB TEVA PHARMS 0.09MG/INH

A073272 001 Dec 28, 1995

PLIVA 0.09MG/INH

A074072 001 Aug 01, 1996

PROVENTIL

0.09MG/INH

N017559 001

SCHERING

0.09MG/INH

VENTOLIN

0.09MG/INH

N018473 001

GLAXOSMITHKLINE 0.09MG/INH

ALBUTEROL SULFATE

CAPSULE;INHALATION

VENTOLIN ROTACAPS

EQ 0.2MG BASE

N019489 001 May 04, 1988

SOLUTION;INHALATION

ALBUTEROL SULFATE

ACTAVIS MID ATLANTIC EQ 0.083% BASE

A073533 001 Sep 26, 1995

APOTEX INC EQ 0.021% BASE

A078623 001 Apr 05, 2010

EQ 0.042% BASE

A078623 002 Apr 05, 2010

EQ 0.083% BASE

A075717 001 Feb 02, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-13(of 375)

\*\* See List Footnote

ALBUTEROL SULFATESOLUTION; INHALATION  
ALBUTEROL SULFATE

BAUSCH AND LOMB	EQ 0.5% BASE	A076391 001	Apr 01, 2003
COPLEY PHARM	EQ 0.083% BASE	A075358 001	Mar 29, 2000
	EQ 0.083% BASE	A073495 001	May 28, 1993
	EQ 0.5% BASE	A073307 001	Nov 27, 1991
HI TECH PHARMA	EQ 0.083% BASE	A075063 001	Feb 09, 1999
LANDELA PHARM	EQ 0.083% BASE	A077569 001	Apr 04, 2006
MYLAN SPECLT	EQ 0.083% BASE **	A072652 001	Feb 21, 1992
ROXANE	EQ 0.083% BASE	A075129 001	Feb 13, 2001
TEVA PHARMS	EQ 0.083% BASE	A075343 001	Nov 09, 1999
WATSON LABS INC	EQ 0.083% BASE	A076370 001	Nov 24, 2003
WOCKHARDT EU OPERATN	EQ 0.083% BASE	A075394 001	Nov 22, 1999
PROVENTIL			
SCHERING	EQ 0.083% BASE **	N019243 002	Jan 14, 1987
	EQ 0.5% BASE **	N019243 001	Jan 14, 1987
VENTOLIN			
+ GLAXOSMITHKLINE	EQ 0.083% BASE **	N019773 001	Apr 23, 1992
	EQ 0.5% BASE **	N019269 002	Jan 16, 1987
SYRUP; ORAL			
ALBUTEROL SULFATE			
ACTAVIS MID ATLANTIC	EQ 2MG BASE/5ML	A075262 001	Mar 30, 1999
MOVA	EQ 2MG BASE/5ML	A074302 001	Sep 30, 1994
WATSON LABS	EQ 2MG BASE/5ML	A073165 001	Apr 29, 1993
PROVENTIL			
+ SCHERING	EQ 2MG BASE/5ML **	N018062 001	Jan 19, 1983
VENTOLIN			
GLAXOSMITHKLINE	EQ 2MG BASE/5ML	N019621 001	Jun 10, 1987
TABLET; ORAL			
ALBUTEROL SULFATE			
AM THERAP	EQ 2MG BASE	A072449 001	Dec 05, 1989
	EQ 4MG BASE	A072450 001	Dec 05, 1989
COPLEY PHARM	EQ 2MG BASE	A072966 001	Nov 22, 1991
	EQ 4MG BASE	A072967 001	Nov 22, 1991
DAVA PHARMS INC	EQ 2MG BASE	A072860 002	Dec 20, 1989
	EQ 4MG BASE	A072860 001	Dec 20, 1989
FOSUN PHARMA	EQ 2MG BASE	A072151 001	Dec 05, 1989
	EQ 4MG BASE	A072152 001	Dec 05, 1989
PLIVA	EQ 2MG BASE	A072316 001	Dec 05, 1989
	EQ 4MG BASE	A072317 001	Dec 05, 1989
TEVA	EQ 2MG BASE	A072619 001	Dec 05, 1989
	EQ 2MG BASE	A072779 001	Jun 25, 1993
	EQ 4MG BASE	A072938 001	Mar 30, 1990
	EQ 4MG BASE	A072620 001	Dec 05, 1989
	EQ 4MG BASE	A072780 001	Jun 25, 1993
	EQ 4MG BASE	A072939 001	Mar 30, 1990
UCB INC	EQ 2MG BASE	A073120 001	Sep 29, 1992
	EQ 4MG BASE	A073121 001	Sep 29, 1992
WARNER CHILCOTT	EQ 2MG BASE	A072817 001	Jan 09, 1990
	EQ 4MG BASE	A072818 001	Jan 09, 1990
WATSON LABS	EQ 2MG BASE	A072629 001	Jan 31, 1991
	EQ 2MG BASE	A072764 001	Aug 28, 1991
	EQ 4MG BASE	A072630 001	Jan 31, 1991
	EQ 4MG BASE	A072765 001	Aug 28, 1991
PROVENTIL			
+ SCHERING	EQ 2MG BASE **	N017853 001	May 07, 1982
+	EQ 4MG BASE **	N017853 002	May 07, 1982
VENTOLIN			
GLAXOSMITHKLINE	EQ 2MG BASE	N019112 001	Jul 10, 1986
	EQ 4MG BASE	N019112 002	Jul 10, 1986
TABLET, EXTENDED RELEASE; ORAL			
PROVENTIL			
SCHERING	EQ 4MG BASE	N019383 001	Jul 13, 1987
VOLMAX			
MURO	EQ 4MG BASE	N019604 002	Dec 23, 1992
	EQ 8MG BASE	N019604 001	Dec 23, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-14(of 375)

\*\* See List Footnote

ALBUTEROL SULFATE; IPRATROPIUM BROMIDEAEROSOL, METERED; INHALATION  
COMBIVENT

BOEHRINGER INGELHEIM	EQ 0.09MG BASE/INH;0.018MG/INH	N020291 001 Oct 24, 1996
SOLUTION; INHALATION		
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE		
APOTEX INC	EQ 0.083% BASE;0.017%	A077117 001 Dec 31, 2007
SANDOZ INC	EQ 0.083% BASE;0.017%	A076867 001 Dec 21, 2006
DUONEB		
+ MYLAN SPECIALITY LP	EQ 0.083% BASE;0.017% **	N020950 001 Mar 21, 2001

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL		
ACLOVATE		
+ FOUGERA PHARMS	0.05% **	N018707 001 Dec 14, 1982
OINTMENT; TOPICAL		
ACLOVATE		
+ FOUGERA PHARMS	0.05% **	N018702 001 Dec 14, 1982

ALCOHOL

INJECTABLE; INJECTION		
ALCOHOL 5% IN DEXTROSE 5%		
MILES	5ML/100ML	A083483 001

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION		
ALCOHOL 10% AND DEXTROSE 5%		
B BRAUN	10ML/100ML;5GM/100ML	N004589 006
ALCOHOL 5% AND DEXTROSE 5%		
B BRAUN	5ML/100ML;5GM/100ML	N004589 004
ALCOHOL 5% IN D5-W		
HOSPIRA	5ML/100ML;5GM/100ML	A083263 001
ALCOHOL 5% IN DEXTROSE 5% IN WATER		
BAXTER HLTHCARE	5ML/100ML;5GM/100ML	A083256 001

ALENDRONATE SODIUM

SOLUTION; ORAL		
FOSAMAX		
+ MERCK	EQ 70MG BASE/75ML **	N021575 001 Sep 17, 2003
TABLET; ORAL		
ALENDRONATE SODIUM		
MYLAN	EQ 35MG BASE	A078638 001 Aug 04, 2008
	EQ 70MG BASE	A078638 002 Aug 04, 2008
TEVA PHARMS	EQ 35MG BASE	A076184 002 Aug 04, 2008
	EQ 70MG BASE	A076184 001 Feb 06, 2008
UPSHER-SMITH LABS	EQ 5MG BASE	A075871 001 Apr 22, 2009
	EQ 10MG BASE	A075871 002 Apr 22, 2009
	EQ 35MG BASE	A075871 004 Apr 22, 2009
	EQ 40MG BASE	A075871 003 Apr 22, 2009
	EQ 70MG BASE	A075871 005 Apr 22, 2009
FOSAMAX		
+ MERCK AND CO INC	EQ 5MG BASE **	N020560 003 Apr 25, 1997
+	EQ 10MG BASE **	N020560 001 Sep 29, 1995
+	EQ 35MG BASE **	N020560 004 Oct 20, 2000
+	EQ 40MG BASE **	N020560 002 Sep 29, 1995

ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL		
ALFUZOSIN HYDROCHLORIDE		
WOCKHARDT LTD	10MG	A090221 001 Aug 10, 2012

ALGLUCERASE

INJECTABLE; INJECTION		
CEREDASE		
GENZYME	10 UNITS/ML	N020057 004 May 08, 1992
	80 UNITS/ML	N020057 003 Apr 05, 1991

**DISCONTINUED DRUG PRODUCT LIST**

6-15(of 375)

\*\* See List Footnote

ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATETABLET;ORAL  
TEKAMLO

NOVARTIS

EQ 150MG BASE;EQ 5MG BASE  
EQ 150MG BASE;EQ 10MG BASE  
EQ 300MG BASE;EQ 5MG BASE  
EQ 300MG BASE;EQ 10MG BASEN022545 001 Aug 26, 2010  
N022545 002 Aug 26, 2010  
N022545 003 Aug 26, 2010  
N022545 004 Aug 26, 2010ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDETABLET;ORAL  
AMTURNIDE

NOVARTIS

EQ 150MG BASE;EQ 5MG BASE;12.5MG  
EQ 300MG BASE;EQ 5MG BASE;12.5MG  
EQ 300MG BASE;EQ 5MG BASE;25MG  
EQ 300MG BASE;EQ 10MG BASE;12.5MG  
EQ 300MG BASE;EQ 10MG BASE;25MGN200045 001 Dec 21, 2010  
N200045 002 Dec 21, 2010  
N200045 003 Dec 21, 2010  
N200045 004 Dec 21, 2010  
N200045 005 Dec 21, 2010ALISKIREN HEMIFUMARATE; VALSARTANTABLET;ORAL  
VALTURNA

NOVARTIS

EQ 150MG BASE;160MG  
EQ 300MG BASE;320MGN022217 001 Sep 16, 2009  
N022217 002 Sep 16, 2009ALKAVERVIRTABLET;ORAL  
VERILOID

3M

2MG

N007336 002

3MG

N007336 003

ALLOPURINOLTABLET;ORAL  
ALLOPURINOL

MUTUAL PHARM

100MG

A070466 001 Dec 24, 1985

300MG

A070467 001 Dec 24, 1985

PURACAP PHARM

100MG

A070150 001 Dec 10, 1985

300MG

A070147 001 Dec 10, 1985

PUREPAC PHARM

100MG

A070579 001 Apr 14, 1986

300MG

A070580 001 Apr 14, 1986

SANDOZ

100MG

A070268 001 Dec 31, 1985

300MG

A070269 001 Dec 31, 1985

SUPERPHARM

100MG

A070950 001 Nov 30, 1988

300MG

A070951 001 Nov 30, 1988

WATSON LABS

100MG

N018241 001 Nov 16, 1984

100MG

N018785 001 Sep 28, 1984

300MG

N018241 002 Nov 16, 1984

300MG

N018785 002 Sep 28, 1984

LOPURIN

ABBOTT

100MG

N018297 001

300MG

N018297 002

ALPRAZOLAMSOLUTION;ORAL  
ALPRAZOLAM

ROXANE

0.5MG/5ML

A074314 001 Oct 31, 1993

TABLET;ORAL  
ALPRAZOLAM

ANI PHARMS INC

0.25MG

A074085 001 Feb 16, 1994

0.5MG

A074085 002 Feb 16, 1994

1MG

A074085 003 Feb 16, 1994

2MG

A074085 004 Feb 26, 1996

IVAX SUB TEVA PHARMS

0.25MG

A074294 001 Jul 29, 1994

0.5MG

A074294 002 Jul 29, 1994

1MG

A074294 003 Jul 29, 1994

2MG

A074294 004 Jul 29, 1994

MYLAN PHARMS INC

0.25MG

A074046 001 Oct 19, 1993

0.5MG

A074046 002 Oct 19, 1993

1MG

A074046 003 Oct 19, 1993

2MG

A074046 004 May 07, 1997

ROXANE

0.25MG

A074199 001 Oct 19, 1993

0.5MG

A074199 002 Oct 19, 1993

1MG

A074199 003 Oct 19, 1993

WATSON LABS

0.25MG

A074456 001 Aug 31, 1995

**DISCONTINUED DRUG PRODUCT LIST**

6-16(of 375)

\*\* See List Footnote

**ALPRAZOLAM**TABLET;ORAL  
ALPRAZOLAM

0.25MG	A074479 001	Jan 21, 1997
0.5MG	A074456 002	Aug 31, 1995
0.5MG	A074479 002	Jan 21, 1997
1MG	A074456 003	Aug 31, 1995
1MG	A074479 003	Jan 21, 1997

TABLET, EXTENDED RELEASE;ORAL

ALPRAZOLAM

ACTAVIS LABS FL INC	0.5MG	A077198 001	May 13, 2010
	1MG	A077198 002	May 13, 2010
	2MG	A077198 003	May 13, 2010
	3MG	A077198 004	May 13, 2010
ANI PHARMS INC	0.5MG	A077979 001	Feb 28, 2007
	1MG	A077979 002	Feb 28, 2007
	2MG	A077979 003	Feb 28, 2007
	3MG	A077979 004	Feb 28, 2007
IMPAX LABS	0.5MG	A077968 004	May 24, 2007
	1MG	A077968 003	May 24, 2007
	2MG	A077968 002	May 24, 2007
	3MG	A077968 001	May 24, 2007
IMPAX LABS INC	0.5MG	A077996 001	Jan 31, 2007
	1MG	A077996 002	Jan 31, 2007
	2MG	A077996 003	Jan 31, 2007
	3MG	A077996 004	Jan 31, 2007
MYLAN	0.5MG	A077391 002	Jan 26, 2006
	1MG	A077391 003	Jan 26, 2006
	2MG	A077391 004	Jan 26, 2006
	3MG	A077391 001	Jan 26, 2006
SANDOZ INC	0.5MG	A077777 001	Jun 30, 2006
	1MG	A077777 002	Jun 30, 2006
	2MG	A077777 003	Jun 30, 2006
	3MG	A077777 004	Jun 30, 2006
VINTAGE PHARMS	0.5MG	A078442 001	Oct 15, 2007
	1MG	A078442 002	Oct 15, 2007
	2MG	A078442 003	Oct 15, 2007
	3MG	A078442 004	Oct 15, 2007

TABLET, ORALLY DISINTEGRATING;ORAL

NIRAVAM

+ UCB INC	0.25MG **	N021726 001	Jan 19, 2005
+	0.5MG **	N021726 002	Jan 19, 2005
+	1MG **	N021726 003	Jan 19, 2005
+	2MG **	N021726 004	Jan 19, 2005

**ALPROSTADIL**

INJECTABLE;INJECTION

CAVERJECT

PFIZER	0.005MG/ML	N020755 001	Oct 31, 1997
	0.01MG/ML	N020755 002	Oct 01, 1997
	0.02MG/ML	N020755 003	Oct 01, 1997
EDEX			
AUXILIUM PHARMS LLC	0.005MG/VIAL	N020649 001	Jun 12, 1997

**ALSEROXYLON**

TABLET;ORAL

RAUTENSIN

NOVARTIS	2MG	N009215 001
RAUWILOID		

3M

2MG	N008867 001
-----	-------------

**ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE**

TABLET, CHEWABLE;ORAL

ALUMINUM HYDROXIDE AND MAGNESIUM TRISILICATE

PENNEX	80MG;20MG	A089449 001	Nov 27, 1987
FOAMCOAT			
GUARDIAN DRUG	80MG;20MG	A071793 001	Sep 04, 1987

FOAMICON

NOVARTIS

80MG;20MG	A072687 001	Jun 28, 1989
-----------	-------------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-17(of 375)

\*\* See List Footnote

AMANTADINE HYDROCHLORIDE

CAPSULE;ORAL

AMANTADINE HYDROCHLORIDE

ACTAVIS ELIZABETH 100MG A077659 001 Feb 23, 2006

WATSON LABS 100MG A071382 001 Jan 21, 1987

SYMADINE

SOLVAY 100MG A071000 001 Sep 04, 1986

SYMMETREL

+ ENDO PHARMS 100MG \*\* N016020 001

SYRUP;ORAL

AMANTADINE HYDROCHLORIDE

G AND W LABS INC 50MG/5ML A072655 001 Oct 30, 1990

SILARX 50MG/5ML A076352 001 Sep 10, 2004

TEVA PHARMS 50MG/5ML A073115 001 Aug 23, 1991

VINTAGE 50MG/5ML A077992 001 Dec 12, 2006

SYMMETREL

+ ENDO PHARMS 50MG/5ML \*\* N016023 002

TABLET;ORAL

SYMMETREL + ENDO PHARMS 100MG \*\* N018101 001

AMBENONIUM CHLORIDE

TABLET;ORAL

MYTELASE SANOFI AVENTIS US 10MG N010155 002

AMCINONIDE

CREAM;TOPICAL

CYCLOCORT + ASTELLAS 0.025% N018116 001

+ 0.1% N018116 002

LOTION;TOPICAL

CYCLOCORT + ASTELLAS 0.1% N019729 001 Jun 13, 1988

OINTMENT;TOPICAL

CYCLOCORT + ASTELLAS 0.1% N018498 001

AMDINOCILLIN

INJECTABLE;INJECTION

COACTIN ROCHE 250MG/VIAL N050565 001 Dec 21, 1984

500MG/VIAL N050565 002 Dec 21, 1984

1GM/VIAL N050565 003 Dec 21, 1984

AMIFOSTINE

INJECTABLE;INJECTION

ETHYOL CLINIGEN HLTHCARE 375MG/VIAL N020221 002 Sep 10, 1999

AMIKACIN SULFATE

INJECTABLE;INJECTION

AMIKACIN SULFATE ABBOTT EQ 250MG BASE/ML A063265 001 Nov 30, 1994

EQ 250MG BASE/ML A063266 001 Oct 31, 1994

HOSPIRA EQ 50MG BASE/ML A063263 001 Nov 30, 1994

EQ 50MG BASE/ML A063350 001 Jul 30, 1993

EQ 62.5MG BASE/ML A063283 001 Oct 31, 1994

EQ 250MG BASE/ML A063264 001 Nov 30, 1994

EQ 250MG BASE/ML A063350 002 Jul 30, 1993

EQ 250MG BASE/ML A064098 001 Jun 26, 1995

EQ 250MG BASE/ML A064099 001 Jun 20, 1995

IGI LABS INC EQ 50MG BASE/ML A063167 001 Dec 14, 1995

EQ 250MG BASE/ML A063169 001 Dec 14, 1995

TEVA PHARMS USA EQ 50MG BASE/ML A064045 001 Sep 28, 1993

WEST-WARD PHARMS INT EQ 50MG BASE/ML A063274 001 May 18, 1992

EQ 250MG BASE/ML A063275 001 May 18, 1992

AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER HOSPIRA EQ 500MG BASE/100ML A064146 001 Apr 02, 1997

AMIKIN APOTHECON EQ 50MG BASE/ML A062311 001

EQ 50MG BASE/ML A062562 001 Sep 20, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-18(of 375)

\*\* See List Footnote

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKIN

+

EQ 50MG BASE/ML \*\*

N050495 001

EQ 250MG BASE/ML

A062311 002

EQ 250MG BASE/ML

A062562 002 Sep 20, 1984

+

EQ 250MG BASE/ML \*\*

N050495 002

AMIKIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

APOTHECON

EQ 5MG BASE/ML

N050618 002 Nov 30, 1987

EQ 10MG BASE/ML

N050618 001 Nov 30, 1987

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

FOSUN PHARMA

EQ 5MG ANHYDROUS; 50MG

A073357 001 Nov 27, 1991

TEVA

EQ 5MG ANHYDROUS; 50MG

A070795 001 Apr 17, 1988

WATSON LABS

EQ 5MG ANHYDROUS; 50MG

A073334 001 Jul 19, 1991

HYDRO-RIDE

PAR PHARM

EQ 5MG ANHYDROUS; 50MG

A070347 001 Dec 25, 1990

MODURETIC 5-50

+ MERCK

EQ 5MG ANHYDROUS; 50MG \*\*

N018201 001

AMINO ACIDS

INJECTABLE; INJECTION

AMINESS 5.2% ESSENTIAL AMINO ACIDS W/ HISTADINE

HOSPIRA

5.2% (5.2GM/100ML)

N018901 001 Apr 06, 1984

AMINOSYN 3.5% IN PLASTIC CONTAINER

ABBOTT

3.5% (3.5GM/100ML)

N018804 001 May 15, 1984

3.5% (3.5GM/100ML)

N018875 001 Aug 08, 1984

AMINOSYN II 3.5%

ICU MEDICAL INC

3.5% (3.5GM/100ML)

N019438 001 Apr 03, 1986

AMINOSYN II 3.5% IN PLASTIC CONTAINER

ABBOTT

3.5% (3.5GM/100ML)

N019491 001 Oct 10, 1986

AMINOSYN II 5%

ICU MEDICAL INC

5% (5GM/100ML)

N019438 002 Apr 03, 1986

AMINOSYN-HBC 7% IN PLASTIC CONTAINER

ABBOTT

7% (7GM/100ML)

N019400 001 Jul 23, 1986

BRANCHAMIN 4%

BAXTER HLTHCARE

4% (4GM/100ML)

N018678 001 Sep 28, 1984

BRANCHAMIN 4% IN PLASTIC CONTAINER

BAXTER HLTHCARE

4% (4GM/100ML)

N018684 001 Sep 28, 1984

FREAMINE 8.5%

B BRAUN

8.5% (8.5GM/100ML)

N016822 001

FREAMINE II 8.5%

B BRAUN

8.5% (8.5GM/100ML)

N016822 002

HEPATASOL 8%

BAXTER HLTHCARE

8% (8GM/100ML)

A020360 001 Apr 04, 1996

NEOPHAM 6.4%

HOSPIRA

6.4% (6.4GM/100ML)

N018792 001 Jan 17, 1984

NOVAMINE 11.4%

HOSPIRA INC

11.4% (11.4GM/100ML)

N017957 003 Aug 09, 1982

NOVAMINE 15%

HOSPIRA INC

15% (75GM/500ML)

N017957 004 Nov 28, 1986

NOVAMINE 15% SULFITE FREE IN PLASTIC CONTAINER

BAXTER HLTHCARE

15% (15GM/100ML)

N020107 001 Feb 05, 1993

NOVAMINE 8.5%

HOSPIRA INC

8.5% (8.5GM/100ML)

N017957 002 Aug 09, 1982

RENAMIN W/O ELECTROLYTES

BAXTER HLTHCARE

6.5% (6.5GM/100ML)

N017493 007 Oct 15, 1982

TRAVASOL 10% W/O ELECTROLYTES

BAXTER HLTHCARE

10% (10GM/100ML)

N017493 006

TRAVASOL 5.5% W/O ELECTROLYTES

BAXTER HLTHCARE

5.5% (5.5GM/100ML)

N017493 004

TRAVASOL 8.5% W/O ELECTROLYTES

BAXTER HLTHCARE

8.5% (8.5GM/100ML)

N017493 005

**DISCONTINUED DRUG PRODUCT LIST**

6-19(of 375)

\*\* See List Footnote

**AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE**

## INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	3.5%;36.8MG/100ML;25GM/100ML;51MG/100ML ;22.4MG/100ML;261MG/100ML;205MG/100ML	N019714	001	Sep 12, 1988
HOSPIRA INC	3.5%;36.8MG/100ML;25GM/100ML;51MG/100ML ;22.4MG/100ML;261MG/100ML;205MG/100ML	N019683	001	Nov 07, 1988
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	4.25%;36.8MG/100ML;20GM/100ML;51MG/100M L;22.4MG/100ML;261MG/100ML;205MG/100ML	N019714	002	Sep 12, 1988
HOSPIRA INC	4.25%;36.8MG/100ML;20GM/100ML;51MG/100M L;22.4MG/100ML;261MG/100ML;205MG/100ML	N019683	002	Nov 07, 1988
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	4.25%;36.8MG/100ML;25GM/100ML;51MG/100M L;22.4MG/100ML;261MG/100ML;205MG/100ML	N019714	004	Sep 12, 1988
HOSPIRA INC	4.25%;36.8MG/100ML;25GM/100ML;51MG/100M L;22.4MG/100ML;261MG/100ML;205MG/100ML	N019683	003	Nov 07, 1988
AMINOSYN II 5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER				
ABBOTT	5%;36.8MG/100ML;25GM/100ML;51MG/100ML; 2.4MG/100ML;261MG/100ML;205MG/100ML	N019714	003	Sep 12, 1988
HOSPIRA INC	5%;36.8MG/100ML;25GM/100ML;51MG/100ML; 2.4MG/100ML;261MG/100ML;205MG/100ML	N019683	004	Nov 07, 1988

**AMINO ACIDS; DEXTROSE**

## INJECTABLE; INJECTION

AMINOSYN 3.5% W/ DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	3.5%;25GM/100ML	N019118	001	Oct 11, 1984
AMINOSYN 3.5% W/ DEXTROSE 5% IN PLASTIC CONTAINER				
ABBOTT	3.5%;5GM/100ML	N019120	001	Oct 11, 1984
AMINOSYN 4.25% W/ DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	4.25%;25GM/100ML	N019119	001	Oct 11, 1984
AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	3.5%;25GM/100ML	N019505	002	Nov 07, 1986
	3.5%;25GM/100ML	N019713	006	Sep 09, 1988
HOSPIRA	3.5%;25GM/100ML	N019681	001	Nov 01, 1988
AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER				
ABBOTT	3.5%;5GM/100ML	N019506	001	Nov 07, 1986
	3.5%;5GM/100ML	N019713	002	Sep 09, 1988
HOSPIRA	3.5%;5GM/100ML	N019681	002	Nov 01, 1988
AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER				
ABBOTT	4.25%;10GM/100ML	N019713	001	Sep 09, 1988
HOSPIRA	4.25%;10GM/100ML	N019681	004	Nov 01, 1988
AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER				
ABBOTT	4.25%;20GM/100ML	N019713	004	Sep 09, 1988
HOSPIRA	4.25%;20GM/100ML	N019681	005	Nov 01, 1988
AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	4.25%;25GM/100ML	N019504	002	Nov 07, 1986
	4.25%;25GM/100ML	N019713	005	Sep 09, 1988
HOSPIRA	4.25%;25GM/100ML	N019681	003	Nov 01, 1988
AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER				
ABBOTT	5%;25GM/100ML	N019565	001	Dec 17, 1986
	5%;25GM/100ML	N019713	003	Sep 09, 1988
HOSPIRA	5%;25GM/100ML	N019681	006	Nov 01, 1988
TRAVASOL 2.75% IN DEXTROSE 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;10GM/100ML	N019520	002	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 15% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;15GM/100ML	N019520	003	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 20% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;20GM/100ML	N019520	004	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 25% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;25GM/100ML	N019520	005	Sep 23, 1988
TRAVASOL 2.75% IN DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2.75%;5GM/100ML	N019520	001	Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4.25%;10GM/100ML	N019520	007	Sep 23, 1988

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-20(of 375)

\*\* See List Footnote

AMINO ACIDS; DEXTROSE

## INJECTABLE; INJECTION

TRAVASOL 4.25% IN DEXTROSE 15% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;15GM/100ML	N019520 008 Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;20GM/100ML	N019520 009 Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;25GM/100ML	N019520 010 Sep 23, 1988
TRAVASOL 4.25% IN DEXTROSE 5% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;5GM/100ML	N019520 006 Sep 23, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

## INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECT AND ADJUSTED PHOSPHATE IN DEXTROSE 10% IN PLASTIC CONTAINER ABBOTT	4.25%;10GM/100ML;51MG/100ML;176.5MG/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML	N019712 002 Sep 08, 1988
HOSPIRA INC	4.25%;10GM/100ML;51MG/100ML;176.5MG/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML	N019682 003 Nov 01, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

## INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER ABBOTT	3.5%;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML	N019564 002 Dec 16, 1986
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER ABBOTT	4.25%;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML	N019564 004 Dec 16, 1986

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

## INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER ABBOTT	3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N019564 001 Dec 16, 1986
HOSPIRA INC	3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N019682 001 Nov 01, 1988
AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER ABBOTT	4.25%;10GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N019564 003 Dec 16, 1986
HOSPIRA INC	4.25%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N019682 002 Nov 01, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

## INJECTABLE; INJECTION

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER BAXTER HLTHCARE	2.75%;10GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML	N020147 002 Oct 23, 1995
TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER BAXTER HLTHCARE	2.75%;15GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML	N020147 003 Oct 23, 1995
TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER BAXTER HLTHCARE	2.75%;20GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML	N020147 004 Oct 23, 1995
TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER BAXTER HLTHCARE	2.75%;25GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML	N020147 005 Oct 23, 1995
TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER BAXTER HLTHCARE	2.75%;5GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML	N020147 001 Oct 23, 1995
TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;10GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML	N020147 007 Oct 23, 1995
TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;15GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML	N020147 008 Oct 23, 1995
TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER BAXTER HLTHCARE	4.25%;20GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML	N020147 009 Oct 23, 1995

**DISCONTINUED DRUG PRODUCT LIST**

6-21(of 375)

\*\* See List Footnote

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 4.25%;25GM/100ML;51MG/100ML;261MG/100ML N020147 010 Oct 23, 1995  
     ;297MG/100ML;77MG/100ML  
 TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 4.25%;5GM/100ML;51MG/100ML;261MG/100ML; N020147 006 Oct 23, 1995  
     297MG/100ML;77MG/100ML

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M IN PLASTIC CONTAINER  
 ABBOTT 3.5%;21MG/100ML;40MG/100ML;128MG/100ML; N018804 002 May 15, 1984  
     234MG/100ML  
 3.5%;21MG/100ML;40MG/100ML;128MG/100ML; N018875 002 Aug 08, 1984  
     234MG/100ML

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M  
 ICU MEDICAL INC 3.5%;21MG/100ML;128MG/100ML;234MG/100ML N017789 005

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN PLASTIC CONTAINER  
 ABBOTT 3.5%;32MG/100ML;128MG/100ML;222MG/100ML N019493 001 Oct 16, 1986  
     ;49MG/100ML

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

VEINAMINE 8%  
 HOSPIRA INC 8%;61MG/100ML;211MG/100ML;56MG/100ML;38 N017957 001  
     8MG/100ML

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 7% W/ ELECTROLYTES  
 ICU MEDICAL INC 7%;102MG/100ML;45MG/100ML;522MG/100ML;4 N019437 006 Apr 03, 1986  
     10MG/100ML

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M  
 ICU MEDICAL INC 3.5%;30MG/100ML;97MG/100ML;120MG/100ML; N019437 007 Apr 03, 1986  
     49MG/100ML

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 3.5%;51MG/100ML;131MG/100ML;218MG/100ML N020177 001 Oct 23, 1995  
     ;35MG/100ML

TRAVASOL 3.5% W/ ELECTROLYTES  
 BAXTER HLTHCARE 3.5%;51MG/100ML;131MG/100ML;218MG/100ML N017493 003  
     ;35MG/100ML

TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 5.5%;102MG/100ML;522MG/100ML;431MG/100M N020173 001 Oct 27, 1995  
     L;224MG/100ML

TRAVASOL 5.5% W/ ELECTROLYTES  
 BAXTER HLTHCARE 5.5%;102MG/100ML;522MG/100ML;431MG/100M N017493 001  
     L;224MG/100ML

TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 8.5%;102MG/100ML;522MG/100ML;594MG/100M N020173 002 Oct 27, 1995  
     L;154MG/100ML

TRAVASOL 8.5% W/ ELECTROLYTES  
 BAXTER HLTHCARE 8.5%;102MG/100ML;522MG/100ML;594MG/100M N017493 002  
     L;154MG/100ML

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMICAR  
 + CLOVER PHARMS 250MG/ML \*\* N015229 002  
 AMINOCAPROIC ACID  
 ABRAXIS PHARM 250MG/ML A070522 001 Jun 17, 1986  
 BAXTER HLTHCARE 250MG/ML N018590 001 Oct 29, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-22(of 375)

\*\* See List Footnote

AMINOCAPROIC ACID

INJECTABLE; INJECTION  
 AMINOCAPROIC ACID  
 HOSPIRA 250MG/ML A070888 001 Jun 16, 1988

AMINOGLUTETHIMIDE

TABLET; ORAL  
 CYTADREN  
 NOVARTIS 250MG N018202 001

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION  
 AMINOHIPPURATE SODIUM  
 MERCK 20% N005619 001

AMINOPHYLLINE

ENEMA; RECTAL  
 SOMOPHYLLIN  
 FISONS 300MG/5ML N018232 001 Apr 02, 1982

INJECTABLE; INJECTION  
 AMINOPHYLLIN  
 GD SEARLE LLC 25MG/ML A087243 001 May 24, 1982  
 25MG/ML A087621 001 May 24, 1982

AMINOPHYLLINE  
 ABRAXIS PHARM 25MG/ML A084568 001  
 25MG/ML A087200 001  
 25MG/ML A087250 001 Jan 06, 1982  
 25MG/ML A087886 001 Aug 30, 1983  
 25MG/ML A088407 001 Jan 25, 1984  
 ELKINS SINK 25MG/ML A087239 001  
 HOSPIRA 25MG/ML A087601 001 Jul 23, 1982  
 INTL MEDICATION 25MG/ML A087209 001 Feb 01, 1982  
 25MG/ML A087867 001 Nov 10, 1983  
 25MG/ML A087868 001 Nov 10, 1983  
 KING PHARMS 25MG/ML A086606 001  
 LUITPOLD 25MG/ML A087240 001  
 LYPHOMED 25MG/ML A087431 001  
 PHARMA SERVE NY 25MG/ML A087387 001 Jun 03, 1983  
 25MG/ML A087392 001 Dec 15, 1983  
 SMITH AND NEPHEW 25MG/ML A088429 001 May 30, 1985  
 25MG/ML A088749 001 May 30, 1985  
 TEVA PARENTERAL 25MG/ML A081142 001 Sep 25, 1991

AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%  
 HOSPIRA 100MG/100ML A088147 002 May 03, 1983  
 200MG/100ML A088147 003 May 03, 1983

AMINOPHYLLINE IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER  
 HOSPIRA 100MG/100ML N018924 001 Dec 12, 1984  
 200MG/100ML N018924 002 Dec 12, 1984  
 400MG/100ML N018924 003 Dec 12, 1984  
 500MG/100ML N018924 004 Dec 12, 1984

SOLUTION; ORAL  
 AMINOPHYLLINE  
 MORTON GROVE 105MG/5ML A088156 001 Dec 05, 1983  
 ROXANE 105MG/5ML A088126 001 Aug 19, 1983

AMINOPHYLLINE DYE FREE  
 ACTAVIS MID ATLANTIC 105MG/5ML A087727 001 Apr 16, 1982

SOMOPHYLLIN  
 FISONS 105MG/5ML A086466 001  
 SOMOPHYLLIN-DF  
 FISONS 105MG/5ML A087045 001

SUPPOSITORY; RECTAL  
 TRUPHYLLINE  
 G AND W LABS 250MG A085498 001 Mar 23, 1983  
 500MG A085498 002 Jan 03, 1983

TABLET; ORAL  
 AMINOPHYLLIN  
 GD SEARLE LLC 100MG N002386 002  
 200MG N002386 003

AMINOPHYLLINE  
 ASCOT 100MG A087522 001 Feb 12, 1982  
 200MG A087523 001 Feb 12, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-23(of 375)

\*\* See List Footnote

AMINOPHYLLINE

TABLET;ORAL

## AMINOPHYLLINE

BARR	100MG	A088297 001	Aug 19, 1983
	200MG	A088298 001	Aug 19, 1983
DURAMED PHARMS BARR	100MG	A088182 001	Mar 31, 1983
	200MG	A088183 001	Mar 31, 1983
HALSEY	100MG	A084674 001	
HIKMA INTL PHARMS	100MG	A084540 001	
	200MG	A085003 001	
IDT AUSTRALIA LTD	100MG	A085261 003	
	100MG	A085262 002	
	200MG	A085261 002	
IMPAX LABS	100MG	A084574 001	
	200MG	A084576 001	
KV PHARM	100MG	A085284 001	
	200MG	A085289 001	
LANNETT	100MG	A084588 001	
	200MG	A084588 002	
PAL PAK	100MG	A084533 001	
PANRAY	100MG	A084552 001	
	200MG	A084552 002	
PUREPAC PHARM	100MG	A084699 001	
	200MG	A085333 001	
ROXANE	100MG	A087500 001	Feb 09, 1982
	200MG	A087501 001	Feb 09, 1982
VALEANT PHARM INTL	200MG	A084563 001	
VANGARD	100MG	A088314 001	Oct 03, 1983
	200MG	A088319 001	Oct 03, 1983
VINTAGE PHARMS	100MG	A085409 001	
	200MG	A085410 001	
WATSON LABS	100MG	A085567 001	
	200MG	A085564 001	

TABLET, DELAYED RELEASE;ORAL

## AMINOPHYLLINE

IMPAX LABS	100MG	A084577 001	
	200MG	A084575 001	
TABLICAPS	100MG	A084632 002	
VALE	100MG	A084531 001	
	200MG	A084530 001	

TABLET, EXTENDED RELEASE;ORAL

## PHYLLOCONTIN

PHARM RES ASSOC	225MG	A086760 001	
-----------------	-------	-------------	--

AMINOSALICYLATE SODIUM

POWDER;ORAL

## P.A.S. SODIUM

CENTURY PHARMS	4GM/PACKET	A080947 001	
HEXCEL	100%	A080097 001	

TABLET;ORAL

## PARASAL SODIUM

PANRAY	500MG	N006811 006	
	1GM	N006811 011	

## SODIUM P.A.S.

LANNETT	500MG	A080138 002	
---------	-------	-------------	--

## TEEBACIN

CONSOLIDATED MIDLAND	500MG	N007320 002	
----------------------	-------	-------------	--

AMINOSALICYLATE SODIUM; AMINOSALICYLIC ACID

TABLET;ORAL

## NEOPASALATE

MEDPOINTE PHARM HLC	846MG;112MG	A080059 002	
---------------------	-------------	-------------	--

AMINOSALICYLIC ACID

TABLET;ORAL

## PARASAL

PANRAY	500MG	N006811 001	
	1GM	N006811 002	

**DISCONTINUED DRUG PRODUCT LIST**

6-24(of 375)

\*\* See List Footnote

AMINOSALICYLIC ACID RESIN COMPLEXPOWDER;ORAL  
REZIPAS

BRISTOL MYERS SQUIBB EQ 500MG BASE/GM

N009052 001

AMIODARONE HYDROCHLORIDE

INJECTABLE;INJECTION

AMIODARONE HYDROCHLORIDE

BEDFORD	50MG/ML	A076018 001 Oct 15, 2002
BEDFORD LABS	50MG/ML	A076299 001 Oct 24, 2002
BEN VENUE	50MG/ML	A076088 001 Oct 15, 2002
HOSPIRA	50MG/ML	A076108 001 Oct 15, 2002
INTL MEDICATION SYS	50MG/ML	N021594 001 Feb 04, 2004
PAR STERILE PRODUCTS	50MG/ML	A076394 001 Apr 25, 2003
TEVA PHARMS USA	50MG/ML	A076163 001 Sep 05, 2003

CORDARONE

+ WYETH PHARMS INC 50MG/ML \*\*

N020377 001 Aug 03, 1995

TABLET;ORAL

AMIODARONE HYDROCHLORIDE

MYLAN	200MG	A075188 001 Feb 24, 1999
TEVA	200MG	A074895 001 Apr 16, 1999

CORDARONE

+ WYETH PHARMS INC 200MG \*\*

N018972 001 Dec 24, 1985

AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE;ORAL

ENDEP

ROCHE 40MG/ML

A085749 001

INJECTABLE;INJECTION

AMITRIPTYLINE HYDROCHLORIDE

WATSON LABS 10MG/ML

A085594 001

ELAVIL

ASTRAZENECA 10MG/ML

N012704 001

TABLET;ORAL

AMITID

BRISTOL MYERS SQUIBB	10MG	A086454 001
	25MG	A086454 002
	50MG	A086454 003
	75MG	A086454 004
	100MG	A086454 005

AMITRIL

WARNER CHILCOTT	10MG	A083939 001
	25MG	A083937 001
	50MG	A083938 002
	75MG	A084957 001
	100MG	A085093 001
	150MG	A086295 001

AMITRIPTYLINE HYDROCHLORIDE

AM THERAP	25MG	A088672 001 Nov 20, 1984
	50MG	A088673 001 Nov 20, 1984
	75MG	A088674 001 Nov 20, 1984
	100MG	A088675 001 Nov 20, 1984

ANI PHARMS INC

	10MG	A085031 002
	25MG	A085031 001
	50MG	A085031 003
	75MG	A085031 004

COPELY PHARM

	10MG	A088421 001 Apr 30, 1984
	25MG	A088422 001 Apr 30, 1984
	50MG	A088423 001 Apr 30, 1984
	75MG	A088424 001 Apr 30, 1984
	100MG	A088425 001 Apr 30, 1984
	150MG	A088426 001 Apr 30, 1984

HALSEY

	10MG	A085923 001
	25MG	A085922 001
	50MG	A085925 001
	75MG	A087557 001 Mar 05, 1982

LEDERLE

	100MG	A085926 001 May 20, 1983
	10MG	A085927 001 May 20, 1983
	10MG	A086744 001

	10MG	A087366 001 Jan 04, 1982
--	------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-25(of 375)

\*\* See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

	25MG	A086746 001
	25MG	A087367 001 May 03, 1982
	50MG	A086743 001
	50MG	A087181 001 Jan 04, 1982
	75MG	A086745 001
	75MG	A087369 001 Jan 04, 1982
	100MG	A086747 001
	100MG	A087368 001 May 03, 1982
	150MG	A087370 001 Jan 04, 1982
MUTUAL PHARM	10MG	A085744 001
	25MG	A085627 001
	50MG	A085745 001
	75MG	A085743 001
	100MG	A085742 002 May 11, 1982
	150MG	A089423 001 Feb 17, 1987
PAR PHARM	10MG	A088697 001 Sep 25, 1984
	25MG	A088698 001 Sep 25, 1984
	50MG	A088699 001 Sep 25, 1984
	75MG	A088700 001 Sep 25, 1984
	100MG	A088701 001 Sep 25, 1984
	150MG	A088702 001 Sep 25, 1984
PLIVA	10MG	A088883 001 Sep 26, 1984
	25MG	A088884 001 Sep 26, 1984
	50MG	A088885 001 Sep 26, 1984
	75MG	A088886 001 Sep 26, 1984
	100MG	A088887 001 Sep 26, 1984
	150MG	A088888 001 Sep 26, 1984
PUREPAC PHARM	10MG	A088075 001 Sep 16, 1983
	10MG	A088084 001 Jul 18, 1983
	25MG	A088076 001 May 20, 1983
	25MG	A088085 001 Jul 18, 1983
	50MG	A088077 001 Sep 16, 1983
	50MG	A088105 001 Jul 18, 1983
	75MG	A088078 001 Sep 16, 1983
	75MG	A088106 001 Jul 18, 1983
	100MG	A088079 001 Sep 16, 1983
	100MG	A088107 001 Jul 18, 1983
ROXANE	10MG	A086002 001
	10MG	A086144 001
	25MG	A085944 001
	25MG	A086145 001
	50MG	A085945 001
	50MG	A086143 001
	75MG	A086004 001
	75MG	A086147 001
	100MG	A086003 001
	100MG	A086146 001
	150MG	A086090 001
	150MG	A086148 001
SUN PHARM INDS INC	10MG	A040816 002 Jun 27, 2008
	10MG	A089399 002 Jul 14, 1987
	25MG	A040816 001 Jun 27, 2008
	25MG	A089399 001 Jul 14, 1987
	50MG	A040816 003 Jun 27, 2008
	50MG	A089399 003 Jul 14, 1987
	75MG	A040816 004 Jun 27, 2008
	75MG	A089399 004 Jul 14, 1987
	100MG	A040816 005 Jun 27, 2008
	100MG	A089399 005 Jul 14, 1987
	150MG	A040816 006 Jun 27, 2008
	150MG	A089399 006 Jul 14, 1987
SUPERPHARM	10MG	A088853 001 Nov 13, 1984
	25MG	A088854 001 Nov 13, 1984
	50MG	A088855 001 Nov 13, 1984
	75MG	A088856 001 Nov 13, 1984
	100MG	A088857 001 Nov 13, 1984

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-26(of 375)

\*\* See List Footnote

AMITRIPTYLINE HYDROCHLORIDE

TABLET;ORAL

## AMITRIPTYLINE HYDROCHLORIDE

TEVA	10MG	A086610 001
	25MG	A086859 001
	50MG	A086857 001
	75MG	A086860 001
	100MG	A085836 001
	100MG	A086854 001
	150MG	A086853 001
UCB INC	10MG	A085864 001
	25MG	A085935 001
	50MG	A085936 001
	75MG	A086337 001
	100MG	A086336 001
	150MG	A086335 001
USL PHARMA	25MG	A087775 001 Feb 10, 1982
VANGARD	10MG	A087632 001 Feb 01, 1982
	50MG	A087616 001 Feb 08, 1982
	75MG	A087617 001 Feb 05, 1982
	100MG	A087639 001 Feb 08, 1982
WATSON LABS	10MG	A085816 001
	10MG	A088620 001 Mar 02, 1984
	25MG	A085817 001
	25MG	A088621 001 Mar 02, 1984
	50MG	A085815 001
	50MG	A088622 001 Mar 02, 1984
	75MG	A085819 001
	75MG	A088633 001 Mar 02, 1984
	100MG	A085820 001
	100MG	A088634 001 Mar 02, 1984
	150MG	A085821 001
	150MG	A088635 001 Mar 02, 1984
WEST WARD	10MG	A087647 001 Mar 05, 1982
	25MG	A087278 001
ELAVIL		
+ ASTRAZENECA	10MG **	N012703 001
+	25MG **	N012703 003
+	50MG **	N012703 004
+	75MG **	N012703 005
+	100MG **	N012703 006
+	150MG **	N012703 007
ENDEP		
ROCHE	10MG	A083639 001
	25MG	A083639 002
	50MG	A083639 003
	75MG	A083639 004
	100MG	A083639 005
	150MG	A085303 001

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET;ORAL

## CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

ANDA REPOSITORY	EQ 12.5MG BASE;5MG	A070765 001 Dec 10, 1986
	EQ 25MG BASE;10MG	A070766 001 Dec 10, 1986
PAR PHARM	EQ 12.5MG BASE;5MG	A072277 001 May 09, 1988
	EQ 25MG BASE;10MG	A072278 001 May 09, 1988
USL PHARMA	EQ 12.5MG BASE;5MG	A070477 001 Jan 12, 1988
	EQ 25MG BASE;10MG	A070478 001 Jan 12, 1988
WATSON LABS	EQ 25MG BASE;10MG	A072053 001 Dec 16, 1988
WATSON LABS TEVA	EQ 12.5MG BASE;5MG	A072052 001 Dec 16, 1988
LIMBITROL		
+ HERITAGE PHARMS INC	EQ 12.5MG BASE;5MG **	N016949 001
LIMBITROL DS		
+ HERITAGE PHARMS INC	EQ 25MG BASE;10MG **	N016949 002

**DISCONTINUED DRUG PRODUCT LIST**

6-27(of 375)

\*\* See List Footnote

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

## TABLET;ORAL

ETRAFON 2-10	SCHERING	10MG;2MG **	N014713 007
ETRAFON 2-25	SCHERING	25MG;2MG **	N014713 004
ETRAFON-A	SCHERING	10MG;4MG **	N014713 002
ETRAFON-FORTE	SCHERING	25MG;4MG **	N014713 006
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE			
IVAX SUB TEVA PHARMS	10MG;2MG	A070935 001 Sep 11, 1986	
	10MG;4MG	A070937 001 Sep 11, 1986	
	25MG;2MG	A070936 001 Sep 11, 1986	
	25MG;4MG	A070938 001 Sep 11, 1986	
	50MG;4MG	A070939 001 Sep 12, 1986	
PAR PHARM	10MG;2MG	A070565 001 Sep 11, 1986	
	10MG;4MG	A070620 001 Sep 11, 1986	
	25MG;2MG	A070621 001 Sep 11, 1986	
	25MG;4MG	A070595 001 Sep 11, 1986	
	50MG;4MG	A070574 001 Sep 11, 1986	
SANDOZ	10MG;2MG	A071062 001 Nov 27, 1987	
	10MG;4MG	A071862 001 Dec 21, 1987	
	25MG;2MG	A071063 001 Nov 27, 1987	
	25MG;4MG	A071064 001 Nov 27, 1987	
	50MG;4MG	A071863 001 Dec 21, 1987	
SUN PHARM INDUSTRIES	10MG;2MG	A071077 001 Nov 12, 1986	
	10MG;4MG	A071078 001 Nov 12, 1986	
	25MG;2MG	A070297 001 Nov 12, 1986	
	25MG;4MG	A071079 001 Nov 12, 1986	
WATSON LABS	10MG;2MG	A070373 001 Aug 25, 1986	
	10MG;2MG	A072539 001 Feb 15, 1989	
	10MG;2MG	A073007 001 Oct 17, 1991	
	10MG;4MG	A070375 001 Aug 25, 1986	
	10MG;4MG	A072540 001 Feb 15, 1989	
	25MG;2MG	A073009 001 Oct 17, 1991	
	25MG;2MG	A070374 001 Aug 25, 1986	
	25MG;4MG	A072541 001 Feb 15, 1989	
	25MG;4MG	A073008 001 Oct 17, 1991	
	25MG;4MG	A070376 001 Aug 25, 1986	
	50MG;4MG	A072134 001 Feb 15, 1989	
	50MG;4MG	A073010 001 Oct 17, 1991	
	50MG;4MG	A070377 001 Nov 04, 1986	
	50MG;4MG	A071558 001 Mar 02, 1987	
	50MG;4MG	A072135 001 Feb 15, 1989	
TRIAVIL 2-10			
NEW RIVER	10MG;2MG **	N014715 004	
TRIAVIL 2-25			
NEW RIVER	25MG;2MG **	N014715 002	
TRIAVIL 4-10			
NEW RIVER	10MG;4MG **	N014715 003	
TRIAVIL 4-25			
NEW RIVER	25MG;4MG **	N014715 005	
TRIAVIL 4-50			
NEW RIVER	50MG;4MG **	N014715 006	

AMLEXANOX

## PASTE;DENTAL

## APHTHASOL

ULURU

5%

N020511 001 Dec 17, 1996

## PATCH;TOPICAL

## AMLEXANOX

ULURU

2MG

N021727 001 Sep 29, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-28(of 375)

\*\* See List Footnote

AMLODIPINE BESYLATE

TABLET;ORAL

## AMLODIPINE BESYLATE

FOSUN PHARMA	EQ 2.5MG BASE	A076859 001	Sep 10, 2007
	EQ 5MG BASE	A076859 002	Sep 10, 2007
	EQ 10MG BASE	A076859 003	Sep 10, 2007
GEDEON RICHTER USA	EQ 2.5MG BASE	A077333 001	Jul 17, 2007
	EQ 5MG BASE	A077333 002	Jul 17, 2007
	EQ 10MG BASE	A077333 003	Jul 17, 2007
GENPHARM	EQ 2.5MG BASE	A077362 001	Jul 09, 2007
	EQ 5MG BASE	A077362 002	Jul 09, 2007
	EQ 10MG BASE	A077362 003	Jul 09, 2007
MYLAN PHARMS INC	EQ 2.5MG BASE	A078224 001	Feb 27, 2008
	EQ 5MG BASE	A078224 002	Feb 27, 2008
	EQ 10MG BASE	A078224 003	Feb 27, 2008
PURACAP PHARM	EQ 2.5MG BASE	A078131 001	Sep 04, 2007
	EQ 5MG BASE	A078131 002	Sep 04, 2007
	EQ 10MG BASE	A078131 003	Sep 04, 2007
SOVEREIGN PHARMS	EQ 2.5MG BASE	A204900 001	Jul 23, 2015
	EQ 5MG BASE	A204900 002	Jul 23, 2015
	EQ 10MG BASE	A204900 003	Jul 23, 2015
SUN PHARM INDUSTRIES	EQ 2.5MG BASE	A078081 001	Jan 31, 2008
	EQ 5MG BASE	A078081 002	Jan 31, 2008
	EQ 10MG BASE	A078081 003	Jan 31, 2008
SYNTHON PHARMS	EQ 2.5MG BASE	A077080 001	Jun 27, 2007
	EQ 5MG BASE	A077080 002	Jun 27, 2007
	EQ 10MG BASE	A077080 003	Jun 27, 2007
TABLET, ORALLY DISINTEGRATING;ORAL			
AMLODIPINE BESYLATE			
SYNTHON PHARMS	EQ 2.5MG BASE	N022026 001	Sep 27, 2007
	EQ 5MG BASE	N022026 002	Sep 27, 2007
	EQ 10MG BASE	N022026 003	Sep 27, 2007

AMLODIPINE MALEATE

TABLET;ORAL

## AMVAZ

DR REDDYS LABS INC	2.5MG	N021435 001	Oct 31, 2003
	5MG	N021435 002	Oct 31, 2003
	10MG	N021435 003	Oct 31, 2003

AMMONIA N-13

INJECTABLE; INTRAVENOUS

## AMMONIA N 13

CENTRAL RADIOPHARM	3.75-260mCi/ML	A204539 001	Jun 23, 2015
--------------------	----------------	-------------	--------------

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

## AMMONIUM CHLORIDE

ABBOTT	5MEQ/ML	A083130 001	
GD SEARLE LLC	3MEQ/ML	A086205 001	
AMMONIUM CHLORIDE 0.9% IN NORMAL SALINE			
MCGAW	900MG/100ML	N006580 001	
AMMONIUM CHLORIDE 2.14%			
B BRAUN	40MEQ/100ML	A085734 001	

AMMONIUM LACTATE

CREAM;TOPICAL

## LAC-HYDRIN

RANBAXY	EQ 12% BASE **	N020508 001	Aug 29, 1996
---------	----------------	-------------	--------------

LOTION;TOPICAL

## LAC-HYDRIN

+ RANBAXY	EQ 12% BASE **	N019155 001	Apr 24, 1985
-----------	----------------	-------------	--------------

AMODIAQUINE HYDROCHLORIDE

TABLET;ORAL

## CAMOQUIN HYDROCHLORIDE

PARKE DAVIS	EQ 200MG BASE	N006441 001
-------------	---------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-29(of 375)

\*\* See List Footnote

AMOXAPINETABLET;ORAL  
AMOXAPINE

UPSHER-SMITH LABS	25MG	A072943 001	Jun 28, 1991
	50MG	A072944 001	Jun 28, 1991
	100MG	A072878 001	Jun 28, 1991
	150MG	A072879 001	Jun 28, 1991
WATSON PHARMS TEVA	25MG	A072418 001	May 11, 1989
	50MG	A072419 001	May 11, 1989
	100MG	A072420 001	May 11, 1989
	150MG	A072421 001	May 11, 1989
ASENDIN			
LEDERLE	25MG	N018021 001	
	50MG	N018021 002	
	100MG	N018021 003	
	150MG	N018021 004	

AMOXICILLINCAPSULE;ORAL  
AMOXICILLIN

LABS ATRAL	250MG	A062528 001	Aug 07, 1985
	500MG	A062528 002	Aug 07, 1985
MYLAN	250MG	A062067 001	
	500MG	A062067 002	
SUN PHARM IND S LTD	250MG	A065016 001	Apr 08, 1999
	500MG	A065016 002	Apr 08, 1999
TEVA	250MG	A062853 001	Dec 22, 1987
	250MG	A063030 001	Feb 28, 1989
	500MG	A062854 001	Dec 22, 1987
	500MG	A063031 001	Feb 28, 1989

## AMOXIL

+ GLAXOSMITHKLINE	250MG **	N050459 001
+	500MG **	N050459 002

## TRIMOX

APOTHECON	250MG	A061885 001	
	250MG	A062098 001	
	250MG	A062152 001	
	250MG	A063099 001	Mar 20, 1992
	500MG	A061885 002	
	500MG	A062098 002	
	500MG	A062152 002	
	500MG	A063099 002	Mar 20, 1992

## UTIMOX

PARKE DAVIS	250MG	A062107 001	
	500MG	A062107 002	

## WYMOX

WYETH AYERST	250MG	A062120 001	
	500MG	A062120 002	

## FOR SUSPENSION;ORAL

## AMOXICILLIN

AM ANTIBIOTICS	125MG/5ML	A062059 001	
	250MG/5ML	A062059 002	
MYLAN	125MG/5ML	A062090 001	
	250MG/5ML	A062090 002	
SUN PHARM IND S LTD	200MG/5ML	A065113 001	Nov 29, 2002
	400MG/5ML	A065113 002	Nov 29, 2002
TEVA	125MG/5ML	A062946 001	Nov 01, 1988
	250MG/5ML	A063001 001	Jan 06, 1989

## AMOXIL

+ DR REDDYS LABS INC	200MG/5ML **	N050760 001	Apr 15, 1999
+	400MG/5ML **	N050760 002	Apr 15, 1999
+ GLAXOSMITHKLINE	50MG/ML **	N050460 005	
+	125MG/5ML **	N050460 001	
+	250MG/5ML **	N050460 002	

## LAROTID

+ GLAXOSMITHKLINE	50MG/ML **	N050460 006	
-------------------	------------	-------------	--

## POLYMOX

APOTHECON	125MG/5ML	A061851 001	
	125MG/5ML	A062323 001	
	250MG/5ML	A061851 002	

**DISCONTINUED DRUG PRODUCT LIST**

6-30(of 375)

\*\* See List Footnote

**AMOXICILLIN**FOR SUSPENSION;ORAL  
POLYMOX

	250MG/5ML	A062323 002
TRIMOX		
APOTHECON	50MG/ML	A061886 001
	125MG/5ML	A061886 002
	125MG/5ML	A062099 001
	125MG/5ML	A062154 001
	125MG/5ML	A062885 001 Mar 08, 1988
	250MG/5ML	A061886 003
	250MG/5ML	A062099 002
	250MG/5ML	A062154 002
	250MG/5ML	A062885 002 Mar 08, 1988

UTIMOX

PARKE DAVIS	125MG/5ML	A062127 001
	250MG/5ML	A062127 002

WYMOX

WYETH AYERST	125MG/5ML	A062131 001
	250MG/5ML	A062131 002

TABLET;ORAL

AMOXICILLIN

DAVA PHARMS INC	875MG	A065344 001 Jan 15, 2009
SUN PHARM INDS LTD	500MG	A065059 001 Nov 24, 2000
	875MG	A065059 002 Nov 24, 2000

AMOXIL

+ DR REDDYS LABS INC	500MG **	N050754 002 Jul 10, 1998
+	875MG **	N050754 001 Jul 10, 1998

TABLET, CHEWABLE;ORAL

AMOXICILLIN

APOTHECON	125MG	A064131 001 May 06, 1996
	250MG	A064131 002 May 06, 1996
DAVA PHARMS INC	125MG	A064139 001 Jan 29, 1996
	250MG	A064139 002 Jan 29, 1996
SUN PHARM INDS LTD	125MG	A065021 001 Dec 23, 1999
	200MG	A065060 001 Nov 29, 2000
	250MG	A065021 002 Dec 23, 1999
	400MG	A065060 002 Nov 29, 2000
TEVA	125MG	A064031 001 Dec 19, 1996
	250MG	A064031 002 Dec 19, 1996

AMOXIL

+ DR REDDYS LABS INC	125MG **	N050542 002
	200MG	N050761 001 Apr 15, 1999
+	250MG **	N050542 001
	400MG	N050761 002 Apr 15, 1999

TABLET, FOR SUSPENSION;ORAL

AMOXICILLIN

AUROBINDO PHARMA LTD	200MG	A065324 001 Jan 17, 2007
	400MG	A065324 002 Jan 17, 2007

DISPERMOX

RANBAXY LABS LTD	200MG	A065080 002 Aug 11, 2003
	400MG	A065080 001 Aug 11, 2003
	600MG	A065159 001 Dec 04, 2003

**AMOXICILLIN; CLAVULANATE POTASSIUM**

FOR SUSPENSION;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

SUN PHARM INDS LTD	200MG/5ML;EQ 28.5MG BASE/5ML	A065132 001 Mar 19, 2003
	400MG/5ML;EQ 57MG BASE/5ML	A065132 002 Mar 19, 2003
	600MG/5ML;EQ 42.9MG BASE/5ML	A065207 002 Jan 30, 2007

AUGMENTIN '200'

+ DR REDDYS LABS INC	200MG/5ML;EQ 28.5MG BASE/5ML	N050725 001 May 31, 1996
----------------------	------------------------------	--------------------------

AUGMENTIN '400'

+ DR REDDYS LABS INC	400MG/5ML;EQ 57MG BASE/5ML	N050725 002 May 31, 1996
----------------------	----------------------------	--------------------------

AUGMENTIN ES-600

+ DR REDDYS LABS INC	600MG/5ML;EQ 42.9MG BASE/5ML	N050755 001 Jun 22, 2001
----------------------	------------------------------	--------------------------

TABLET;ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

APOTEX INC	250MG;EQ 125MG BASE	A065333 001 Feb 24, 2009
	500MG;EQ 125MG BASE	A065333 002 Feb 24, 2009

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-31(of 375)

\*\* See List Footnote

AMOXICILLIN; CLAVULANATE POTASSIUM

TABLET;ORAL

## AMOXICILLIN AND CLAVULANATE POTASSIUM

SUN PHARM INDS LTD	875MG;EQ 125MG BASE 500MG;EQ 125MG BASE 875MG;EQ 125MG BASE	A065317 003 Oct 20, 2008 A065109 001 Nov 04, 2002 A065102 001 Sep 17, 2002
AUGMENTIN '250' + DR REDDYS LABS INC	250MG;EQ 125MG BASE **	N050564 001 Aug 06, 1984
AUGMENTIN '500' + DR REDDYS LABS INC	500MG;EQ 125MG BASE **	N050564 002 Aug 06, 1984
TABLET, CHEWABLE;ORAL		
AMOXICILLIN AND CLAVULANATE POTASSIUM		
SANDOZ	200MG;EQ 28.5MG BASE 400MG;EQ 57MG BASE	A065065 001 Apr 18, 2002 A065065 002 Apr 18, 2002
SUN PHARM INDS LTD	200MG;EQ 28.5MG BASE 400MG;EQ 57MG BASE	A065161 001 Dec 03, 2003 A065161 002 Dec 03, 2003
AUGMENTIN '125' + DR REDDYS LABS INC	125MG;EQ 31.25MG BASE **	N050597 001 Jul 22, 1985
AUGMENTIN '200' + DR REDDYS LABS INC	200MG;EQ 28.5MG BASE	N050726 001 May 31, 1996
AUGMENTIN '250' + DR REDDYS LABS INC	250MG;EQ 62.5MG BASE **	N050597 002 Jul 22, 1985
AUGMENTIN '400' + DR REDDYS LABS INC	400MG;EQ 57MG BASE	N050726 002 May 31, 1996

AMPHETAMINE ADIPATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE ADIPATE; DEXTROAMPHETAMINE SULFATE

CAPSULE;ORAL

## DELCOBESE

TEVA	1.25MG;1.25MG;1.25MG;1.25MG ** 2.5MG;2.5MG;2.5MG;2.5MG ** 3.75MG;3.75MG;3.75MG;3.75MG ** 5MG;5MG;5MG;5MG **	A083564 001 A083564 002 A083564 003 A083564 004
------	--	--

TABLET;ORAL

## DELCOBESE

TEVA	1.25MG;1.25MG;1.25MG;1.25MG 2.5MG;2.5MG;2.5MG;2.5MG 3.75MG;3.75MG;3.75MG;3.75MG 5MG;5MG;5MG;5MG	A083563 004 A083563 003 A083563 002 A083563 001
------	--	--

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET;ORAL

## ADDERALL 10

+ TEVA WOMENS	2.5MG;2.5MG;2.5MG;2.5MG **	N011522 007 Feb 13, 1996
---------------	----------------------------	--------------------------

## ADDERALL 12.5

+ TEVA WOMENS	3.125MG;3.125MG;3.125MG;3.125MG **	N011522 012 Aug 31, 2000
---------------	------------------------------------	--------------------------

## ADDERALL 15

+ TEVA WOMENS	3.75MG;3.75MG;3.75MG;3.75MG **	N011522 013 Aug 31, 2000
---------------	--------------------------------	--------------------------

## ADDERALL 20

+ TEVA WOMENS	5MG;5MG;5MG;5MG **	N011522 008 Feb 13, 1996
---------------	--------------------	--------------------------

## ADDERALL 30

+ TEVA WOMENS	7.5MG;7.5MG;7.5MG;7.5MG **	N011522 010 May 12, 1997
---------------	----------------------------	--------------------------

## ADDERALL 5

+ TEVA WOMENS	1.25MG;1.25MG;1.25MG;1.25MG **	N011522 009 May 12, 1997
---------------	--------------------------------	--------------------------

## ADDERALL 7.5

+ TEVA WOMENS	1.875MG;1.875MG;1.875MG;1.875MG **	N011522 011 Aug 31, 2000
---------------	------------------------------------	--------------------------

## DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

TEVA PHARMS	1.25MG;1.25MG;1.25MG;1.25MG 2.5MG;2.5MG;2.5MG;2.5MG 5MG;5MG;5MG;5MG 7.5MG;7.5MG;7.5MG;7.5MG	A040472 001 Sep 30, 2003 A040472 002 Sep 30, 2003 A040472 003 Sep 30, 2003 A040472 004 Sep 30, 2003
-------------	--	--

AMPHETAMINE RESIN COMPLEX; DEXTROAMPHETAMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE;ORAL

## BIPHETAMINE 12.5

UCB INC	EQ 6.25MG BASE;EQ 6.25MG BASE	N010093 007
---------	-------------------------------	-------------

## BIPHETAMINE 20

UCB INC	EQ 10MG BASE;EQ 10MG BASE	N010093 003
---------	---------------------------	-------------

## BIPHETAMINE 7.5

UCB INC	EQ 3.75MG BASE;EQ 3.75MG BASE	N010093 009
---------	-------------------------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-32(of 375)

\*\* See List Footnote

AMPHETAMINE SULFATE

TABLET;ORAL

AMPHETAMINE SULFATE

LANNETT

5MG

10MG

A083901 001 Aug 31, 1984

A083901 002 Aug 31, 1984

AMPHOTERICIN B

CREAM;TOPICAL

FUNGIZONE

APOTHECON

3%

N050314 001

INJECTABLE;INJECTION

AMPHOTERICIN B

ABBOTT

50MG/VIAL

A064141 001 Dec 23, 1996

ABRAXIS PHARM

50MG/VIAL

A062728 001 Apr 13, 1987

TEVA PARENTERAL

50MG/VIAL

A064062 001 Mar 31, 1995

FUNGIZONE

APOTHECON

50MG/VIAL

A060517 001

INJECTABLE, LIPID COMPLEX;INJECTION

AMPHOTEC

ALKOPHARMA USA

50MG/VIAL

N050729 001 Nov 22, 1996

100MG/VIAL

N050729 002 Nov 22, 1996

LOTION;TOPICAL

FUNGIZONE

APOTHECON

3%

A060570 001

OINTMENT;TOPICAL

FUNGIZONE

APOTHECON

3%

N050313 001

SUSPENSION;ORAL

FUNGIZONE

BRISTOL MYERS SQUIBB

100MG/ML

N050341 003

AMPICILLIN SODIUM

INJECTABLE;INJECTION

AMPICILLIN SODIUM

APOTHECON

EQ 125MG BASE/VIAL

A062860 001 Feb 05, 1988

EQ 250MG BASE/VIAL

A062860 002 Feb 05, 1988

EQ 500MG BASE/VIAL

A062860 003 Feb 05, 1988

EQ 1GM BASE/VIAL

A062860 004 Feb 05, 1988

EQ 2GM BASE/VIAL

A062860 005 Feb 05, 1988

CONSOLIDATED PHARM

EQ 125MG BASE/VIAL

A061936 005

EQ 250MG BASE/VIAL

A061936 001

EQ 500MG BASE/VIAL

A061936 002

EQ 1GM BASE/VIAL

A061936 003

EQ 2GM BASE/VIAL

A061936 004

HANFORD GC

EQ 125MG BASE/VIAL

A062772 005 Apr 15, 1993

EQ 500MG BASE/VIAL

A062772 008 Apr 15, 1993

EQ 1GM BASE/VIAL

A062772 002 Apr 15, 1993

EQ 2GM BASE/VIAL

A062772 004 Apr 15, 1993

INTL MEDICATION

EQ 1GM BASE/VIAL

A062634 002 Jan 09, 1987

EQ 2GM BASE/VIAL

A062634 003 Jan 09, 1987

LILLY

EQ 500MG BASE/VIAL

A062565 001 Apr 04, 1985

EQ 1GM BASE/VIAL

A062565 002 Apr 04, 1985

EQ 2GM BASE/VIAL

A062565 003 Jun 24, 1986

WATSON LABS INC

EQ 125MG BASE/VIAL

A062816 001 Oct 24, 1988

EQ 250MG BASE/VIAL

A062816 002 Oct 24, 1988

EQ 500MG BASE/VIAL

A062816 003 Oct 24, 1988

EQ 1GM BASE/VIAL

A062816 004 Oct 24, 1988

EQ 2GM BASE/VIAL

A062816 005 Oct 24, 1988

EQ 10GM BASE/VIAL

A062994 001 Sep 15, 1988

WEST-WARD PHARMS INT

EQ 125MG BASE/VIAL

A062692 001 Jun 24, 1986

EQ 250MG BASE/VIAL

A062692 002 Jun 24, 1986

EQ 500MG BASE/VIAL

A062692 003 Jun 24, 1986

EQ 1GM BASE/VIAL

A062692 004 Jun 24, 1986

EQ 2GM BASE/VIAL

A062692 005 Jun 24, 1986

EQ 10GM BASE/VIAL

A062692 006 Jun 24, 1986

OMNIPEN-N

WYETH AYERST

EQ 125MG BASE/VIAL

A060626 001

EQ 125MG BASE/VIAL

A062718 001 Dec 16, 1986

EQ 250MG BASE/VIAL

A060626 002

EQ 250MG BASE/VIAL

A062718 002 Dec 16, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-33(of 375)

\*\* See List Footnote

AMPICILLIN SODIUMINJECTABLE; INJECTION  
OMNIPEN-N

PENBRITIN-S			
+ WYETH AYERST	EQ 125MG BASE/VIAL **	N050072 001	
+ EQ 250MG BASE/VIAL **	N050072 002		
+ EQ 500MG BASE/VIAL **	N050072 003		
+ EQ 1GM BASE/VIAL **	N050072 004		
+ EQ 2GM BASE/VIAL **	N050072 005		
+ EQ 4GM BASE/VIAL **	N050072 006		
POLYCILLIN-N			
BRISTOL	EQ 125MG BASE/VIAL **	N050309 001	
EQ 250MG BASE/VIAL **	N050309 002		
EQ 500MG BASE/VIAL **	N050309 003		
EQ 1GM BASE/VIAL **	N050309 004		
EQ 2GM BASE/VIAL **	N050309 005		
TOTACILLIN-N			
GLAXOSMITHKLINE	EQ 125MG BASE/VIAL	A060677 001	
EQ 250MG BASE/VIAL	A060677 002		
EQ 500MG BASE/VIAL	A060677 003		
EQ 1GM BASE/VIAL	A060677 004		
EQ 2GM BASE/VIAL	A062727 001 Dec 19, 1986		
EQ 2GM BASE/VIAL	A060677 005		
EQ 10GM BASE/VIAL	A062727 002 Dec 19, 1986		
A060677 006			

AMPICILLIN SODIUM; SULBACTAM SODIUMINJECTABLE; INJECTION  
UNASYN

PFIZER	EQ 500MG BASE/VIAL; EQ 250MG BASE/VIAL	N050608 003 Dec 31, 1986
--------	--	--------------------------

AMPICILLIN/AMPICILLIN TRIHYDRATECAPSULE; ORAL  
AMCILL

PARKE DAVIS	EQ 250MG BASE	A062041 001
	EQ 500MG BASE	A062041 002
AMPICILLIN TRIHYDRATE		
AM ANTIBIOTICS	EQ 250MG BASE	A061602 001
	EQ 500MG BASE	A061602 002
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A060765 001
	EQ 500MG BASE	A060765 002
LEDERLE	EQ 250MG BASE	A062208 001
	EQ 500MG BASE	A062208 002
MYLAN	EQ 250MG BASE	A061755 001
	EQ 500MG BASE	A061755 002
PUREPAC PHARM	EQ 250MG BASE	A061853 001
	EQ 500MG BASE	A061853 002
TEVA	EQ 250MG BASE	A061502 001
	EQ 500MG BASE	A061502 002
VITARINE	EQ 250MG BASE	A061387 001
	EQ 500MG BASE	A061387 003
OMNIPEN (AMPICILLIN)		
WYETH AYERST	250MG	A060624 001
	500MG	A060624 002
PENBRITIN		
WYETH AYERST	EQ 250MG BASE	A060908 001
	EQ 500MG BASE	A060908 002
PFIZERPEN-A		
PFIZER	EQ 250MG BASE	A062050 001
	EQ 500MG BASE	A062050 002
POLYCILLIN		
BRISTOL	EQ 250MG BASE	N050310 001
	EQ 500MG BASE	N050310 002

**DISCONTINUED DRUG PRODUCT LIST**

6-34(of 375)

\*\* See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE;ORAL

PRINCIPEN

APOTHECON	EQ 250MG BASE	A062888 001 Mar 04, 1988
	EQ 500MG BASE	A062888 002 Mar 04, 1988
BRISTOL MYERS SQUIBB	EQ 250MG BASE	A061392 001
	EQ 500MG BASE	A061392 002

PRINCIPEN '250'

APOTHECON	EQ 250MG BASE	A062157 002
	EQ 250MG BASE	N050056 001

PRINCIPEN '500'

APOTHECON	EQ 500MG BASE	A062157 001
	EQ 500MG BASE	N050056 002

TOTACILLIN

GLAXOSMITHKLINE	EQ 250MG BASE	A060060 001
	EQ 250MG BASE	A062212 001
	EQ 500MG BASE	A060060 002
	EQ 500MG BASE	A062212 002

FOR SUSPENSION;ORAL

AMCILL

PARKE DAVIS	EQ 125MG BASE/5ML	A062030 001
	EQ 250MG BASE/5ML	A062030 002

AMPICILLIN TRIHYDRATE

AM ANTIBIOTICS	EQ 125MG BASE/5ML	A061601 001
	EQ 250MG BASE/5ML	A061601 002
MYLAN	EQ 125MG BASE/5ML	A061829 002
	EQ 250MG BASE/5ML	A061829 001
PUREPAC PHARM	EQ 125MG BASE/5ML	A061980 001
	EQ 250MG BASE/5ML	A061980 002
TEVA	EQ 125MG BASE/5ML	A061370 001
	EQ 250MG BASE/5ML	A061370 002

OMNIPEN (AMPICILLIN)

WYETH AYERST	100MG/ML	A060625 001
	125MG/5ML	A060625 002
	250MG/5ML	A060625 003
	500MG/5ML	A060625 004

PENBRITIN

WYETH AYERST	EQ 100MG BASE/ML	N050019 001
	EQ 125MG BASE/5ML	N050019 002
	EQ 250MG BASE/5ML	N050019 003

PFIZERPEN-A

PFIZER	EQ 125MG BASE/5ML	A062049 001
	EQ 250MG BASE/5ML	A062049 002

POLYCILLIN

APOTHECON	EQ 125MG BASE/5ML	A062297 001
	EQ 250MG BASE/5ML	A062297 002
BRISTOL	EQ 100MG BASE/ML	N050308 004
	EQ 125MG BASE/5ML	N050308 001
	EQ 250MG BASE/5ML	N050308 002
	EQ 500MG BASE/5ML	N050308 003

PRINCIPEN

APOTHECON	EQ 100MG BASE/ML	A061394 001
	EQ 125MG BASE/5ML	A061394 002
	EQ 250MG BASE/5ML	A061394 003

PRINCIPEN '125'

APOTHECON	EQ 125MG BASE/5ML	A060127 002
	EQ 125MG BASE/5ML	A062151 001

PRINCIPEN '250'

APOTHECON	EQ 250MG BASE/5ML	A060127 001
	EQ 250MG BASE/5ML	A062151 002

TOTACILLIN

GLAXOSMITHKLINE	EQ 125MG BASE/5ML	A060666 001
	EQ 125MG BASE/5ML	A062223 001
	EQ 250MG BASE/5ML	A060666 002
	EQ 250MG BASE/5ML	A062223 002

TABLET, CHEWABLE;ORAL

POLYCILLIN

BRISTOL	EQ 125MG BASE	N050093 001
---------	---------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-35(of 375)

\*\* See List Footnote

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

CAPSULE;ORAL PRINCIPEN W/ PROBENECID APOTHECON	EQ 389MG BASE;111MG EQ 389MG BASE;111MG	A062150 001 N050488 001
FOR SUSPENSION;ORAL POLYCILLIN-PRB APOTHECON BRISTOL	EQ 3.5GM BASE/BOT;1GM/BOT EQ 3.5GM BASE/BOT;1GM/BOT	A061898 001 N050457 001
PROBAMPACIN G AND W LABS INC	EQ 3.5GM BASE/BOT;1GM/BOT	A061741 001

AMPRENAVIR

CAPSULE;ORAL AGENERASE GLAXOSMITHKLINE	50MG 150MG	N021007 001 Apr 15, 1999 N021007 002 Apr 15, 1999
SOLUTION;ORAL AGENERASE + GLAXOSMITHKLINE	15MG/ML **	N021039 001 Apr 15, 1999

ANAGRELIDE HYDROCHLORIDE

CAPSULE;ORAL AGRYLIN + SHIRE LLC	EQ 1MG BASE **	N020333 002 Mar 14, 1997
ANAGRELIDE HYDROCHLORIDE MYLAN PHARMS INC	EQ 0.5MG BASE EQ 0.5MG BASE EQ 1MG BASE EQ 1MG BASE	A076811 001 Apr 18, 2005 A077613 001 Jun 27, 2006 A076811 002 Apr 18, 2005 A077613 002 Jun 27, 2006
ROXANE	EQ 0.5MG BASE EQ 1MG BASE	A076489 001 Apr 18, 2005 A076489 002 Apr 18, 2005
UPSHER-SMITH LABS	EQ 0.5MG BASE EQ 1MG BASE	A076683 001 Apr 18, 2005 A076683 002 Apr 18, 2005
WATSON LABS	EQ 0.5MG BASE EQ 1MG BASE	A076417 001 Apr 18, 2005 A076417 002 Apr 18, 2005

ANASTROZOLE

TABLET;ORAL ANASTROZOLE IMPAX LABS INC	1MG	A091242 001 May 31, 2012
KREMERS URBAN PHARMS	1MG	A091331 001 Jan 05, 2011
SANDOZ	1MG	A079007 001 Jun 28, 2010
SUN PHARM IND LTD	1MG	A091177 001 Jul 15, 2011
SYNTON PHARMS	1MG	A078322 001 Jun 28, 2010
WATSON LABS TEVA	1MG	A078984 001 Jun 28, 2010

ANILERIDINE HYDROCHLORIDE

TABLET;ORAL LERITINE MERCK	EQ 25MG BASE	N010585 002

ANILERIDINE PHOSPHATE

INJECTABLE; INJECTION LERITINE MERCK	25MG/ML	N010520 003

ANISINDIONE

TABLET;ORAL MIRADON SCHERING	50MG	N010909 003

ANISOTROPINE METHYLBROMIDE

TABLET;ORAL ANISOTROPINE METHYLBROMIDE WATSON LABS	50MG	A086046 001
VALPIN 50 ENDO PHARMS	50MG	N013428 001

**DISCONTINUED DRUG PRODUCT LIST**

6-36(of 375)

\*\* See List Footnote

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

VASOCON-A

NOVARTIS

0.5%;0.05%

N018746 002 Jul 11, 1994

APOMORPHINE HYDROCHLORIDE

INJECTABLE;SUBCUTANEOUS

APOKYN

US WORLDMEDS

20MG/2ML (10MG/ML)

N021264 001 Apr 20, 2004

APROTININ

INJECTABLE;INJECTION

TRASYLOL

BAYER HLTHCARE

10,000KIU/ML

N020304 001 Dec 29, 1993

ARbutamine Hydrochloride

INJECTABLE;INJECTION

GENESA

GENSIA AUTOMEDICS

0.05MG/ML

N020420 001 Sep 12, 1997

ARDEPARIN SODIUM

INJECTABLE;INJECTION

NORMIFLO

+ PHARMACIA AND UPJOHN

5,000 UNITS/0.5ML \*\*

N020227 002 May 23, 1997

+

10,000 UNITS/0.5ML \*\*

N020227 001 May 23, 1997

ARGATROBAN

SOLUTION;IV (INFUSION)

ARGATROBAN IN DEXTROSE

SANDOZ

125MG/125ML (1MG/ML)

N201743 001 May 09, 2011

ARIPIPRAZOLE

INJECTABLE;INTRAMUSCULAR

ABILIFY

OTSUKA

9.75MG/1.3ML (7.5MG/ML)

N021866 001 Sep 20, 2006

SOLUTION;ORAL

ABILIFY

+ OTSUKA

1MG/ML \*\*

N021713 001 Dec 10, 2004

TABLET, ORALLY DISINTEGRATING;ORAL

ABILIFY

OTSUKA

10MG \*\*

N021729 002 Jun 07, 2006

15MG \*\*

N021729 003 Jun 07, 2006

+

20MG \*\*

N021729 004 Jun 07, 2006

+

30MG \*\*

N021729 005 Jun 07, 2006

ARMODAFINIL

TABLET;ORAL

ARMODAFINIL

WATSON LABS INC

100MG

A200156 002 Aug 29, 2012

200MG

A200156 004 Aug 29, 2012

NUVIGIL

+ CEPHALON

100MG \*\*

N021875 002 Mar 26, 2009

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANthenol; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; PALMITATE; VITAMIN E

INJECTABLE;INJECTION

BEROCCA PN

ROCHE

50MG/ML;0.03MG/ML;0.0025MG/ML;7.5MG/ML;

N006071 003 Oct 10, 1985

100

IU/ML;0.2MG/ML;20MG/ML;2MG/ML;1.8MG/ML;

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANthenol; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE;INJECTION

M.V.C. 9+3

ABRAXIS PHARM

10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;2

N018440 002 Aug 08, 1985

0

IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/

M.V.I.-12 ADULT

HOSPIRA

10MG/ML;0.006MG/ML;0.5MCG/ML;1.5MG/ML;2

N008809 004 Aug 08, 1985

0

IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/

+

20MG/ML;0.006MG/ML;0.05MCG/ML;1.5MG/ML;

N008809 006 Sep 09, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-37(of 375)

\*\* See List Footnote

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 ADULT

0.0005MG/ML; 0.06MG/ML; 4MG/ML; 0.6MG/ML; 0  
.36MG/ML; 0.6MG/ML; 0.1MG/ML; 1MG/ML

MVC PLUS

WATSON LABS

10MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2  
0  
IU/ML; 0.04MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 ADULT

HOSPIRA

20MG/ML; 0.006MG/ML; 0.5MCG/ML; 1.5MG/ML; 2  
0  
IU/ML; 0.6MG/ML; 4MG/ML; 0.4MG/ML; 0.36MG/

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN 5'-PHOSPHATE SODIUM; THIAMINE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 LYOPHILIZED

TELIGENT PHARMA INC

100MG/VIAL; 0.06MG/VIAL; 0.005MG/VIAL; 15M  
G/VIAL; 5MCG/VIAL; 0.4MG/VIAL; 40MG/VIAL; 4  
MG/VIAL; 3.6MG/VIAL; 3MG/VIAL; 1MG/VIAL; 10  
MG/VIAL

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

VITAMED

HOSPIRA

N/A, 80MG/VIAL; N/A, 0.02MG/VIAL; N/A, 0.001  
MG/VIAL; 400  
IU/10ML, N/A; N/A, 0.14MG/VIAL; N/A, 17MG/VI  
AL; N/A, 5MG/VIAL; 0.2MG/10ML, N/A; N/A, 1MG/  
VIAL; N/A, 1.4MG/VIAL; N/A, 1.2MG/VIAL; EQ  
2,300 UNITS BASE/10ML, N/A; 7 IU/10ML, N/A

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

PEG-3350, SODIUM SULFATE, SODIUM CHLORIDE, POTASSIUM CHLORIDE, SODIUM ASCORBATE AND ASCORBIC NOVEL LABS INC 4.7GM; 100GM; 1.015GM; 5.9GM; 2.691GM; 7.5GM A090145 001 Jan 25, 2012

ASPIRIN

TABLET; ORAL

BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN

BAYER 500MG

N021317 001 Oct 18, 2001

TABLET, EXTENDED RELEASE; ORAL

8-HOUR BAYER

BAYER 650MG

N016030 001

MEASURIN

BAYER 650MG

N016030 002

ASPIRIN; BUTALBITAL

TABLET; ORAL

AXOTAL

SAVAGE LABS 650MG; 50MG

A088305 001 Oct 13, 1983

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

NOSTRUM LABS INC 325MG; 50MG; 40MG

A078149 001 Jun 13, 2007

WATSON LABS 325MG; 50MG; 40MG

A086231 002 Feb 12, 1985

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

ACTAVIS ELIZABETH 325MG; 50MG; 40MG

A086710 002 Aug 23, 1983

HALSEY 325MG; 50MG; 40MG

A089448 001 Dec 01, 1986

IVAX PHARMS 325MG; 50MG; 40MG

A085441 002 Oct 31, 1984

PURACAP PHARM 325MG; 50MG; 40MG

A087048 002 Dec 09, 1983

QUANTUM PHARMICS 325MG; 50MG; 40MG

A088972 001 Jun 18, 1985

SANDOZ 325MG; 50MG; 40MG

A086398 002 Apr 06, 1984

WATSON LABS 325MG; 50MG; 40MG

A086237 002 Mar 23, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-38(of 375)

\*\* See List Footnote

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET;ORAL

FIORINAL

+ ALLERGAN SALES LLC 325MG;50MG;40MG \*\*

N017534 003 Apr 16, 1986

LANORINAL

LANNETT 325MG;50MG;40MG

A086986 002 Oct 18, 1985

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE;ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

VINTAGE PHARMS LLC 325MG;50MG;40MG;30MG

A075351 001 Mar 05, 1999

WATSON LABS 325MG;50MG;40MG;30MG

A074359 001 Aug 31, 1995

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET;ORAL

INVAGESIC

SANDOZ 385MG;30MG;25MG

A074817 001 Nov 27, 1996

INVAGESIC FORTE

SANDOZ 770MG;60MG;50MG

A074817 002 Nov 27, 1996

NORGESIC

+ MEDICIS 385MG;30MG;25MG \*\*

N013416 003 Oct 27, 1982

NORGESIC FORTE

+ MEDICIS 770MG;60MG;50MG \*\*

N013416 004 Oct 27, 1982

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

STEVENS J 385MG;30MG;25MG

A074988 001 Apr 30, 1999

770MG;60MG;50MG

A074988 002 Apr 30, 1999

ORPHEGESIC

PRINSTON INC 385MG;30MG;25MG

A075141 001 May 29, 1998

ORPHEGESIC FORTE

PRINSTON INC 770MG;60MG;50MG

A075141 002 May 29, 1998

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE;ORAL

COMPOUND 65

ALRA 389MG;32.4MG;65MG

A084553 002 Aug 17, 1983

DARVON COMPOUND

XANODYNE PHARM 389MG;32.4MG;32MG

N010996 006 Mar 08, 1983

DARVON COMPOUND-65

XANODYNE PHARM 389MG;32.4MG;65MG

N010996 007 Mar 08, 1983

PROPOXYPHENE COMPOUND 65

IVAX SUB TEVA PHARMS 389MG;32.4MG;65MG

A083077 002 Dec 07, 1984

SANDOZ 389MG;32.4MG;65MG

A080044 002 Sep 16, 1983

TEVA 389MG;32.4MG;65MG

A089025 001 Mar 29, 1985

PROPOXYPHENE COMPOUND-65

SANDOZ 389MG;32.4MG;65MG

A083101 002 Jun 24, 1985

PROPOXYPHENE HYDROCHLORIDE W/ ASPIRIN AND CAFFEINE

WATSON LABS 389MG;32.4MG;65MG

A085732 002 Sep 03, 1984

ASPIRIN; CARISOPRODOL

TABLET;ORAL

CARISOPRODOL COMPOUND

WATSON LABS 325MG;200MG

A088809 001 Oct 03, 1985

SOMA COMPOUND

MEDA PHARMS 325MG;200MG \*\*

N012365 005 Jul 11, 1983

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET;ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

OXFORD PHARMS 325MG;200MG;16MG

A040283 001 Dec 29, 1998

SOMA COMPOUND W/ CODEINE

MEDA PHARMS 325MG;200MG;16MG \*\*

N012366 002 Jul 11, 1983

ASPIRIN; HYDROCODONE BITARTRATE

TABLET;ORAL

AZDONE

SCHWARZ PHARMA 500MG;5MG \*\*

A089420 001 Jan 25, 1988

VICOPRIN

ABBOTT 500MG;5MG

A086333 001 Sep 14, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-39(of 375)

\*\* See List Footnote

**ASPIRIN; MEPROBAMATE**

TABLET;ORAL			
EQUAGESIC			
SUN PHARM INDUSTRIES	325MG;200MG	N011702	003 Dec 29, 1983
MEPRO-ASPIRIN			
SANDOZ	325MG;200MG	A089127	001 Mar 02, 1987
MEPROBAMATE AND ASPIRIN			
PAR PHARM	325MG;200MG	A089126	001 Aug 19, 1986
MICRAININ			
MEDPOINTE PHARM HLC	325MG;200MG	A084978	001
Q-GESIC			
QUANTUM PHARMICS	325MG;200MG	A088740	001 Jun 01, 1984

**ASPIRIN; METHOCARBAMOL**

TABLET;ORAL			
METHOCARBAMOL AND ASPIRIN			
IVAX SUB TEVA PHARMS	325MG;400MG	A087211	001 Dec 22, 1982
MCNEIL	325MG;400MG	A089193	001 Feb 12, 1986
PAR PHARM	325MG;400MG	A089657	001 Nov 04, 1988
ROBAXISAL			
ROBINS AH	325MG;400MG	N012281	001

**ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE**

TABLET;ORAL			
CODOXY			
HALSEY	325MG;4.5MG;0.38MG	A087464	001 Jul 01, 1982
OXYCODONE AND ASPIRIN			
SUN PHARM INDUSTRIES	325MG;4.5MG;0.38MG	A040260	001 Jul 17, 1998
	325MG;4.5MG;0.38MG	A087794	001 May 26, 1982
WATSON LABS	325MG;4.5MG;0.38MG	A040255	001 Feb 27, 1998
OXYCODONE AND ASPIRIN (HALF-STRENGTH)			
ROXANE	325MG;2.25MG;0.19MG	A087742	001 Jun 04, 1982
PERCODAN			
ENDO PHARMS	325MG;4.5MG;0.38MG **	N007337	006
PERCODAN-DEMI			
ENDO PHARMS	325MG;2.25MG;0.19MG **	N007337	005
ROXIPRIN			
ROXANE	325MG;4.5MG;0.38MG	A087743	001 Jun 04, 1982

**ASPIRIN; PENTAZOCINE HYDROCHLORIDE**

TABLET;ORAL			
TALWIN COMPOUND			
+ SANOFI AVENTIS US	325MG;EQ 12.5MG BASE **	N016891	001

**ASPIRIN; PRAVASTATIN SODIUM**

TABLET, TABLET;ORAL			
PRAVIGARD PAC (COPACKAGED)			
BRISTOL MYERS SQUIBB	325MG,N/A;N/A,80MG	N021387	006 Jun 24, 2003
TABLET, TABLET, TABLET;ORAL			
PRAVIGARD PAC (COPACKAGED)			
BRISTOL MYERS SQUIBB	81MG,N/A;N/A,20MG	N021387	001 Jun 24, 2003
	81MG,N/A;N/A,40MG	N021387	002 Jun 24, 2003
	81MG,N/A;N/A,80MG	N021387	003 Jun 24, 2003
	325MG,N/A;N/A,20MG	N021387	004 Jun 24, 2003
	325MG,N/A;N/A,40MG	N021387	005 Jun 24, 2003

**ASPIRIN; PROPOXYPHENE HYDROCHLORIDE**

CAPSULE;ORAL			
DARVON W/ ASA			
XANODYNE PHARM	325MG;65MG	N010996	005

**ASPIRIN; PROPOXYPHENE NAPSYLATE**

CAPSULE;ORAL			
DARVON-N W/ ASA			
AAIPHARMA LLC	325MG;100MG	N016829	001
TABLET;ORAL			
DARVON-N W/ ASA			
AAIPHARMA LLC	325MG;100MG	N016863	001

**DISCONTINUED DRUG PRODUCT LIST**

6-40(of 375)

\*\* See List Footnote

ATAZANAVIR SULFATE

CAPSULE;ORAL

REYATAZ

+ BRISTOL MYERS SQUIBB EQ 100MG BASE \*\*

N021567 001 Jun 20, 2003

ATENOLOL

INJECTABLE;INJECTION

TENORMIN

+ ASTRAZENECA 0.5MG/ML \*\*

N019058 001 Sep 13, 1989

TABLET;ORAL

ATENOLOL

ABLE	25MG	A076907 001 Jul 30, 2004
	50MG	A076907 002 Jul 30, 2004
	100MG	A076907 003 Jul 30, 2004
APOTHECON	50MG	A073317 001 Mar 20, 1992
	100MG	A073318 001 Mar 20, 1992
DAVA PHARMS INC	25MG	A074099 001 Apr 28, 1992
MYLAN	25MG	A074126 003 Aug 26, 1998
	50MG	A074126 001 Mar 23, 1994
	100MG	A074126 002 Mar 23, 1994
NORTHSTAR HLTHCARE	25MG	A078254 001 Sep 25, 2009
	50MG	A078254 002 Sep 25, 2009
	100MG	A078254 003 Sep 25, 2009
NOSTRUM LABS	50MG	A074127 001 Feb 21, 1995
	100MG	A074127 002 Feb 21, 1995
PLIVA	25MG	A074101 001 Jul 17, 1997
	50MG	A074101 002 Jul 17, 1997
	100MG	A074101 003 Jul 17, 1997
SANDOZ	25MG	A074265 001 Feb 28, 1994
	50MG	A074265 002 Feb 28, 1994
	100MG	A074265 003 Feb 28, 1994
SCS	50MG	A073676 001 Oct 30, 1992
	100MG	A073676 002 Oct 30, 1992
TEVA	50MG	A073315 001 May 28, 1993
	100MG	A073316 001 May 28, 1993
TEVA PHARMS	50MG	A074120 001 Feb 24, 1995
	100MG	A074120 002 Feb 24, 1995
WATSON LABS	50MG	A073352 001 Dec 27, 1991
WATSON LABS TEVA	100MG	A073353 001 Dec 27, 1991

ATENOLOL; CHLORTHALIDONE

TABLET;ORAL

ATENOLOL AND CHLORTHALIDONE

NOSTRUM LABS	50MG;25MG	A074404 001 May 14, 1998
	100MG;25MG	A074404 002 May 14, 1998
PLIVA	50MG;25MG	A074107 001 Sep 24, 1997
	100MG;25MG	A074107 002 Sep 24, 1997

ATOMOXETINE HYDROCHLORIDE

CAPSULE;ORAL

ATOMOXETINE HYDROCHLORIDE

ZYDUS PHARMS USA INC	18MG	A079017 001 Sep 17, 2010
	25MG	A079017 002 Sep 17, 2010
	40MG	A079017 003 Sep 17, 2010
	60MG	A079017 004 Sep 17, 2010
	80MG	A079017 005 Sep 17, 2010
	100MG	A079017 006 Sep 17, 2010

STRATTERA

LILLY 5MG N021411 001 Nov 26, 2002

ATORVASTATIN CALCIUM

TABLET;ORAL

ATORVASTATIN CALCIUM

TEVA PHARMS	EQ 10MG BASE	A078773 001 May 29, 2012
	EQ 20MG BASE	A078773 002 May 29, 2012
	EQ 40MG BASE	A078773 003 May 29, 2012
	EQ 80MG BASE	A078773 004 May 29, 2012

**DISCONTINUED DRUG PRODUCT LIST**

6-41(of 375)

\*\* See List Footnote

ATORVASTATIN CALCIUM; EZETIMIBE

TABLET;ORAL

LIPTRUZET

+ MERCK SHARP DOHME	EQ 10MG BASE;10MG **	N200153 001	May 03, 2013
+	EQ 20MG BASE;10MG **	N200153 002	May 03, 2013
+	EQ 40MG BASE;10MG **	N200153 003	May 03, 2013
+	EQ 80MG BASE;10MG **	N200153 004	May 03, 2013

ATOVAQUONE

TABLET;ORAL

MEPRON

+ GLAXOSMITHKLINE LLC	250MG **	N020259 001	Nov 25, 1992
-----------------------	----------	-------------	--------------

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

BAXTER HLTHCARE	10MG/ML	A074824 001	Sep 30, 1997
BAXTER HLTHCARE CORP	10MG/ML	A074753 001	Jan 23, 1997
HOSPIRA	10MG/ML	A074632 001	Dec 23, 1996
	10MG/ML	A074740 001	Mar 28, 1997
TEVA PARENTERAL	10MG/ML	A074784 001	Jun 11, 1997
WATSON PHARMS TEVA	10MG/ML	A074945 001	Jul 28, 1998

ATRACURIUM BESYLATE PRESERVATIVE FREE

BAXTER HLTHCARE	10MG/ML	A074825 001	Sep 30, 1997
BAXTER HLTHCARE CORP	10MG/ML	A074768 001	Jan 23, 1997
HOSPIRA	10MG/ML	A074633 001	Dec 23, 1996
	10MG/ML	A074639 001	Mar 25, 1997
	10MG/ML	A074741 001	Mar 28, 1997
WATSON LABS INC	10MG/ML	A074944 001	Jul 28, 1998

TRACRIUM

+ HOSPIRA	10MG/ML **	N018831 002	Jun 20, 1985
-----------	------------	-------------	--------------

TRACRIUM PRESERVATIVE FREE

+ HOSPIRA	10MG/ML **	N018831 001	Nov 23, 1983
-----------	------------	-------------	--------------

ATROPOINE

INJECTABLE; INJECTION

ATROPOINE

ABBVIE	EQ 2MG SULFATE/0.7ML	A071295 001	Jan 30, 1987
--------	----------------------	-------------	--------------

ATROPOINE SULFATE

AEROSOL, METERED; INHALATION

ATROPOINE SULFATE

US ARMY	EQ 0.36MG BASE/INH	N020056 001	Sep 19, 1990
---------	--------------------	-------------	--------------

ATROPOINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET;ORAL

MOTOFEN HALF-STRENGTH

SEBELA IRELAND LTD	0.025MG;0.5MG	N017744 001	
--------------------	---------------	-------------	--

ATROPOINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE;ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPOINE SULFATE

SCHERER RP	0.025MG;2.5MG	A086440 001	
------------	---------------	-------------	--

SOLUTION;ORAL

COLONAID

MEDPOINTE PHARM HLC	0.025MG/5ML;2.5MG/5ML	A085735 001	
---------------------	-----------------------	-------------	--

LOMANATE

ALPHARMA US PHARMS	0.025MG/5ML;2.5MG/5ML	A085746 001	
--------------------	-----------------------	-------------	--

LOMOTIL

GD SEARLE LLC	0.025MG/5ML;2.5MG/5ML	N012699 001	
---------------	-----------------------	-------------	--

TABLET;ORAL

COLONAID

MEDPOINTE PHARM HLC	0.025MG;2.5MG	A085737 001	
---------------------	---------------	-------------	--

DI-ATRO

MD PHARM	0.025MG;2.5MG	A085266 001	
----------	---------------	-------------	--

DIPHENOXYLATE HYDROCHLORIDE AND ATROPOINE SULFATE

ABLE	0.025MG;2.5MG	A040395 001	Nov 27, 2000
------	---------------	-------------	--------------

ASCOT	0.025MG;2.5MG	A087934 001	Jul 19, 1983
-------	---------------	-------------	--------------

HEATHER	0.025MG;2.5MG	A086798 001	
---------	---------------	-------------	--

HIKMA PHARMS	0.025MG;2.5MG	A087765 001	Mar 15, 1982
--------------	---------------	-------------	--------------

INWOOD LABS	0.025MG;2.5MG	A085509 001	
-------------	---------------	-------------	--

KV PHARM	0.025MG;2.5MG	A085659 001	
----------	---------------	-------------	--

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-42(of 375)

\*\* See List Footnote

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET;ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

LEDERLE	0.025MG;2.5MG	A086950 001
PARKE DAVIS	0.025MG;2.5MG	A087131 001
PVT FORM	0.025MG;2.5MG	A085766 001
R AND S PHARMA	0.025MG;2.5MG	A085035 001
ROXANE	0.025MG;2.5MG	A086057 001
SANDOZ	0.025MG;2.5MG	A086173 001
SUN PHARM INDUSTRIES	0.025MG;2.5MG	A085506 001
USL PHARMA	0.025MG;2.5MG	A087842 001 Mar 29, 1982
VALEANT PHARM INTL	0.025MG;2.5MG	A087195 001 Feb 16, 1982
WATSON LABS	0.025MG;2.5MG	A085876 001
LO-TROL		
VANGARD	0.025MG;2.5MG	A088009 001 Mar 25, 1983
LOGEN		
SUPERPHARM	0.025MG;2.5MG	A088962 001 May 10, 1985
LONOX		
SANDOZ	0.025MG;2.5MG	A085311 002
LOW-QUEL		
HALSEY	0.025MG;2.5MG	A085211 001

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE;INJECTION

ENLON-PLUS

MYLAN INSTITUTIONAL	0.14MG/ML;10MG/ML	N019677 001 Nov 06, 1991
+	0.14MG/ML;10MG/ML	N019678 001 Nov 06, 1991

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE;INJECTION

ATROPINE AND DEMEROL

ABBVIE	0.4MG/ML;50MG/ML	A087853 001 Nov 26, 1982
	0.4MG/ML;75MG/ML	A087847 001 Nov 26, 1982
	0.4MG/ML;100MG/ML	A087848 001 Nov 26, 1982
MEPERIDINE AND ATROPINE SULFATE		
WYETH AYERST	0.4MG/ML;50MG/ML	A085121 001
	0.4MG/ML;75MG/ML	A085121 002
	0.4MG/ML;100MG/ML	A085121 003

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE;INTRAMUSCULAR

ATNAA

US ARMY	2.1MG/0.7ML;600MG/2ML	N021175 001 Jan 17, 2002
---------	-----------------------	--------------------------

AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION;TOPICAL

SHADE UVAGUARD

+ BAYER HEALTHCARE LLC	3%;7.5%;3%	N020045 001 Dec 07, 1992
------------------------	------------	--------------------------

AZATADINE MALEATE

TABLET;ORAL

OPTIMINE

SCHERING	1MG	N017601 001
----------	-----	-------------

AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

TRINALIN

SCHERING	1MG;120MG	N018506 001 Mar 23, 1982
----------	-----------	--------------------------

AZATHIOPRINE

TABLET;ORAL

IMURAN

+ SEBELA IRELAND LTD	25MG **	N016324 002
----------------------	---------	-------------

AZATHIOPRINE SODIUM

INJECTABLE;INJECTION

IMURAN

+ CASPER PHARMA LLC	EQ 100MG BASE/VIAL **	N017391 001
---------------------	-----------------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-43(of 375)

\*\* See List Footnote

AZELASTINE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC OPTIVAR + MYLAN SPECIALITY LP	0.05%	N021127 001 May 22, 2000
SPRAY, METERED;NASAL ASTEPRO MYLAN SPECIALITY LP	EQ 0.125MG BASE/SPRAY	N022203 001 Oct 15, 2008

AZITHROMYCIN

CAPSULE;ORAL ZITHROMAX + PFIZER	EQ 250MG BASE **	N050670 001 Nov 01, 1991
FOR SUSPENSION;ORAL AZITHROMYCIN SANDOZ	EQ 100MG BASE/5ML EQ 200MG BASE/5ML	A065297 001 Sep 18, 2006 A065297 002 Sep 18, 2006
INJECTABLE;INJECTION AZITHROMYCIN CSPC OUYI PHARM CO TEVA PARENTERAL	EQ 500MG BASE/VIAL EQ 500MG BASE/VIAL EQ 2.5GM BASE/VIAL	A065265 001 Jan 18, 2007 N050809 001 Dec 19, 2006 N050809 002 Dec 19, 2006
TABLET;ORAL AZITHROMYCIN MYLAN	EQ 250MG BASE EQ 500MG BASE	A065365 001 May 30, 2007 A065366 001 May 30, 2007

AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATE

FOR SUSPENSION, TABLET;ORAL TROVAN/ZITHROMAX COMPLIANCE PAK PFIZER	EQ 1GM BASE,N/A;N/A, EQ 100MG BASE	N050762 001 Dec 18, 1998
--	------------------------------------	--------------------------

AZLOCILLIN SODIUM

INJECTABLE;INJECTION AZLIN BAYER PHARMS	EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL EQ 3GM BASE/VIAL EQ 3GM BASE/VIAL EQ 3GM BASE/VIAL EQ 4GM BASE/VIAL EQ 4GM BASE/VIAL EQ 4GM BASE/VIAL	A062388 001 Sep 08, 1982 A062417 001 Oct 12, 1982 N050562 001 Sep 03, 1982 A062388 002 Sep 08, 1982 A062417 002 Oct 12, 1982 N050562 002 Sep 03, 1982 A062388 003 Sep 08, 1982 A062417 003 Oct 12, 1982 N050562 003 Sep 03, 1982
---	--	--

AZTREONAM

INJECTABLE;INJECTION AZACTAM BRISTOL MYERS SQUIBB 500MG/VIAL	N050580 001 Dec 31, 1986
AZACTAM IN PLASTIC CONTAINER BRISTOL MYERS SQUIBB 10MG/ML	N050632 003 May 24, 1989
AZTREONAM WEST-WARD PHARMS INT 1GM/VIAL 2GM/VIAL	A065286 001 Mar 23, 2011 A065286 002 Mar 23, 2011

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION;ORAL SPECTROBID PFIZER	125MG/5ML	N050556 001 Mar 23, 1982
TABLET;ORAL SPECTROBID PFIZER	400MG 800MG	N050520 001 N050520 002 Sep 12, 1983

BACITRACIN

INJECTABLE;INJECTION BACITRACIN PFIZER	50,000 UNITS/VIAL	A060282 001
OINTMENT;OPHTHALMIC BACIGUENT PHARMACIA AND UPJOHN	500 UNITS/GM	A060734 001
BACITRACIN LILLY	500 UNITS/GM	A060687 001

**DISCONTINUED DRUG PRODUCT LIST**

6-44(of 375)

\*\* See List Footnote

**BACITRACIN**

OINTMENT;OPHTHALMIC		
BACITRACIN		
PHARMADERM	500 UNITS/GM	A062158 001
PHARMAFAIR	500 UNITS/GM	A062453 001 Mar 28, 1984
OINTMENT;TOPICAL		
BACITRACIN		
COMBE	500 UNITS/GM	A062799 001 May 14, 1987
NASKA	500 UNITS/GM	A062857 001 Nov 13, 1987
POWDER;FOR RX COMPOUNDING		
BACI-RX		
X GEN PHARMS	5,000,000 UNITS/BOT	A061580 001
BACITRACIN		
APOTHEKERNES	5,000,000 UNITS/BOT	A061699 001
PADDOCK LLC	5,000,000 UNITS/BOT	A062456 001 Jul 27, 1983

**BACITRACIN ZINC**

POWDER;FOR RX COMPOUNDING		
ZIBA-RX		
X GEN PHARMS	500,000 UNITS/BOT	A061737 001

**BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE**

OINTMENT;OPHTHALMIC		
CORTISPORIN		
+ CASPER PHARMA LLC	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM **	N050416 002
ZINC BACITRACIN,NEOMYCIN SULFATE,POLYMYXIN B SULFATE & HYDROCORTISONE		
PHARMAFAIR	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062389 001 Jul 02, 1982
OINTMENT;TOPICAL		
NEOMYCIN & POLYMYXIN B SULFATES & BACITRACIN ZINC & HYDROCORTISONE		
PHARMAFAIR	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM	A062381 001 Sep 06, 1985

**BACITRACIN ZINC; LIDOCAINE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE**

OINTMENT;TOPICAL		
LANABIOTIC		
COMBE	400 UNITS/GM;40MG/GM;EQ 5MG BASE/GM;5,000 UNITS/GM	A062499 001 Jun 03, 1985

**BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE**

OINTMENT;OPHTHALMIC		
BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE		
PHARMAFAIR	400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A062386 001 Sep 09, 1982
BACITRACIN-NEOMYCIN-POLYMYXIN		
PHARMADERM	400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM	A062167 001
NEO-POLYCYN		
DOW PHARM	500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM	A060647 001
OINTMENT;TOPICAL		
BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE		
NASKA	400 UNITS/GM;EQ 3.5MG BASE/GM;5,000 UNITS/GM	A062833 001 Nov 09, 1987

**BACITRACIN ZINC; POLYMYXIN B SULFATE**

AEROSOL;TOPICAL		
POLYSPORIN		
GLAXOSMITHKLINE	10,000 UNITS/GM;2,000,000 UNITS/GM	N050167 002 Mar 01, 1985
OINTMENT;OPHTHALMIC		
OCUMYCIN		
PHARMAFAIR	500 UNITS/GM;10,000 UNITS/GM	A062430 001 Apr 08, 1983
POLYSPORIN		
MONARCH PHARMS	500 UNITS/GM;10,000 UNITS/GM **	A061229 001
OINTMENT;TOPICAL		
BACITRACIN ZINC-POLYMYXIN B SULFATE		
NASKA	500 UNITS/GM;10,000 UNITS/GM	A062849 001 Nov 13, 1987

**DISCONTINUED DRUG PRODUCT LIST**

6-45(of 375)

\*\* See List Footnote

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

ALTANA 400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM A060731 002

BACITRACIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT;OPHTHALMIC

MYCITRACIN

PHARMACIA AND UPJOHN 500 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM A061048 001

BACITRACIN; POLYMYXIN B SULFATE

DISC;TOPICAL

LANABIOTIC

COMBE 500 UNITS/GM;5,000 UNITS/GM N050598 001 Sep 22, 1986

BACLOFEN

TABLET;ORAL

BACLOFEN

MYLAN	10MG	A077181 001	Jul 29, 2005
TEVA	10MG	A073043 001	Feb 27, 1992
	20MG	A073044 001	Feb 27, 1992
USL PHARMA	10MG	A071260 001	May 06, 1988
	20MG	A071261 001	May 06, 1988
WATSON LABS	10MG	A072824 001	Sep 18, 1991
	10MG	A073092 001	Jan 28, 1994
	20MG	A074698 001	Aug 20, 1996
	20MG	A072825 001	Sep 18, 1991
	20MG	A073093 001	Jan 28, 1994
	20MG	A074698 002	Aug 20, 1996

LIORESAL

+ NOVARTIS 10MG \*\*  
+ 20MG \*\*N017851 001  
N017851 003 Jan 20, 1982

TABLET, ORALLY DISINTEGRATING;ORAL

KEMSTRO

UCB INC 10MG N021589 001 Oct 30, 2003  
20MG N021589 002 Oct 30, 2003BARIUM SULFATE

FOR SUSPENSION;ORAL

E-Z-CAT DRY

+ BRACCO 40% (9GM/POUCH)

N208036 003 Jan 03, 2017

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED;INHALATION

BECLOVENT

GLAXOSMITHKLINE 0.042MG/INH N018153 001

VANCERIL

SCHERING 0.042MG/INH N017573 001

VANCERIL DOUBLE STRENGTH

SCHERING 0.084MG/INH N020486 001 Dec 24, 1996

AEROSOL, METERED;NASAL

BECONASE

GLAXOSMITHKLINE 0.042MG/INH N018584 001

VANCENASE

SCHERING 0.042MG/INH N018521 001

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED;NASAL

VANCENASE AQ

SCHERING EQ 0.042MG DIPROP/SPRAY N019589 001 Dec 23, 1987  
EQ 0.084MG DIPROP/SPRAY N020469 001 Jun 26, 1996BENAZEPRIL HYDROCHLORIDE

TABLET;ORAL

BENAZEPRIL HYDROCHLORIDE

ACTAVIS LABS FL INC	5MG	A076267 001	Feb 11, 2004
	10MG	A076267 002	Feb 11, 2004
	20MG	A076267 003	Feb 11, 2004
	40MG	A076267 004	Feb 11, 2004
GENPHARM	5MG	A076476 001	Feb 11, 2004
	10MG	A076476 002	Feb 11, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-46(of 375)

\*\* See List Footnote

BENAZEPRIL HYDROCHLORIDE

TABLET;ORAL

BENAZEPRIL HYDROCHLORIDE

20MG	A076476 003	Feb 11, 2004
40MG	A076476 004	Feb 11, 2004

BENAZEPRIL HYDROCHLORIDE; HYDROCHLORTIAZIDE

TABLET;ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLORTIAZIDE

ACTAVIS LABS FL INC	5MG;6.25MG	A076342 001	Feb 11, 2004
	10MG;12.5MG	A076342 002	Feb 11, 2004
	20MG;12.5MG	A076342 003	Feb 11, 2004
	20MG;25MG	A076342 004	Feb 11, 2004
IVAX SUB TEVA PHARMS	5MG;6.25MG	A076348 001	Feb 11, 2004
	10MG;12.5MG	A076348 002	Feb 11, 2004
	20MG;12.5MG	A076348 003	Feb 11, 2004
	20MG;25MG	A076348 004	Feb 11, 2004
MYLAN PHARMS INC	5MG;6.25MG	A076612 001	Feb 11, 2004
	10MG;12.5MG	A076612 002	Feb 11, 2004
	20MG;12.5MG	A076612 003	Feb 11, 2004
	20MG;25MG	A076612 004	Feb 11, 2004
SUN PHARM INDs LTD	5MG;6.25MG	A077483 001	Sep 08, 2005
	10MG;12.5MG	A077483 002	Sep 08, 2005
	20MG;12.5MG	A077483 003	Sep 08, 2005
	20MG;25MG	A077483 004	Sep 08, 2005
LOTENSIN HCT			
+ US PHARMS HOLDINGS I	5MG;6.25MG	N020033 001	May 19, 1992

BENDROFLUMETHIAZIDE

TABLET;ORAL

NATURETIN-10

APOTHECON	10MG	N012164 003
-----------	------	-------------

NATURETIN-2.5

APOTHECON	2.5MG	N012164 001
-----------	-------	-------------

NATURETIN-5

APOTHECON	5MG	N012164 002
-----------	-----	-------------

BENDROFLUMETHIAZIDE; NADOLOL

TABLET;ORAL

NADOLOL AND BENDROFLUMETHIAZIDE

MYLAN	5MG;40MG	A078688 001	Feb 15, 2008
	5MG;80MG	A078688 002	Feb 15, 2008

BENOXINATE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

BENOXINATE HYDROCHLORIDE

SOLA BARNES HIND	0.4%	A084149 001
------------------	------	-------------

BENTIROMIDE

SOLUTION;ORAL

CHYMEX

SAVAGE LABS	500MG/7.5ML	N018366 001	Dec 29, 1983
-------------	-------------	-------------	--------------

BENZONATATE

CAPSULE;ORAL

BENZONATATE

NESHER PHARMS	100MG	A040795 001	Oct 31, 2007
	200MG	A040795 002	Oct 31, 2007

TESSALON

+ PFIZER	200MG **	N011210 003	Jun 25, 1999
----------	----------	-------------	--------------

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL;TOPICAL

BENZACLIN

VALEANT BERMUDA	5%;EQ 1% BASE	N050756 002	Apr 20, 2007
-----------------	---------------	-------------	--------------

BENZPHETAMINE HYDROCHLORIDE

TABLET;ORAL

BENZPHETAMINE HYDROCHLORIDE

EPIC PHARMA LLC	50MG	A040714 001	Oct 29, 2007
IMPAK LABS	50MG	A040845 001	Nov 18, 2008

DIDREX

+ PHARMACIA AND UPJOHN	25MG **	N012427 003
------------------------	---------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-47(of 375)

\*\* See List Footnote

BENZPHETAMINE HYDROCHLORIDE

TABLET;ORAL

DIDREX

+

50MG \*\*

N012427 002

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE;INJECTION

EMETE-CON

PFIZER

EQ 50MG BASE/VIAL

N016820 001

SUPPOSITORY;RECTAL

EMETE-CON

ROERIG

EQ 100MG BASE

N016818 006

BENZTHIAZIDE

TABLET;ORAL

AQUATAG

SOLVAY

25MG

N016001 001

50MG

N016001 002

BENZTHIAZIDE

PVT FORM

50MG

A083206 001

EXNA

AH ROBINS INC

50MG

N012489 001

FOVANE

PFIZER

50MG

N012128 002

URESE

PFIZER

25MG

N012128 003

BENZTROPINE MESYLATE

TABLET;ORAL

BENZTROPINE MESYLATE

ANDA REPOSITORY

1MG

A081264 001 Jan 23, 1992

2MG

A081265 001 Jan 23, 1992

LANNETT HOLDINGS INC

0.5MG \*\*

A088877 001 Apr 11, 1985

1MG \*\*

A088894 001 Apr 11, 1985

2MG \*\*

A088895 001 Apr 11, 1985

OXFORD PHARMS

2MG

A040706 001 Feb 14, 2008

QUANTUM PHARMICS

0.5MG

A088514 001 Jan 31, 1984

1MG

A088510 001 Jan 31, 1984

2MG

A088511 001 Jan 31, 1984

USL PHARMA

0.5MG

A089211 001 Jun 14, 1988

1MG

A089212 001 Jun 14, 1988

2MG

A089213 001 Jun 14, 1988

COGENTIN

+ MERCK

0.5MG \*\*

N009193 004

+ +

1MG \*\*

N009193 003

+ +

2MG \*\*

N009193 002

BENZYL BENZOATE

EMULSION;TOPICAL

BENZYL BENZOATE

LANNETT

50%

A084535 001

BEPRIDIL HYDROCHLORIDE

TABLET;ORAL

BEPADIN

MEDPOINTE PHARM HLC

200MG

N019001 001 Dec 28, 1990

300MG

N019001 002 Dec 28, 1990

400MG

N019001 003 Dec 28, 1990

VASCOR

JOHNSON AND JOHNSON

200MG

N019002 001 Dec 28, 1990

300MG

N019002 002 Dec 28, 1990

400MG

N019002 003 Dec 28, 1990

BETA CAROTENE

CAPSULE;ORAL

SOLATENE

ROCHE

30MG

N017589 001

**DISCONTINUED DRUG PRODUCT LIST**

6-48(of 375)

\*\* See List Footnote

BETAMETHASONE

CREAM;TOPICAL		
CELESTONE		
SCHERING	0.2%	N014762 001
SYRUP;ORAL		
CELESTONE		
MERCK SHARP DOHME	0.6MG/5ML	N014215 002
TABLET;ORAL		
CELESTONE		
SCHERING	0.6MG	N012657 003

BETAMETHASONE BENZOATE

CREAM;TOPICAL		
UTICORT		
PARKE DAVIS	0.025%	N016998 002
GEL;TOPICAL		
UTICORT		
PARKE DAVIS	0.025%	N017244 001
LOTION;TOPICAL		
UTICORT		
PARKE DAVIS	0.025%	N017528 001
OINTMENT;TOPICAL		
UTICORT		
PARKE DAVIS	0.025%	N018089 001

BETAMETHASONE DIPROPIONATE

CREAM;TOPICAL		
ALPHATREX		
SAVAGE LABS	EQ 0.05% BASE	N019138 001 Jun 26, 1984
BETAMETHASONE DIPROPIONATE		
PERRIGO NEW YORK	EQ 0.05% BASE	A072536 001 Jan 31, 1990
	EQ 0.05% BASE	A074579 001 Nov 26, 1997
PHARMADERM	EQ 0.05% BASE	N019136 001 Jun 26, 1984
TARO	EQ 0.05% BASE	A071143 001 Jun 17, 1987
TEVA	EQ 0.05% BASE	A071476 001 Aug 10, 1987
DIPROSONE		
SCHERING	EQ 0.05% BASE	N017536 001
CREAM, AUGMENTED;TOPICAL		
DIPROLENE		
SCHERING	EQ 0.05% BASE	N019408 001 Jan 31, 1986
DISC;TOPICAL		
DIPROSONE		
SCHERING	EQ 0.1% BASE	N017829 001
GEL, AUGMENTED;TOPICAL		
DIPROLENE		
SCHERING	EQ 0.05% BASE	N019408 002 Nov 22, 1991
LOTION;TOPICAL		
ALPHATREX		
SAVAGE LABS	EQ 0.05% BASE	A070273 001 Aug 12, 1985
BETAMETHASONE DIPROPIONATE		
ALPHARMA US PHARMS	EQ 0.05% BASE	A071085 001 Feb 03, 1987
G AND W LABS INC	EQ 0.05% BASE	A071882 001 Jun 06, 1988
PHARMADERM	EQ 0.05% BASE	A070274 001 Aug 12, 1985
TARO	EQ 0.05% BASE	A072276 001 Aug 24, 1988
	EQ 0.05% BASE	A074272 001 Sep 30, 1994
DIPROSONE		
+ SCHERING	EQ 0.05% BASE **	N017781 001
OINTMENT;TOPICAL		
ALPHATREX		
SAVAGE LABS	EQ 0.05% BASE	N019143 001 Sep 04, 1984
BETAMETHASONE DIPROPIONATE		
PERRIGO NEW YORK	EQ 0.05% BASE	A072526 001 Jan 31, 1990
PHARMADERM	EQ 0.05% BASE	N019140 001 Sep 04, 1984
TEVA	EQ 0.05% BASE	A071477 001 Aug 10, 1987
DIPROSONE		
SCHERING	EQ 0.05% BASE	N017691 001

**DISCONTINUED DRUG PRODUCT LIST**

6-49(of 375)

\*\* See List Footnote

BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

BETAMETHASONE SODIUM PHOSPHATE

WATSON LABS EQ 3MG BASE/ML  
CELESTONE  
+ SCHERING EQ 3MG BASE/ML \*\*

A085738 001

N017561 001

BETAMETHASONE VALERATE

CREAM; TOPICAL

BETADERM

ROACO EQ 0.1% BASE N018839 001 Jun 30, 1983  
BETAMETHASONE VALERATE PERRIGO NEW YORK EQ 0.1% BASE A070053 001 Jun 10, 1986  
PHARMADERM EQ 0.1% BASE N018860 002 Aug 31, 1983  
PHARMAFAIR EQ 0.1% BASE A070485 001 May 29, 1987  
TARO EQ 0.1% BASE A070062 001 May 14, 1985  
BETATREX SAVAGE LABS EQ 0.1% BASE N018862 001 Aug 31, 1983  
VALISONE SCHERING EQ 0.01% BASE N016322 002  
EQ 0.1% BASE N016322 001

LOTION; TOPICAL

BETAMETHASONE VALERATE PHARMADERM EQ 0.1% BASE N018870 001 Aug 31, 1983  
PHARMAFAIR EQ 0.1% BASE A070484 001 May 29, 1987  
TEVA PHARMS EQ 0.1% BASE A071883 001 Apr 22, 1988  
BETATREX SAVAGE LABS EQ 0.1% BASE N018867 001 Aug 31, 1983  
VALISONE SCHERING EQ 0.1% BASE N016932 001

OINTMENT; TOPICAL

BETAMETHASONE VALERATE PERRIGO NEW YORK EQ 0.1% BASE A071478 001 Dec 23, 1987  
PHARMADERM EQ 0.1% BASE N018864 001 Aug 31, 1983  
PHARMAFAIR EQ 0.1% BASE A070486 001 May 29, 1987  
BETATREX SAVAGE LABS EQ 0.1% BASE N018863 001 Aug 31, 1983  
VALISONE SCHERING EQ 0.1% BASE N016740 001BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

APOTEX INC EQ 0.5% BASE

A075446 001 Sep 28, 2000

TABLET; ORAL

KERLONE

SANOFI AVENTIS US 10MG  
20MGN019507 001 Oct 27, 1989  
N019507 002 Oct 27, 1989BETAXOLOL HYDROCHLORIDE; CHLORTHALIDONE

TABLET; ORAL

KERLEDEX

SANOFI AVENTIS US 5MG;12.5MG  
10MG;12.5MGN019807 001 Oct 30, 1992  
N019807 002 Oct 30, 1992BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC PILO

ALCON EQ 0.25% BASE;1.75%

N020619 001 Apr 17, 1997

BETAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

HISTALOG

LILLY 50MG/ML

N009344 001

BETHANECHOL CHLORIDE

INJECTABLE; INJECTION

URECHOLINE

+ ODYSSEY PHARMS 5MG/ML \*\*

N006536 001

TABLET; ORAL

BETHANECHOL CHLORIDE

ABLE 5MG

A040492 001 Jul 27, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-50(of 375)

\*\* See List Footnote

BETHANECHOL CHLORIDE

TABLET;ORAL

BETHANECHOL CHLORIDE

	10MG	A040483 001 Jul 27, 2004
	25MG	A040485 001 Jul 27, 2004
	50MG	A040509 001 Jul 27, 2004
ACTAVIS ELIZABETH	5MG	A040552 001 Oct 28, 2004
	10MG	A040553 001 Oct 28, 2004
	25MG	A040554 001 Oct 28, 2004
	50MG	A040551 001 Oct 28, 2004
ASCOT	10MG	A088288 001 Jun 08, 1983
	25MG	A088289 001 Jun 08, 1983
IMPAX LABS	5MG	A040721 001 Nov 01, 2006
	10MG	A040721 002 Nov 01, 2006
	25MG	A040721 003 Nov 01, 2016
	50MG	A040721 004 Nov 01, 2006
IVAX SUB TEVA PHARMS	25MG	A084689 001
LANNETT	5MG	A084702 001
	10MG	A084712 001
	25MG	A084074 001
SANDOZ	5MG	A084353 001
	10MG	A084378 001
	25MG	A084379 001
	25MG	A084383 001
	25MG	A084384 001
SUN PHARM INDS INC	5MG	A040897 001 Apr 22, 2009
	10MG	A040897 002 Apr 22, 2009
	25MG	A040897 003 Apr 22, 2009
	50MG	A040897 004 Apr 22, 2009
WATSON LABS	5MG	A084402 001
	5MG	A085230 002
	5MG	A085841 001
	10MG	A084408 001
	10MG	A085228 001
	10MG	A085842 001
	25MG	A084441 001
	25MG	A085229 001
	25MG	A085839 001
	50MG	A087397 001
	50MG	A087444 001
MYOTONACHOL		
GLENWOOD	5MG	A084188 001
	10MG	A084188 003
	25MG	A084188 004
URECHOLINE		
+ ODYSSEY PHARMS	5MG **	N006536 003
+	10MG **	N006536 002
+	25MG **	N006536 004
+	50MG **	N006536 005

BETHANIDINE SULFATE

TABLET;ORAL

TENATHAN

ROBINS AH

10MG

N017675 001

25MG

N017675 002

BICALUTAMIDE

TABLET;ORAL

BICALUTAMIDE

KUDCO IRELAND

50MG

A077995 001 Jul 06, 2009

ROXANE

50MG

A078285 001 Mar 24, 2011

SYNTHON PHARMS

50MG

A077973 001 Jul 06, 2009

BIMATOPROST

SOLUTION/DROPS;OPHTHALMIC

LUMIGAN

+ ALLERGAN

0.03% \*\*

N021275 001 Mar 16, 2001

**DISCONTINUED DRUG PRODUCT LIST**

6-51(of 375)

\*\* See List Footnote

BIPERIDEN HYDROCHLORIDE

TABLET;ORAL

AKINETON

ABBVIE

2MG

N012003 001

BIPERIDEN LACTATE

INJECTABLE;INJECTION

AKINETON

ABBVIE

5MG/ML

N012418 002

BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION, TABLET, DELAYED RELEASE;ORAL

HALFLYTLY

+ BRAINTREE

5MG,N/A;N/A,210GM;N/A,0.74GM;N/A,2.86GM N021551 003 Jul 16, 2010  
;N/A,5.6GM \*\*BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE;ORAL

HELDAC

+ CASPER PHARMA LLC

262.4MG,N/A,N/A;N/A,250MG,N/A;N/A,N/A,5 N050719 001 Aug 15, 1996  
00MG \*\*BISOPROLOL FUMARATE

TABLET;ORAL

BISOPROLOL FUMARATE

ANDA REPOSITORY

5MG

A075474 001 Oct 25, 2002

10MG

A075474 002 Oct 25, 2002

ZEBETA

TEVA WOMENS

5MG \*\*

N019982 002 Jul 31, 1992

10MG \*\*

N019982 001 Jul 31, 1992

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

ACTAVIS ELIZABETH

2.5MG;6.25MG

A075672 001 Sep 25, 2000

5MG;6.25MG

A075672 002 Sep 25, 2000

10MG;6.25MG

A075672 003 Sep 25, 2000

APOTHECON

2.5MG;6.25MG

A075642 002 Dec 27, 2000

5MG;6.25MG

A075642 001 Dec 27, 2000

10MG;6.25MG

A075642 003 Dec 27, 2000

IVAX SUB TEVA PHARMS

2.5MG;6.25MG

A075632 001 Sep 27, 2000

5MG;6.25MG

A075632 002 Sep 27, 2000

10MG;6.25MG

A075632 003 Sep 27, 2000

SANDOZ

2.5MG;6.25MG

A075527 001 Sep 25, 2000

5MG;6.25MG

A075527 003 Sep 25, 2000

10MG;6.25MG

A075527 002 Sep 25, 2000

TEVA

2.5MG;6.25MG

A075686 001 Jan 19, 2001

5MG;6.25MG

A075686 002 Jan 19, 2001

10MG;6.25MG

A075686 003 Jan 19, 2001

WATSON LABS TEVA

2.5MG;6.25MG

A075469 001 Sep 25, 2000

5MG;6.25MG

A075469 002 Sep 25, 2000

10MG;6.25MG

A075469 003 Sep 25, 2000

BITOLTEROL MESYLATE

AEROSOL, METERED;INHALATION

TORNALATE

SANOFI AVENTIS US

0.37MG/INH

N018770 001 Dec 28, 1984

SOLUTION;INHALATION

TORNALATE

SANOFI AVENTIS US

0.2%

N019548 001 Feb 19, 1992

BLEOMYCIN SULFATE

INJECTABLE;INJECTION

BLENOXANE

+ BRISTOL MYERS SQUIBB

EQ 15 UNITS BASE/VIAL \*\*

N050443 001

+

EQ 30 UNITS BASE/VIAL \*\*

N050443 002 Sep 07, 1995

BLEOMYCIN SULFATE

PHARMACHEMIE BV

EQ 15 UNITS BASE/VIAL

A065201 001 Dec 13, 2007

TEVA PARENTERAL

EQ 15 UNITS BASE/VIAL

A064084 001 Jun 01, 1996

EQ 30 UNITS BASE/VIAL

A064084 002 Jun 01, 1996

**DISCONTINUED DRUG PRODUCT LIST**

6-52(of 375)

\*\* See List Footnote

BOCEPREVIRCAPSULE;ORAL  
VICTRELIS

MERCK SHARP DOHME 200MG

N202258 001 May 13, 2011

BRETYLIUM TOSYLATE

INJECTABLE;INJECTION

BRETYLIUM TOSYLATE

ABRAXIS PHARM	50MG/ML	A070134 001	Apr 29, 1986
	100MG/ML	A071298 001	Feb 13, 1987
ASTRAZENECA	50MG/ML	A071151 001	Aug 10, 1987
	50MG/ML	A071152 001	Aug 10, 1987
	50MG/ML	A071153 001	Aug 10, 1987
EUROHLTH INTL SARL	50MG/ML	A070546 001	May 14, 1986
+ HOSPIRA	50MG/ML **	N019030 001	Apr 29, 1986
	50MG/ML	N019033 001	Apr 29, 1986
INTL MEDICATION	50MG/ML	A070119 001	Apr 29, 1986
LUITPOLD	50MG/ML	A070891 001	Jul 26, 1988
WEST-WARD PHARMS INT	50MG/ML	A070545 001	May 14, 1986
BRETYLIUM TOSYLATE IN DEXTROSE 5%			
ABBOTT	200MG/100ML	N019005 002	Apr 29, 1986
	400MG/100ML	N019005 003	Apr 29, 1986
	800MG/100ML	N019005 001	Apr 29, 1986
BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	100MG/100ML	N019121 001	Apr 29, 1986
	200MG/100ML	N019121 002	Apr 29, 1986
	400MG/100ML	N019121 003	Apr 29, 1986
BAXTER HLTHCARE	200MG/100ML	N019837 002	Apr 12, 1989
	400MG/100ML	N019837 001	Apr 12, 1989
HOSPIRA INC	200MG/100ML	N019008 002	Apr 29, 1986
	400MG/100ML	N019008 003	Apr 29, 1986
	800MG/100ML	N019008 001	Apr 29, 1986
BRETYLOL			
HOSPIRA	50MG/ML	N017954 001	

BRIMONIDINE TARTRATE

SOLUTION/DROPS;OPHTHALMIC

ALPHAGAN

+ ALLERGAN	0.2% **	N020613 001	Sep 06, 1996
	0.5%	N020490 001	Mar 13, 1997

BRIMONIDINE TARTRATE

TEVA PARENTERAL 0.2%

A076372 001 Sep 10, 2004

BROMFENAC SODIUM

SOLUTION/DROPS;OPHTHALMIC

BROMDAY

+ BAUSCH AND LOMB INC	EQ 0.09% ACID **	N021664 002	Oct 16, 2010
BROMFENAC SODIUM		A201211 001	May 11, 2011
COASTAL PHARMS	EQ 0.09% ACID		

XIBROM

+ BAUSCH AND LOMB INC	EQ 0.09% ACID **	N021664 001	Mar 24, 2005
-----------------------	------------------	-------------	--------------

BROMOCRIPTINE MESYLATE

CAPSULE;ORAL

BROMOCRIPTINE MESYLATE

LEK PHARM EQ 5MG BASE

A075100 001 Dec 10, 1998

BROMODIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE;ORAL

AMBODRYL

PARKE DAVIS 25MG

N007984 001

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP;ORAL

AMBENYL

FOREST LABS 12.5MG/5ML;10MG/5ML

N009319 006 Jan 10, 1984

BROMANYL

ALPHARMA US PHARMS 12.5MG/5ML;10MG/5ML

A088343 001 Aug 15, 1984

BROMODIPHENHYDRAMINE HYDROCHLORIDE AND CODEINE PHOSPHATE

WOCKHARDT 12.5MG/5ML;10MG/5ML

A088626 001 Oct 12, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-53(of 375)

\*\* See List Footnote

BROMPHENIRAMINE MALEATE

ELIXIR;ORAL

BROMPHENIRAMINE MALEATE

ALPHARMA US PHARMS

2MG/5ML

A086936 001

KV PHARM

2MG/5ML

A085466 001

PHARM ASSOC

2MG/5ML

A087517 001

USL PHARMA

2MG/5ML

A087964 001 Jan 25, 1983

INJECTABLE;INJECTION

BROMPHENIRAMINE MALEATE

WATSON LABS

10MG/ML

A083821 001

100MG/ML

A083820 001

DIMETANE-TEN

WYETH AYERST

10MG/ML

N011418 002

TABLET;ORAL

BROMPHENIRAMINE MALEATE

BARR

4MG

A084468 001

IVAX SUB TEVA PHARMS

4MG

A084351 001

NEWTRON PHARMS

4MG

A086987 001

NEXGEN PHARMA INC

4MG

A086187 001

PAR PHARM

4MG

A087009 001

PIONEER PHARMS

4MG

A088604 001 Jul 13, 1984

UPSHER-SMITH LABS

4MG

A083215 001

VITARINE

4MG

A085850 001

WATSON LABS

4MG

A083123 001

4MG

A085769 001

DIMETANE

WYETH CONS

4MG

N010799 003

TABLET, EXTENDED RELEASE;ORAL

DIMETANE

WYETH CONS

8MG

N010799 010 Jun 10, 1983

12MG

N010799 011 Jun 10, 1983

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP;ORAL

BROMANATE DM

ALPHARMA US PHARMS

2MG/5ML;10MG/5ML;30MG/5ML

A088722 001 Mar 07, 1985

BROMFED-DM

WOCKHARDT

2MG/5ML;10MG/5ML;30MG/5ML

A089681 001 Dec 22, 1988

DIMETANE-DX

+ ROBINS AH

2MG/5ML;10MG/5ML;30MG/5ML \*\*

N019279 001 Aug 24, 1984

BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

ALZA

16MG;240MG

N019672 001 Mar 29, 1996

BUCLIZINE HYDROCHLORIDE

TABLET;ORAL

BUCLADIN-S

STUART PHARMS

50MG

N010911 006

BUDESONIDE

AEROSOL, METERED;NASAL

RHINOCORT

ASTRAZENECA

0.032MG/INH

N020233 001 Feb 14, 1994

POWDER, METERED;INHALATION

PULMICORT

ASTRAZENECA

0.16MG/INH

N020441 002 Jun 24, 1997

0.32MG/INH

N020441 003 Jun 24, 1997

BUMETANIDE

INJECTABLE;INJECTION

BUMETANIDE

HOSPIRA

0.25MG/ML

A074160 001 Oct 30, 1997

TEVA PARENTERAL

0.25MG/ML

A074613 001 Nov 18, 1997

BUMEX

+ VALIDUS PHARMS

0.25MG/ML \*\*

N018226 001 Feb 28, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-54(of 375)

\*\* See List Footnote

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

## BUPIVACAINE HYDROCHLORIDE

HOSPIRA	0.75%	A070587 001 Mar 03, 1987
BUPIVACAINE HYDROCHLORIDE KIT		
HOSPIRA	0.075%	N019978 001 Sep 03, 1992
	0.114%	N019978 002 Sep 03, 1992
	0.23%	N019978 003 Sep 03, 1992
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE		
INTL MEDICATED	0.25%	A076012 001 Jan 09, 2002
	0.5%	A076012 002 Jan 09, 2002
	0.75%	A076012 003 Jan 09, 2002

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

## BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

HOSPIRA	0.25%;0.005MG/ML	A071166 001 Jun 16, 1988
	0.5%;0.005MG/ML	A071169 001 Jun 16, 1988
	0.75%;0.005MG/ML	A071171 001 Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

## DUOCAINE

AMPHASTAR PHARMS INC	EQ 0.375% (37.5MG/10ML);EQ 1% (100MG/10ML)	N021496 001 May 23, 2003
----------------------	---	--------------------------

BUPRENORPHINE HYDROCHLORIDE

TABLET; SUBLINGUAL

## SUBUTEX

+ INDIVIOR INC	EQ 2MG BASE **	N020732 002 Oct 08, 2002
+	EQ 8MG BASE **	N020732 003 Oct 08, 2002

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE

TABLET; SUBLINGUAL

## SUBOXONE

+ INDIVIOR INC	EQ 2MG BASE;EQ 0.5MG BASE **	N020733 001 Oct 08, 2002
+	EQ 8MG BASE;EQ 2MG BASE	N020733 002 Oct 08, 2002

BUPROPION HYDROCHLORIDE

TABLET; ORAL

## BUPROPION HYDROCHLORIDE

SANDOZ	75MG	A075613 002 Oct 10, 2000
	100MG	A075613 001 Oct 10, 2000
TEVA	75MG	A075310 001 Nov 29, 1999
	100MG	A075310 002 Nov 29, 1999
WELLBUTRIN		
+ GLAXOSMITHKLINE	50MG **	N018644 001 Dec 30, 1985
+	75MG **	N018644 002 Dec 30, 1985
+	100MG **	N018644 003 Dec 30, 1985

TABLET, EXTENDED RELEASE; ORAL

## BUPROPION HYDROCHLORIDE

ACTAVIS LABS FL INC	300MG	A077715 002 Jun 13, 2007
IMPAX LABS	300MG	A077415 002 Dec 15, 2006
SANDOZ	100MG	A076845 001 Jul 14, 2005
	150MG	A076834 001 Jul 14, 2005
	150MG	A076845 002 Jul 14, 2005
WOCKHARDT LTD	100MG	A201331 001 Aug 30, 2012
	150MG	A201331 002 Aug 30, 2012
	200MG	A201331 003 Aug 30, 2012

## WELLBUTRIN SR

GLAXOSMITHKLINE	50MG	N020358 001 Oct 04, 1996
-----------------	------	--------------------------

## ZYBAN

GLAXOSMITHKLINE	100MG	N020711 002 May 14, 1997
-----------------	-------	--------------------------

BUSPIRONE HYDROCHLORIDE

CAPSULE; ORAL

## BUSPAR

BRISTOL MYERS SQUIBB	5MG	N021190 001 Dec 20, 2000
	7.5MG	N021190 002 Dec 20, 2000
	10MG	N021190 003 Dec 20, 2000
	15MG	N021190 004 Dec 20, 2000

**DISCONTINUED DRUG PRODUCT LIST**

6-55(of 375)

\*\* See List Footnote

BUSPIRONE HYDROCHLORIDE

## TABLET;ORAL

## BUPSPAR

+ BRISTOL MYERS SQUIBB	5MG **	N018731 001	Sep 29, 1986
+	10MG **	N018731 002	Sep 29, 1986
+	15MG **	N018731 003	Apr 22, 1996
+	30MG **	N018731 004	Apr 22, 1996
<b>BUSPIRONE HYDROCHLORIDE</b>			
APOTEX	5MG	A075521 001	Apr 05, 2002
	10MG	A075521 002	Apr 05, 2002
	15MG	A075521 003	Apr 05, 2002
EGIS	5MG	A075119 001	Mar 14, 2002
	10MG	A075119 002	Mar 14, 2002
	15MG	A075119 003	Jan 23, 2003
IVAX SUB TEVA PHARMS	5MG **	A075385 001	Mar 01, 2002
	10MG **	A075385 002	Mar 01, 2002
	15MG **	A075385 003	Mar 01, 2002
MYLAN	5MG	A075467 001	Feb 28, 2002
	10MG	A075467 003	Feb 28, 2002
	15MG	A075467 004	Feb 28, 2002
NESHER PHARMS	5MG	A075572 001	Feb 27, 2002
	10MG	A075572 002	Feb 27, 2002
	15MG	A075572 003	Feb 27, 2002
OXFORD PHARMS	5MG	A075388 001	May 09, 2002
	10MG	A075388 002	May 09, 2002
	15MG	A075388 003	May 09, 2002
SANDOZ	5MG	A075413 001	Mar 19, 2002
	10MG	A075413 002	Mar 19, 2002
	15MG	A075413 003	Mar 19, 2002

BUTABARBITAL SODIUM

## CAPSULE;ORAL

## BUTICAPS

MEDPOINTE PHARM HLC	15MG	A085381 001
	30MG	A085381 002
	50MG	A085381 003
	100MG	A085381 004

## ELIXIR;ORAL

## BUTABARB

ALPHARMA US PHARMS	30MG/5ML	A085873 001
--------------------	----------	-------------

## BUTABARBITAL SODIUM

WOCKHARDT	30MG/5ML	A085383 001
-----------	----------	-------------

## BUTALAN

LANNETT	33.3MG/5ML	A085880 001
---------	------------	-------------

## BUTISOL SODIUM

MEDA PHARMS	30MG/5ML	A085380 001
-------------	----------	-------------

## SARISOL

HALSEY	30MG/5ML	A084723 001
--------	----------	-------------

## TABLET;ORAL

## BUTABARBITAL

BUNDY	30MG	A085550 001
-------	------	-------------

## BUTABARBITAL SODIUM

SANDOZ	15MG	A084292 003 Feb 09, 1982
	15MG	A085938 001
	30MG	A084272 002
	30MG	A085934 001

SOLVAY	16.2MG	A083606 001
	32.4MG	A083898 001
	48.6MG	A083897 001
	97.2MG	A083896 001

TEVA	15MG	A088632 001 May 18, 1985
	30MG	A088631 001 May 01, 1985

WATSON LABS	15MG	A085764 001
	30MG	A085772 001

WHITEWORTH TOWN PLSN	15MG	A083325 002
	30MG	A083337 001

## BUTISOL SODIUM

MYLAN SPECIALITY LP	15MG	N000793 002
	50MG	N000793 003
	100MG	N000793 005

**DISCONTINUED DRUG PRODUCT LIST**

6-56(of 375)

\*\* See List Footnote

BUTABARBITAL SODIUM

TABLET;ORAL

SARISOL NO. 1 HALSEY	15MG	A084719 001
SARISOL NO. 2 HALSEY	30MG	A084719 002
SODIUM BUTABARBITAL HIKMA PHARMS	15MG	A085418 001
	30MG	A085432 001
IVAX SUB TEVA PHARMS	15MG	A083484 001
	30MG	A084040 001
LANNETT	15MG	A085849 001
	30MG	A085866 001
	100MG	A085881 001
MARSHALL PHARMA	16.2MG	A083524 001
	32.4MG	A083858 001

BUTENAFINE HYDROCHLORIDE

CREAM;TOPICAL

MENTAX-TC

MYLAN

1%

N021408 001 Oct 17, 2002

BUTOCONAZOLE NITRATE

CREAM;VAGINAL

BUTOCONAZOLE NITRATE

PERRIGO PHARMA INTL

2%

N019881 001 Feb 07, 1997

FEMSTAT

ROCHE PALO

2%

N019215 001 Nov 25, 1985

FEMSTAT 3

+ BAYER

2%

N020421 001 Dec 21, 1995

SUPPOSITORY;VAGINAL

FEMSTAT

ROCHE PALO

100MG

N019359 001 Nov 25, 1985

BUTORPHANOL TARTRATE

INJECTABLE;INJECTION

BUTORPHANOL TARTRATE

BAXTER HLTHCARE CORP

2MG/ML

A075697 001 Oct 23, 2001

HIKMA FARMACEUTICA

2MG/ML

A078247 001 Apr 29, 2009

HOSPIRA

1MG/ML

A075342 001 Nov 04, 1999

1MG/ML

A075559 001 Mar 20, 2000

2MG/ML

A075342 002 Nov 04, 1999

2MG/ML

A075559 002 Mar 20, 2000

BUTORPHANOL TARTRATE PRESERVATIVE FREE

BAXTER HLTHCARE CORP

1MG/ML

A075695 001 Oct 23, 2001

2MG/ML

A075695 002 Oct 23, 2001

HOSPIRA

1MG/ML

A074620 001 Jan 22, 1997

1MG/ML

A075170 001 Sep 28, 1998

2MG/ML

A074620 002 Jan 22, 1997

2MG/ML

A075170 002 Sep 28, 1998

STADOL

APOTHECON

2MG/ML \*\*

N017857 004

STADOL PRESERVATIVE FREE

APOTHECON

1MG/ML \*\*

N017857 001

2MG/ML \*\*

N017857 002

SPRAY, METERED;NASAL

STADOL

BRISTOL MYERS SQUIBB

1MG/SPRAY \*\*

N019890 001 Dec 12, 1991

CABERGOLINE

TABLET;ORAL

CABERGOLINE

IMPAX LABS INC

0.5MG

A077843 001 Jul 03, 2007

DOSTINEX

+ PHARMACIA AND UPJOHN

0.5MG \*\*

N020664 001 Dec 23, 1996

**DISCONTINUED DRUG PRODUCT LIST**

6-57(of 375)

\*\* See List Footnote

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY;RECTAL

CAFERGOT

+ NOVARTIS

100MG;2MG \*\*

N009000 002

TABLET;ORAL

CAFERGOT

NOVARTIS

100MG;1MG

N006620 001

WIGRAINE

ORGANON USA INC

100MG;1MG

A086562 001

CALCIFEDIOL

CAPSULE;ORAL

CALDEROL

ORGANON USA INC

0.02MG

N018312 001

0.05MG

N018312 002

CALCIPOTRIENE

OINTMENT;TOPICAL

DOVONEX

+ LEO PHARMA AS

0.005% \*\*

N020273 001 Dec 29, 1993

SOLUTION;TOPICAL

DOVONEX

+ LEO PHARM

0.005% \*\*

N020611 001 Mar 03, 1997

CALCITONIN HUMAN

INJECTABLE;INJECTION

CIBACALCIN

NOVARTIS

0.5MG/VIAL

N018470 001 Oct 31, 1986

CALCITONIN SALMON

INJECTABLE;INJECTION

CALCIMAR

SANOFI AVENTIS US

200 IU/ML

N017769 001

400 IU/VIAL

N017497 001

CALCITONIN-SALMON

IGI LABS INC

200 IU/ML

A073690 001 Apr 14, 1995

MIACALCIN

MYLAN IRELAND LTD

100 IU/ML

N017808 001 Jul 03, 1986

SPRAY, METERED;NASAL

MIACALCIN

+ MYLAN IRELAND LTD

200 IU/SPRAY

N020313 002 Aug 17, 1995

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED;NASAL

FORTICAL

UPSHER-SMITH LABS

200 IU/SPRAY

N021406 001 Aug 12, 2005

CALCITRIOL

INJECTABLE;INJECTION

CALCIJEX

+ ABBVIE

0.001MG/ML \*\*

N018874 001 Sep 25, 1986

+

0.002MG/ML \*\*

N018874 002 Sep 25, 1986

CALCITRIOL

AKORN

0.002MG/ML

A078066 002 Jan 29, 2008

FRESENIUS KABI USA

0.001MG/ML

A075836 001 Dec 31, 2002

0.002MG/ML

A075836 002 Dec 31, 2002

FRESENIUS MEDCL

0.001MG/ML

A075766 001 Feb 20, 2003

0.002MG/ML

A075766 002 Feb 20, 2003

HOSPIRA

0.001MG/ML

A075816 001 Jan 16, 2004

0.002MG/ML

A075816 002 Jan 16, 2004

LUITPOLD

0.001MG/ML

A075746 001 Sep 26, 2003

0.002MG/ML

A075746 002 Sep 26, 2003

ROCKWELL MEDCL

0.001MG/ML

A076206 001 Sep 17, 2003

SAGENT PHARMS

0.001MG/ML

A077102 001 Feb 08, 2006

TEVA PARENTERAL

0.001MG/ML

A075823 001 Mar 31, 2003

0.002MG/ML

A075823 002 Mar 31, 2003

**DISCONTINUED DRUG PRODUCT LIST**

6-58(of 375)

\*\* See List Footnote

CALCIUM ACETATECAPSULE;ORAL  
PHOSLOFRESENIUS MEDCL 333.5MG N021160 001 Apr 02, 2001  
667MG N021160 002 Apr 02, 2001

TABLET;ORAL

CALCIUM ACETATE  
WEST-WARD PHARMS INT 667MG A077693 001 Jan 30, 2008  
PHOSLO  
+ FRESENIUS MEDCL 667MG \*\* N019976 001 Dec 10, 1990CALCIUM CARBONATE; RISEDRONATE SODIUMTABLET, TABLET;ORAL  
ACTONEL WITH CALCIUM (COPACKAGED)  
+ WARNER CHILCOTT EQ 500MG BASE,N/A;N/A,35MG \*\* N021823 001 Aug 12, 2005CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

PRISMASOL B22GK 2/2.5 IN PLASTIC CONTAINER  
BAXTER HLTHCARE CORP 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;  
3.05GM/1000ML;0.157GM/1000ML;2.21GM/100  
0ML;7.07GM/1000ML (5000ML) N021703 012 Oct 10, 2008PRISMASOL BGK 4/0 IN PLASTIC CONTAINER  
BAXTER HLTHCARE CORP N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.0  
5GM/1000ML;0.314GM/1000ML;3.09GM/1000ML  
;6.46GM/1000ML (5000ML) N021703 005 Oct 25, 2006PRISMASOL BGK 4/3.5 IN PLASTIC CONTAINER  
BAXTER HLTHCARE CORP 5.15GM/1000ML;20GM/1000ML;5.4GM/1000ML;  
2.03GM/1000ML;0.314GM/1000ML;3.09GM/100  
0ML;6.46GM/1000ML (5000ML) N021703 008 Oct 25, 2006PRISMASOL BK 0/0 IN PLASTIC CONTAINER  
BAXTER HLTHCARE CORP N/A/1000ML;N/A/1000ML;5.4GM/1000ML;3.05  
GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46  
GM/1000ML (5000ML) N021703 007 Oct 25, 2006PRISMASOL BK 4/2.5 IN PLASTIC CONTAINER  
BAXTER HLTHCARE CORP 3.68GM/1000ML;N/A/1000ML;5.4GM/1000ML;3  
.05GM/1000ML;0.314GM/1000ML;3.09GM/1000  
ML;6.46GM/1000ML (5000ML) N021703 009 Oct 25, 2006CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION;IRRIGATION

NAVSTEL  
ALCON PHARMS LTD 0.154MG/ML;0.92MG/ML;0.2MG/ML;0.184MG/M  
L;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/M  
L N022193 001 Jul 24, 2008CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDEINJECTABLE;INJECTION  
ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER  
B BRAUN 37MG/100ML;5GM/100ML;31MG/100ML;120MG/1  
00ML;330MG/100ML;88MG/100ML N019864 001 Jun 10, 1993ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER  
B BRAUN 37MG/100ML;5GM/100ML;31MG/100ML;120MG/1  
00ML;330MG/100ML;88MG/100ML N018271 001CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATEINJECTABLE;INJECTION  
ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER  
B BRAUN 35MG/100ML;5GM/100ML;30MG/100ML;74MG/10  
0ML;640MG/100ML;500MG/100ML;74MG/100ML N019867 001 Dec 20, 1993ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER  
B BRAUN 35MG/100ML;5GM/100ML;30MG/100ML;74MG/10  
0ML;640MG/100ML;500MG/100ML;74MG/100ML N018269 002 Jan 17, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-59(of 375)

\*\* See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

## INJECTABLE; INJECTION

PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER  
 BAXTER HLTHCARE 37MG/100ML;5GM/100ML;30MG/100ML;119MG/100ML;161MG/100ML;94MG/100ML;138MG/100ML N017390 001

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

## SOLUTION; INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER  
 B BRAUN 510MG/100ML;30GM/100ML;200MG/100ML;9.2G M/100ML;9.6GM/100ML N018807 001 Aug 26, 1983  
 510MG/100ML;30GM/100ML;200MG/100ML;9.4G M/100ML;11GM/100ML N018807 003 Aug 26, 1983

DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER  
 B BRAUN 510MG/100ML;50GM/100ML;200MG/100ML;9.2G M/100ML;9.6GM/100ML N018807 002 Aug 26, 1983  
 510MG/100ML;50GM/100ML;200MG/100ML;9.4G M/100ML;11GM/100ML N018807 004 Aug 26, 1983

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER  
 B BRAUN 29MG/100ML;2.5GM/100ML;15MG/100ML;610MG /100ML;560MG/100ML N018460 006 Jan 29, 1986

DIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER  
 B BRAUN 29MG/100ML;1.5GM/100ML;15MG/100ML;610MG /100ML;560MG/100ML N018460 001

DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER  
 B BRAUN 29MG/100ML;4.25GM/100ML;15MG/100ML;610M G/100ML;560MG/100ML N018460 003

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

## SOLUTION; INTRAPERITONEAL

DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;1.5GM/100ML;15.2MG/100ML;5 67MG/100ML;392MG/100ML N018379 002

DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;2.5GM/100ML;15.2MG/100ML;5 67MG/100ML;392MG/100ML N018379 003

DELFLEX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;3.5GM/100ML;15.2MG/100ML;5 67MG/100ML;392MG/100ML N018379 007 Jun 24, 1988

DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;4.25GM/100ML;15.2MG/100ML;5 567MG/100ML;392MG/100ML N018379 001

DELFLEX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;1.5GM/100ML;5.08MG/100ML;5 38MG/100ML;448MG/100ML N018379 004 Jul 07, 1982

DELFLEX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5 38MG/100ML;448MG/100ML N018379 005 Jul 07, 1982

DELFLEX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5 38MG/100ML;448MG/100ML N018379 008 Jun 24, 1988

DELFLEX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER  
 FRESENIUS MEDCL 25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;5 538MG/100ML;448MG/100ML N018379 006 Jul 07, 1982

DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER  
 B BRAUN 26MG/100ML;1.5GM/100ML;5MG/100ML;530MG/ 100ML;450MG/100ML N018460 007 Jan 29, 1986  
 26MG/100ML;1.5GM/100ML;15MG/100ML;560MG /100ML;390MG/100ML N018460 002

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER  
 B BRAUN 26MG/100ML;2.5GM/100ML;5MG/100ML;530MG/ 100ML;450MG/100ML N018460 005 Nov 02, 1983  
 26MG/100ML;5GM/100ML;5MG/100ML;530MG/100ML;450MG/100ML N018460 008 Jan 29, 1986

DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER  
 B BRAUN 26MG/100ML;4.25GM/100ML;5MG/100ML;530MG /100ML;450MG/100ML N018460 009 Jan 29, 1986  
 26MG/100ML;4.25GM/100ML;15MG/100ML;560M G/100ML;390MG/100ML N018460 004

INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER  
 FRESENIUS 18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5 38MG/100ML;448MG/100ML A020374 001 Jun 13, 1994

**DISCONTINUED DRUG PRODUCT LIST**

6-60(of 375)

\*\* See List Footnote

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER FRESENIUS	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	A020374 002	Jun 13, 1994
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER FRESENIUS	18.4MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 5 38MG/100ML; 448MG/100ML	A020374 003	Jun 13, 1994
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER FRESENIUS	18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML	A020374 004	Jun 13, 1994

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER B BRAUN	20MG/100ML; 5GM/100ML; 30MG/100ML; 380MG/1 00ML; 600MG/100ML	N018258 001
--	---	-------------

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER HOSPIRA	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/1 00ML	N018254 001
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER B BRAUN	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/1 00ML	N018256 001
BAXTER HLTHCARE	33MG/100ML; 5GM/100ML; 30MG/100ML; 860MG/1 00ML	N016695 001

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER B BRAUN	4MG/100ML; 4GM/100ML; 6MG/100ML; 120MG/100 ML; 62MG/100ML	N019634 002	Feb 24, 1988
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER B BRAUN	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/1 00ML; 310MG/100ML	N017510 001	
MILES	20MG/100ML; 5GM/100ML; 30MG/100ML; 600MG/1 00ML; 310MG/100ML	N018499 001	
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/ 100ML; 310MG/100ML	N019685 005	Oct 17, 1988
	20MG/100ML; 5GM/100ML; 179MG/100ML; 600MG/ 100ML; 310MG/100ML	N019685 006	Oct 17, 1988
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/ 100ML; 310MG/100ML	N019685 007	Oct 17, 1988
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 254MG/100ML; 600MG/ 100ML; 310MG/100ML	N019685 003	Oct 17, 1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER ICU MEDICAL INC	20MG/100ML; 5GM/100ML; 104MG/100ML; 600MG/ 100ML; 310MG/100ML	N019685 001	Oct 17, 1988

CALCIUM CHLORIDE; DEXTROSE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

INPERSOL-ZM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 538MG/100ML; 44 8MG/100ML	N019395 001	Mar 26, 1986
INPERSOL-ZM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 538MG/100ML; 44 8MG/100ML	N019395 002	Mar 26, 1986
INPERSOL-ZM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 538MG/100ML; 4 48MG/100ML	N019395 003	Mar 26, 1986

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER ABBOTT	16.5MG/ML; 25.4MG/ML; 74.6MG/ML; 121MG/ML; 16.1MG/ML	N019399 001	Jun 16, 1986
---	---	-------------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-61(of 375)

\*\* See List Footnote

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER

B BRAUN	35MG/100ML;30MG/100ML;74MG/100ML;640MG/ 100ML;500MG/100ML;74MG/100ML 35MG/100ML;30MG/100ML;74MG/100ML;640MG/ 100ML;500MG/100ML;74MG/100ML	N018899 001 Oct 31, 1983
		N019718 001 Sep 29, 1989

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE R IN PLASTIC CONTAINER

BAXTER HLTHCARE	36.8MG/100ML;30.5MG/100ML;74.6MG/100ML; 640MG/100ML;496MG/100ML;89.6MG/100ML	N017438 001
-----------------	---	-------------

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ACETATED RINGER'S IN PLASTIC CONTAINER

B BRAUN	20MG/100ML;30MG/100ML;380MG/100ML;600MG/ 100ML	N018725 001 Nov 29, 1982
---------	---	--------------------------

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

B BRAUN	33MG/100ML;30MG/100ML;860MG/100ML	N018721 001 Nov 09, 1982
---------	-----------------------------------	--------------------------

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

ABBOTT	33MG/100ML;30MG/100ML;860MG/100ML	N018462 001
--------	-----------------------------------	-------------

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

ABBOTT	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ 100ML	N019485 001 Oct 24, 1985
--------	---	--------------------------

B BRAUN	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ 100ML	N018023 001
---------	---	-------------

MILES	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ 100ML	N018417 001
-------	---	-------------

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

BAXTER HLTHCARE	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ 100ML	N019933 001 Aug 29, 1989
-----------------	---	--------------------------

CALCIUM GLUCEPTATE

INJECTABLE; INJECTION

CALCIUM GLUCEPTATE

ABBOTT	EQ 90MG CALCIUM/5ML	A080001 001
	EQ 90MG CALCIUM/5ML	A083159 001
ABRAXIS PHARM	EQ 90MG CALCIUM/5ML	A089373 001 Apr 30, 1987
LILLY	EQ 90MG CALCIUM/5ML	N006470 001

CALCIUM METRIZOATE; MEGLUMINE METRIZOATE; METRIZOATE MAGNESIUM; METRIZOATE SODIUM

INJECTABLE; INJECTION

ISOPAQUE 440

GE HEALTHCARE	0.78MG/ML;75.9MG/ML;0.15MG/ML;16.6MG/ML	N016847 001
---------------	---	-------------

CALCIUM; MEGLUMINE; METRIZOIC ACID

INJECTABLE; INJECTION

ISOPAQUE 280

GE HEALTHCARE	0.35MG/ML;140.1MG/ML;461.8MG/ML	N017506 001
---------------	---------------------------------	-------------

CANDICIDIN

OINTMENT; VAGINAL

VANOBID

SANOFI AVENTIS US	0.6MG/GM	A061596 001
-------------------	----------	-------------

TABLET; VAGINAL

VANOBID

SANOFI AVENTIS US	3MG	A061613 001
-------------------	-----	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-62(of 375)

\*\* See List Footnote

CAPTOPRIL

TABLET;ORAL

CAPOTEN

+ PAR PHARM	12.5MG **	N018343 005	Jan 17, 1985
+	25MG **	N018343 002	
+	37.5MG **	N018343 006	Sep 17, 1986
+	50MG **	N018343 001	
+	75MG **	N018343 007	Jun 13, 1995
+	100MG **	N018343 003	
+	150MG **	N018343 004	Jun 13, 1995
<b>CAPTOPRIL</b>			
APOTEX	12.5MG	A074737 001	Oct 28, 1998
	25MG	A074737 002	Oct 28, 1998
	50MG	A074737 003	Oct 28, 1998
	100MG	A074737 004	Oct 28, 1998
APOTHECON	12.5MG	A074472 001	Mar 31, 1995
	25MG	A074472 002	Mar 31, 1995
	50MG	A074472 003	Mar 31, 1995
	100MG	A074472 004	Mar 31, 1995
BOSCOGEN	12.5MG	A074677 004	May 30, 1997
	25MG	A074677 002	May 30, 1997
	50MG	A074677 001	May 30, 1997
	100MG	A074677 003	May 30, 1997
DAVA PHARMS INC	12.5MG	A074423 001	Feb 13, 1996
	25MG	A074423 002	Feb 13, 1996
	50MG	A074423 003	Feb 13, 1996
	100MG	A074423 004	Feb 13, 1996
EGIS PHARMS	12.5MG	A074748 004	May 29, 1997
	25MG	A074748 002	May 29, 1997
	50MG	A074748 001	May 29, 1997
	100MG	A074748 003	May 29, 1997
FOSUN PHARMA	12.5MG	A074363 001	Nov 09, 1995
	25MG	A074363 002	Nov 09, 1995
	50MG	A074363 003	Nov 09, 1995
	100MG	A074363 004	Nov 09, 1995
G AND W LABS INC	12.5MG	A074433 001	Feb 13, 1996
	12.5MG	A074462 001	Feb 13, 1996
	12.5MG	A074483 001	Feb 13, 1996
	12.5MG	A074590 004	Aug 30, 1996
	25MG	A074433 002	Feb 13, 1996
	25MG	A074462 002	Feb 13, 1996
	25MG	A074483 002	Feb 13, 1996
	25MG	A074590 002	Aug 30, 1996
	50MG	A074433 003	Feb 13, 1996
	50MG	A074462 003	Feb 13, 1996
	50MG	A074483 003	Feb 13, 1996
	50MG	A074590 001	Aug 30, 1996
	100MG	A074433 004	Feb 13, 1996
	100MG	A074462 004	Feb 13, 1996
	100MG	A074483 004	Feb 13, 1996
	100MG	A074590 003	Aug 30, 1996
PAR PHARM	12.5MG	A074493 001	Feb 13, 1996
	25MG	A074493 002	Feb 13, 1996
	50MG	A074493 003	Feb 13, 1996
	100MG	A074493 004	Feb 13, 1996
PUREPAC PHARM	12.5MG	A074640 001	Mar 31, 1997
	25MG	A074640 002	Mar 31, 1997
	50MG	A074640 003	Mar 31, 1997
	100MG	A074640 004	Mar 31, 1997
SANDOZ	12.5MG	A074481 001	Feb 13, 1996
	12.5MG	A074519 001	Feb 13, 1996
	25MG	A074481 002	Feb 13, 1996
	25MG	A074519 002	Feb 13, 1996
	50MG	A074481 003	Feb 13, 1996
	50MG	A074519 003	Feb 13, 1996
	100MG	A074481 004	Feb 13, 1996
	100MG	A074519 004	Feb 13, 1996
VINTAGE PHARMS LLC	12.5MG	A074418 001	Feb 13, 1996
	25MG	A074418 002	Feb 13, 1996

**DISCONTINUED DRUG PRODUCT LIST**

6-63(of 375)

\*\* See List Footnote

CAPTOPRILTABLET;ORAL  
Captopril

	50MG	A074418 003	Feb 13, 1996
	100MG	A074418 004	Feb 13, 1996
WATSON LABS	12.5MG	A074451 001	Feb 13, 1996
	12.5MG	A074576 001	Apr 23, 1996
	25MG	A074451 002	Feb 13, 1996
	25MG	A074576 002	Apr 23, 1996
	50MG	A074451 003	Feb 13, 1996
	50MG	A074576 003	Apr 23, 1996
	100MG	A074451 004	Feb 13, 1996
	100MG	A074576 004	Apr 23, 1996

CAPTOPRIL; HYDROCHLORTIAZIDE

TABLET;ORAL

CAPOZIDE 25/15	25MG;15MG **	N018709 001	Oct 12, 1984
+ APOTHECON			
CAPOZIDE 25/25	25MG;25MG **	N018709 002	Oct 12, 1984
+ APOTHECON			
CAPOZIDE 50/15	50MG;15MG **	N018709 004	Oct 12, 1984
+ APOTHECON			
CAPOZIDE 50/25	50MG;25MG **	N018709 003	Oct 12, 1984
+ APOTHECON			
CAPTOPRIL AND HYDROCHLORTIAZIDE			
IVAX SUB TEVA PHARMS	25MG;15MG	A075055 001	Jun 18, 1998
	25MG;25MG	A075055 002	Jun 18, 1998
	50MG;15MG	A075055 004	Jun 18, 1998
	50MG;25MG	A075055 003	Jun 18, 1998
VINTAGE PHARMS LLC	25MG;15MG	A074788 001	Dec 29, 1997
	25MG;25MG	A074788 002	Dec 29, 1997
	50MG;15MG	A074788 004	Dec 29, 1997
	50MG;25MG	A074788 003	Dec 29, 1997
WATSON LABS	50MG;25MG	A074832 001	Dec 29, 1997

CARBACHOL

SOLUTION;INTRAOCULAR

CARBACHOL			
PHARMAFAIR	0.01%	A070292 001	May 21, 1986
CARBASTAT			
NOVARTIS	0.01%	A073677 001	Apr 28, 1995

CARBAMAZEPINE

SUSPENSION;ORAL

CARBAMAZEPINE			
TARO	100MG/5ML	A075875 001	Dec 21, 2000

TABLET;ORAL

CARBAMAZEPINE			
ACTAVIS ELIZABETH	200MG	A071696 001	Nov 09, 1987
INWOOD LABS	200MG	A070231 001	Aug 14, 1986
PLIVA	200MG	A071479 001	Jul 24, 1987
USL PHARMA	200MG	A070300 001	May 15, 1986
WARNER CHILCOTT	200MG	A070429 001	Jan 02, 1987

TERIL

TARO	200MG	A076525 001	Sep 26, 2003
------	-------	-------------	--------------

TABLET, CHEWABLE;ORAL

CARBAMAZEPINE			
JUBILANT CADISTA	100MG	A071940 001	Feb 01, 1988

CARBENICILLIN DISODIUM

INJECTABLE;INJECTION

GOPEN			
ROERIG	EQ 1GM BASE/VIAL	N050306 001	
	EQ 2GM BASE/VIAL	N050306 004	
	EQ 5GM BASE/VIAL	N050306 002	
	EQ 10GM BASE/VIAL	N050306 006	
	EQ 30GM BASE/VIAL	N050306 007	

PYOPEN

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	N050298 001	
	EQ 2GM BASE/VIAL	N050298 002	
	EQ 5GM BASE/VIAL	N050298 003	

**DISCONTINUED DRUG PRODUCT LIST**

6-64(of 375)

\*\* See List Footnote

CARBENICILLIN DISODIUMINJECTABLE; INJECTION  
PYOPENEQ 10GM BASE/VIAL  
EQ 20GM BASE/VIALN050298 006  
N050298 007CARBENICILLIN INDANYL SODIUMTABLET; ORAL  
GEOCILLIN

PFIZER EQ 382MG BASE N050435 001

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

IDT AUSTRALIA LTD	10MG;100MG	A073587 002	Jun 29, 1995
	25MG;100MG	A073587 001	Jun 29, 1995
	25MG;250MG	A073587 003	Jun 29, 1995
SCS	10MG;100MG	A074080 001	Mar 25, 1994
	25MG;100MG	A074080 002	Mar 25, 1994
	25MG;250MG	A074080 003	Mar 25, 1994
WATSON LABS	10MG;100MG	A073381 001	Sep 28, 1993
	25MG;100MG	A073382 001	Sep 28, 1993
	25MG;250MG	A073383 001	Sep 28, 1993

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

KV PHARM 50MG;200MG A076663 001 Jun 24, 2004

TABLET, FOR SUSPENSION; ORAL

CARBILEV

RANBAXY	10MG;100MG	A076643 001	Jun 10, 2005
	25MG;100MG	A076643 002	Jun 10, 2005
	25MG;250MG	A076643 003	Jun 10, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

CARBIDOPA AND LEVODOPA

IMPAK LABS	10MG;100MG	A090631 001	Jun 08, 2010
	25MG;100MG	A090631 002	Jun 08, 2010
	25MG;250MG	A090631 003	Jun 08, 2010

PARCOPA

UCB INC	10MG;100MG **	A076699 001	Aug 27, 2004
	25MG;100MG **	A076699 002	Aug 27, 2004
	25MG;250MG **	A076699 003	Aug 27, 2004

CARBINOXAMINE MALEATE

ELIXIR; ORAL

CLISTIN

+ MCNEIL 4MG/5ML \*\*

N008955 001

TABLET; ORAL

CLISTIN

+ ORTHO MCNEIL PHARM 4MG \*\*

N008915 001

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

CIPLA LTD	50MG/VIAL	A077383 001	Jan 27, 2006
	150MG/VIAL	A077383 002	Jan 27, 2006
	450MG/VIAL	A077383 003	Jan 27, 2006
FRESENIUS KABI USA	50MG/VIAL	A076235 001	Oct 14, 2004
	150MG/VIAL	A076235 002	Oct 14, 2004
	450MG/VIAL	A076235 003	Oct 14, 2004
HOSPIRA	50MG/VIAL	A076473 001	Oct 27, 2004
	150MG/VIAL	A076473 002	Oct 27, 2004
	450MG/VIAL	A076473 003	Oct 27, 2004
MYLAN LABS LTD	50MG/VIAL	A091510 001	May 29, 2012
	150MG/VIAL	A091510 002	May 29, 2012
	450MG/VIAL	A091510 003	May 29, 2012
PLIVA	50MG/VIAL	A076602 001	Nov 16, 2004
	150MG/VIAL	A076602 002	Nov 16, 2004
	450MG/VIAL	A076602 003	Nov 16, 2004
SANDOZ	50MG/VIAL	A076959 001	Mar 18, 2005
	150MG/VIAL	A076959 002	Mar 18, 2005
	450MG/VIAL	A076959 003	Mar 18, 2005
WATSON LABS TEVA	50MG/VIAL	A076162 001	Oct 14, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-65(of 375)

\*\* See List Footnote

**CARBOPLATIN**

INJECTABLE; INJECTION  
CARBOPLATIN

WEST-WARD PHARMS INT	150MG/VIAL 450MG/VIAL 50MG/VIAL 150MG/VIAL 450MG/VIAL	A076162 002 Oct 14, 2004 A076162 003 Oct 14, 2004 A076099 001 Oct 14, 2004 A076099 002 Oct 14, 2004 A076099 003 Oct 14, 2004
PARAPLATIN		
+ CORDEN PHARMA	50MG/VIAL **	N019880 001 Mar 03, 1989
+	150MG/VIAL **	N019880 002 Mar 03, 1989
+	450MG/VIAL **	N019880 003 Mar 03, 1989
INJECTABLE; IV (INFUSION)		
CARBOPLATIN		
ACTAVIS TOTOWA	50MG/5ML (10MG/ML) 150MG/15ML (10MG/ML) 450MG/45ML (10MG/ML) 600MG/60ML (10MG/ML)	A078732 001 Feb 06, 2012 A078732 002 Feb 06, 2012 A078732 003 Feb 06, 2012 A078732 004 Feb 06, 2012
FRESENIUS KABI USA	50MG/5ML (10MG/ML) 50MG/5ML (10MG/ML) 150MG/15ML (10MG/ML) 150MG/15ML (10MG/ML)	A077247 001 Oct 21, 2004 A077266 001 Feb 15, 2006 A077247 002 Oct 21, 2004 A077266 002 Feb 15, 2006
PHARMACHEMIE BV	50MG/5ML (10MG/ML) 150MG/15ML (10MG/ML) 450MG/45ML (10MG/ML)	A077679 001 Feb 25, 2009 A077679 002 Feb 25, 2009 A077679 003 Feb 25, 2009
TEVA PARENTERAL	50MG/5ML (10MG/ML) 150MG/15ML (10MG/ML) 450MG/45ML (10MG/ML)	A077389 001 Mar 30, 2007 A077389 002 Mar 30, 2007 A077389 003 Mar 30, 2007
PARAPLATIN		
+ CORDENPHARMA	50MG/5ML (10MG/ML) **	N020452 001 Jul 14, 2003
+	150MG/15ML (10MG/ML) **	N020452 002 Jul 14, 2003
+	450MG/45ML (10MG/ML) **	N020452 003 Jul 14, 2003
+	600MG/60ML (10MG/ML) **	N020452 004 Jan 15, 2004

**CARISOPRODOL**

CAPSULE; ORAL  
SOMA

MYLAN SPECIALITY LP	250MG	N011792 003
---------------------	-------	-------------

TABLET; ORAL

CARISOPRODOL		
ABLE	350MG	A040421 001 Jun 21, 2001
EPIC PHARMA LLC	350MG	A040397 001 Sep 21, 2000
OXFORD PHARMS	350MG	A040188 001 Mar 07, 1997
PIONEER PHARMS	350MG	A089390 001 Oct 13, 1988
SANDOZ	350MG	A081025 001 Apr 13, 1989
	350MG	A089566 001 Aug 30, 1988
SUN PHARM INDS LTD	350MG	A040755 001 Feb 27, 2007
WATSON LABS	350MG	A040152 001 Dec 03, 1996
	350MG	A085433 001
WATSON LABS TEVA	350MG	A086179 001

RELA

SCHERING	350MG	N012155 001
----------	-------	-------------

**CARPHENAZINE MALEATE**

CONCENTRATE; ORAL  
PROKETAZINE

WYETH AYERST	50MG/ML	N014173 001
--------------	---------	-------------

TABLET; ORAL

PROKETAZINE		
WYETH AYERST	12.5MG	N012768 001
	25MG	N012768 002
	50MG	N012768 004

**CARPROFEN**

TABLET; ORAL  
RIMADYL

ROCHE	100MG 150MG	N018550 002 Dec 31, 1987 N018550 003 Dec 31, 1987
-------	----------------	--

**DISCONTINUED DRUG PRODUCT LIST**

6-66(of 375)

\*\* See List Footnote

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

APOTEX INC

1%

A076097 001 Feb 06, 2002

OCUPRESS

+ NOVARTIS

1%

N019972 001 May 23, 1990

TABLET;ORAL

CARTROL

ABBVIE

2.5MG

N019204 001 Dec 28, 1988

5MG

N019204 002 Dec 28, 1988

10MG

N019204 003 Dec 28, 1988

CARVEDILOL

TABLET;ORAL

CARVEDILOL

HIKMA

3.125MG

A077887 001 Sep 07, 2007

6.25MG

A077887 002 Sep 07, 2007

12.5MG

A077887 003 Sep 07, 2007

25MG

A077887 004 Sep 07, 2007

PLIVA HRVATSKA DOO

3.125MG

A078240 001 Oct 30, 2007

6.25MG

A078240 002 Oct 30, 2007

12.5MG

A078240 003 Oct 30, 2007

25MG

A078240 004 Oct 30, 2007

WOCKHARDT LTD

3.125MG

A078786 001 Dec 22, 2009

6.25MG

A078786 002 Dec 22, 2009

12.5MG

A078786 003 Dec 22, 2009

25MG

A078786 004 Dec 22, 2009

CEFACLOR

CAPSULE;ORAL

CECLOR

+ LILLY

EQ 250MG BASE \*\*

N050521 001

+

EQ 500MG BASE \*\*

N050521 002

CEFACLOR

CEPH INTL

EQ 250MG BASE

A062205 001

EQ 500MG BASE

A062205 002

DAVA PHARMS INC

EQ 250MG BASE

A064107 001 Apr 27, 1995

EQ 500MG BASE

A064107 002 Apr 27, 1995

IVAX SUB TEVA PHARMS

EQ 250MG BASE

A064061 001 Apr 27, 1995

EQ 500MG BASE

A064061 002 Apr 27, 1995

RANBAXY

EQ 250MG BASE

A064156 001 Aug 28, 1997

EQ 500MG BASE

A064156 002 Aug 28, 1997

TEVA

EQ 250MG BASE

A064081 001 Sep 16, 1996

EQ 250MG BASE

A064145 001 Jun 24, 1996

EQ 500MG BASE

A064081 002 Sep 16, 1996

EQ 500MG BASE

A064145 002 Jun 24, 1996

WATSON LABS INC

EQ 250MG BASE

A064148 001 May 23, 1996

EQ 500MG BASE

A064148 002 May 23, 1996

FOR SUSPENSION;ORAL

CECLOR

+ LILLY

EQ 125MG BASE/5ML \*\*

N050522 001

+

EQ 250MG BASE/5ML \*\*

N050522 002

CEFACLOR

DAVA PHARMS INC

EQ 125MG BASE/5ML

A064114 001 Apr 28, 1995

EQ 187MG BASE/5ML

A064115 001 Apr 28, 1995

EQ 250MG BASE/5ML

A064116 001 Apr 28, 1995

EQ 375MG BASE/5ML

A064110 001 Apr 28, 1995

FACTA FARMA

EQ 125MG BASE/5ML

A062206 001

EQ 187MG BASE/5ML

A062206 003 Apr 20, 1988

EQ 250MG BASE/5ML

A062206 002

EQ 375MG BASE/5ML

A062206 004 Apr 20, 1988

IVAX SUB TEVA PHARMS

EQ 125MG BASE/5ML

A064087 001 Apr 28, 1995

EQ 187MG BASE/5ML

A064086 001 Apr 28, 1995

EQ 250MG BASE/5ML

A064085 001 Apr 28, 1995

EQ 375MG BASE/5ML

A064070 001 Apr 28, 1995

RANBAXY

EQ 125MG BASE/5ML

A064166 001 Oct 02, 1997

EQ 187MG BASE/5ML

A064165 001 Oct 02, 1997

EQ 250MG BASE/5ML

A064164 001 Oct 02, 1997

EQ 375MG BASE/5ML

A064155 001 Oct 02, 1997

WATSON LABS INC

EQ 125MG BASE/5ML

A064204 001 Feb 18, 1998

**DISCONTINUED DRUG PRODUCT LIST**

6-67(of 375)

\*\* See List Footnote

**CEFACLOR**FOR SUSPENSION;ORAL  
CEFACLOR

EQ 187MG BASE/5ML	A064205	001	Feb 18, 1998
EQ 250MG BASE/5ML	A064206	001	Feb 18, 1998
EQ 375MG BASE/5ML	A064207	001	Feb 18, 1998

TABLET, CHEWABLE;ORAL

RANICLOR

RANBAXY LABS LTD	EQ 125MG BASE	A065092	001	Dec 22, 2003
	EQ 187MG BASE	A065092	002	Dec 22, 2003
	EQ 250MG BASE	A065092	003	Dec 22, 2003
	EQ 375MG BASE	A065092	004	Dec 22, 2003

TABLET, EXTENDED RELEASE;ORAL

CECLOR CD

LILLY	EQ 375MG BASE	N050673	001	Jun 28, 1996
	EQ 500MG BASE	N050673	002	Jun 28, 1996

CEFACLOR

WORLD GEN	EQ 500MG BASE	A065057	001	Jan 05, 2001
-----------	---------------	---------	-----	--------------

**CEFADROXIL/CEFADROXIL HEMIHYDRATE**

CAPSULE;ORAL

CEFADROXIL

IVAX SUB TEVA PHARMS	EQ 500MG BASE	A062766	001	Mar 03, 1987
PUREPAC PHARM	EQ 500MG BASE	A063017	001	Jan 05, 1989
RANBAXY LABS LTD	EQ 500MG BASE	A065015	001	Jun 22, 1999
SANDOZ	EQ 500MG BASE	A062291	001	
TEVA	EQ 500MG BASE	A062695	001	Feb 10, 1989

DURICEF

WARNER CHILCOTT	EQ 250MG BASE	N050512	002	
+	EQ 500MG BASE **	N050512	001	

ULTRACEF

BRISTOL	EQ 500MG BASE	A062378	001	Mar 16, 1982
---------	---------------	---------	-----	--------------

FOR SUSPENSION;ORAL

CEFADROXIL

ANI PHARMS INC	EQ 125MG BASE/5ML	A062698	001	Mar 01, 1989
	EQ 250MG BASE/5ML	A062698	002	Mar 01, 1989
	EQ 250MG BASE/5ML	A065278	001	Jan 20, 2006
	EQ 500MG BASE/5ML	A062698	003	Mar 01, 1989
	EQ 500MG BASE/5ML	A065278	002	Jan 20, 2006

APOTHECON	EQ 125MG BASE/5ML	A062334	001	
	EQ 250MG BASE/5ML	A062334	002	
	EQ 500MG BASE/5ML	A062334	003	

SUN PHARM INDs LTD	EQ 125MG BASE/5ML	A065115	001	Mar 26, 2003
	EQ 250MG BASE/5ML	A065115	002	Mar 26, 2003
	EQ 500MG BASE/5ML	A065115	003	Mar 26, 2003

DURICEF

+	WARNER CHILCOTT	EQ 125MG BASE/5ML **	N050527	002	
+		EQ 250MG BASE/5ML **	N050527	003	
+		EQ 500MG BASE/5ML **	N050527	001	

ULTRACEF

BRISTOL	EQ 125MG BASE/5ML	A062376	001	Mar 16, 1982
	EQ 250MG BASE/5ML	A062376	002	Mar 16, 1982
	EQ 500MG BASE/5ML	A062376	003	Mar 16, 1982

TABLET;ORAL

CEFADROXIL

RANBAXY	EQ 1GM BASE	A065018	001	Apr 23, 1999
---------	-------------	---------	-----	--------------

DURICEF

+	WARNER CHILCOTT	EQ 1GM BASE **	N050528	001	
---	-----------------	----------------	---------	-----	--

ULTRACEF

APOTHECON	EQ 1GM BASE	A062390	001	Jun 10, 1982
BRISTOL	EQ 1GM BASE	A062408	001	Aug 31, 1982

**CEFAMANDOLE NAFATE**

INJECTABLE;INJECTION

MANDOL

LILLY	EQ 500MG BASE/VIAL	N050504	001	
	EQ 1GM BASE/VIAL	A062560	001	Sep 10, 1985
	EQ 1GM BASE/VIAL	N050504	002	
	EQ 2GM BASE/VIAL	A062560	002	Sep 10, 1985
	EQ 2GM BASE/VIAL	N050504	003	

**DISCONTINUED DRUG PRODUCT LIST**

6-68(of 375)

\*\* See List Footnote

CEFAMANDOLE NAFATEINJECTABLE; INJECTION  
MANDOL

EQ 10GM BASE/VIAL

N050504 004

CEFAZOLIN SODIUMINJECTABLE; INJECTION  
ANCEF

+ GLAXOSMITHKLINE	EQ 250MG BASE/VIAL **
+ GLAXOSMITHKLINE	EQ 500MG BASE/VIAL
+ GLAXOSMITHKLINE	EQ 1GM BASE/VIAL **
+ GLAXOSMITHKLINE	EQ 5GM BASE/VIAL **
+ GLAXOSMITHKLINE	EQ 10GM BASE/VIAL **

N050461 001
N050461 002
N050461 003
N050461 004
N050461 005

ANCEF IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 10MG BASE/ML
BAXTER HLTHCARE	EQ 20MG BASE/ML

N050566 003 Jun 08, 1983
N050566 004 Jun 08, 1983

ANCEF IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 10MG BASE/ML
BAXTER HLTHCARE	EQ 20MG BASE/ML

N050566 001 Jun 08, 1983
N050566 002 Jun 08, 1983

CEFAZOLIN AND DEXTROSE

B BRAUN	EQ 500MG BASE/VIAL
---------	--------------------

N050779 001 Jul 27, 2000
--------------------------

CEFAZOLIN SODIUM

ABRAXIS PHARM	EQ 500MG BASE/VIAL
ABRAXIS PHARM	EQ 1GM BASE/VIAL
ABRAXIS PHARM	EQ 10GM BASE/VIAL
ABRAXIS PHARM	EQ 20GM BASE/VIAL

A062688 002 Nov 17, 1986
A062688 003 Nov 17, 1986
A062688 004 Nov 17, 1986
A062688 005 Aug 03, 1987

AUROBINDO PHARMA	EQ 500MG BASE/VIAL
AUROBINDO PHARMA	EQ 1GM BASE/VIAL
AUROBINDO PHARMA	EQ 250MG BASE/VIAL

A065395 001 Aug 08, 2008
A065395 002 Aug 08, 2008

BEDFORD	EQ 500MG BASE/VIAL
BEDFORD	EQ 1GM BASE/VIAL
BEDFORD	EQ 5GM BASE/VIAL

A062894 001 Jul 21, 1988
A062894 002 Jul 21, 1988

CEPHAZONE PHARMA	EQ 10GM BASE/VIAL
CEPHAZONE PHARMA	EQ 500MG BASE/VIAL
CEPHAZONE PHARMA	EQ 1GM BASE/VIAL

A065280 001 Mar 18, 2009
A065280 002 Mar 18, 2009

FACTA FARMA	EQ 10GM BASE/VIAL
FACTA FARMA	EQ 20GM BASE/VIAL
FACTA FARMA	EQ 500MG BASE/VIAL

A063207 001 Dec 27, 1991
A063214 001 Dec 27, 1991

FRESENIUS KABI USA	EQ 10GM BASE/VIAL
FRESENIUS KABI USA	EQ 20GM BASE/VIAL
FRESENIUS KABI USA	EQ 500MG BASE/VIAL **

A064169 001 Aug 14, 1998
A064169 002 Aug 14, 1998

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL
STERI PHARMA	EQ 500MG BASE/VIAL
STERI PHARMA	EQ 1GM BASE/VIAL

A064170 001 Mar 18, 1998
A064170 002 Mar 18, 1998

TEVA PHARMS	EQ 250MG BASE/VIAL
TEVA PHARMS	EQ 500MG BASE/VIAL
TEVA PHARMS	EQ 1GM BASE/VIAL

A063016 001 Mar 14, 1989
A063016 002 Mar 14, 1989

WATSON LABS INC	EQ 5GM BASE/VIAL
WATSON LABS INC	EQ 10GM BASE/VIAL
WATSON LABS INC	EQ 20GM BASE/VIAL

A063016 003 Mar 14, 1989
A063018 001 Mar 05, 1990

WEST-WARD PHARMS INT	EQ 250MG BASE/VIAL
WEST-WARD PHARMS INT	EQ 500MG BASE/VIAL
WEST-WARD PHARMS INT	EQ 1GM BASE/VIAL

A062807 001 Jan 12, 1988
A062807 002 Jan 12, 1988

WEST-WARD PHARMS INT	EQ 5GM BASE/VIAL
WEST-WARD PHARMS INT	EQ 10GM BASE/VIAL
WEST-WARD PHARMS INT	EQ 20GM BASE/VIAL

A062807 003 Jan 12, 1988
A062807 004 Jan 12, 1988

WEST-WARD PHARMS INT	EQ 250MG BASE/VIAL
WEST-WARD PHARMS INT	EQ 500MG BASE/VIAL
WEST-WARD PHARMS INT	EQ 1GM BASE/VIAL

A062807 005 Jan 12, 1988
A062807 006 Jan 12, 1988

KEFZOL

ACS DOBFAR	EQ 250MG BASE/VIAL
ACS DOBFAR	EQ 20GM BASE/VIAL

A061773 001
A061773 005 Sep 08, 1987

LILLY	EQ 500MG BASE/VIAL
LILLY	EQ 1GM BASE/VIAL

A062557 001 Sep 10, 1985
A062557 002 Sep 10, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-69(of 375)

\*\* See List Footnote

CEFDINIR

CAPSULE;ORAL OMNICEF + ABBVIE	300MG **	N050739 001 Dec 04, 1997
FOR SUSPENSION;ORAL OMNICEF + ABBVIE	125MG/5ML **	N050749 001 Dec 04, 1997
+ ABBVIE	250MG/5ML **	N050749 002 Jul 29, 2004

CEFDITOREN PIVOXIL

TABLET;ORAL SPECTRACEF VANSEN PHARMA	200MG 400MG	N021222 001 Aug 29, 2001 N021222 002 Jul 21, 2008
--	----------------	--

CEFEPIME HYDROCHLORIDE

INJECTABLE;INJECTION CEFEPIME HYDROCHLORIDE SANDOZ	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 500MG BASE/VIAL	A090291 002 Dec 21, 2010 A090291 003 Dec 21, 2010 A090291 001 Dec 21, 2010
--	--	--

CEFIXIME

FOR SUSPENSION;ORAL SUPRAX + LEDERLE	100MG/5ML **	N050622 001 Apr 28, 1989
TABLET;ORAL SUPRAX + LEDERLE	200MG ** 400MG **	N050621 001 Apr 28, 1989 N050621 002 Apr 28, 1989

CEFMENOXIME HYDROCHLORIDE

INJECTABLE;INJECTION CEFMAX TAP PHARM	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	N050571 001 Dec 30, 1987 N050571 002 Dec 30, 1987 N050571 003 Dec 30, 1987
---	--	--

CEFMETAZOLE SODIUM

INJECTABLE;INJECTION ZEFAZONE + PHARMACIA AND UPJOHN	EQ 1GM BASE/VIAL ** EQ 2GM BASE/VIAL **	N050637 001 Dec 11, 1989 N050637 002 Dec 11, 1989
ZEFAZONE IN PLASTIC CONTAINER + PHARMACIA AND UPJOHN	EQ 20MG BASE/ML ** EQ 40MG BASE/ML **	N050683 001 Dec 29, 1992 N050683 002 Dec 29, 1992

CEFONICID SODIUM

INJECTABLE;INJECTION MONOCID GLAXOSMITHKLINE	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 10GM BASE/VIAL	N050579 001 May 23, 1984 A063295 001 Jul 26, 1993 N050579 002 May 23, 1984 N050579 003 May 23, 1984 N050579 004 May 23, 1984
--	---	--

CEFOPERAZONE SODIUM

INJECTABLE;INJECTION CEFOBID PFIZER	EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL EQ 10GM BASE/VIAL	A063333 001 Mar 31, 1995 N050551 001 Nov 18, 1982 A063333 002 Mar 31, 1995 N050551 002 Nov 18, 1982 N050551 003 Mar 05, 1990
CEFOBID IN PLASTIC CONTAINER PFIZER	EQ 20MG BASE/ML EQ 40MG BASE/ML	N050613 002 Jul 31, 1987 N050613 001 Jul 23, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-70(of 375)

\*\* See List Footnote

**CEFORANIDE**INJECTABLE; INJECTION  
PRECEF

APOTHECON	500MG/VIAL 1GM/VIAL 2GM/VIAL 10GM/VIAL 20GM/VIAL	A062579 001 Nov 26, 1984 A062579 002 Nov 26, 1984 A062579 003 Nov 26, 1984 A062579 004 Nov 26, 1984 A062579 005 Nov 26, 1984
BRISTOL	500MG/VIAL 1GM/VIAL 2GM/VIAL 10GM/VIAL 20GM/VIAL	N050554 001 May 24, 1984 N050554 002 May 24, 1984 N050554 003 May 24, 1984 N050554 004 May 24, 1984 N050554 005 May 24, 1984

**CEFOTAXIME SODIUM**INJECTABLE; INJECTION  
CEFOTAXIME

FRESENIUS KABI USA	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 10GM BASE/VIAL EQ 20GM BASE/VIAL	A064200 001 Mar 24, 2000 A064200 002 Mar 24, 2000 A064200 003 Mar 24, 2000 A064201 001 Mar 24, 2000 A064201 002 Mar 24, 2000
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER B BRAUN	EQ 2GM BASE	N050792 001 Jul 29, 2004
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER B BRAUN	EQ 1GM BASE	N050792 002 Jul 29, 2004
CEFOTAXIME SODIUM		
AUROBINDO PHARMA	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	A065517 001 Nov 06, 2009 A065517 002 Nov 06, 2009 A065517 003 Nov 06, 2009
AUROBINDO PHARMA LTD	EQ 10GM BASE/VIAL	A065516 001 Nov 06, 2009
CEPHAZONE PHARMA	EQ 10GM BASE/VIAL	A065348 001 Jan 25, 2010
CLAFORAN		
SANOFI AVENTIS US	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	A062659 001 Jan 13, 1987 A062659 002 Jan 13, 1987
CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER US PHARM HOLDINGS	EQ 20MG BASE/ML EQ 40MG BASE/ML	N050596 002 May 20, 1985 N050596 004 May 20, 1985
CLAFORAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER US PHARM HOLDINGS	EQ 20MG BASE/ML EQ 40MG BASE/ML	N050596 001 May 20, 1985 N050596 003 May 20, 1985

**CEFOTETAN DISODIUM**INJECTABLE; INJECTION  
CEFOTAN

TELIGENT	EQ 10GM BASE/VIAL	N050588 003 Apr 25, 1988
TELIGENT PHARMA INC	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	A063293 001 Apr 29, 1993 A063293 002 Apr 29, 1993
CEFOTAN IN PLASTIC CONTAINER TELIGENT	EQ 20MG BASE/ML EQ 40MG BASE/ML	N050694 002 Jul 30, 1993 N050694 001 Jul 30, 1993

**CEFOTIAM HYDROCHLORIDE**INJECTABLE; INJECTION  
CERADON

TAKEDA	EQ 1GM BASE/VIAL	N050601 001 Dec 30, 1988
--------	------------------	--------------------------

**CEFOXITIN SODIUM**INJECTABLE; INJECTION  
CEFOXITIN

ACS DOBFAR SPA	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 10GM BASE/VIAL	A065467 001 Aug 31, 2011 A065467 002 Aug 31, 2011 A065464 001 Aug 31, 2011
FRESENIUS KABI USA	EQ 1GM BASE/VIAL ** EQ 2GM BASE/VIAL ** EQ 10GM BASE/VIAL	A065012 001 Jul 03, 2000 A065012 002 Jul 03, 2000 A065011 001 Jul 03, 2000
MEFOXIN		
MYLAN INSTITUTIONAL	EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL ** EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL **	A062757 001 Jan 08, 1987 N050517 001 A062757 002 Jan 08, 1987 N050517 002

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*



**DISCONTINUED DRUG PRODUCT LIST**

6-72(of 375)

\*\* See List Footnote

CEFTAZIDIMEINJECTABLE; INJECTION  
PENTACEF

	2GM/VIAL	A063322	002	Nov 07,	1995
	2GM/VIAL	A064006	002	Mar 31,	1992
	6GM/VIAL	A064008	001	Mar 31,	1992
	10GM/VIAL	A064008	002	Mar 31,	1992
TAZIDIME					
LILLY	1GM/VIAL	A062655	001	Nov 20,	1985
	2GM/VIAL	A062655	002	Nov 20,	1985
TAZIDIME IN PLASTIC CONTAINER					
LILLY	1GM/VIAL	A062739	001	Jul 10,	1986
	2GM/VIAL	A062739	002	Jul 10,	1986

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

CEFTAZIDIME SODIUM IN PLASTIC CONTAINER					
BAXTER HLTHCARE	EQ 10MG BASE/ML	A063221	001	Apr 29,	1993
	EQ 20MG BASE/ML	A063221	002	Apr 29,	1993
	EQ 40MG BASE/ML	A063221	003	Apr 29,	1993
FORTAZ IN PLASTIC CONTAINER					
TELIGENT	EQ 10MG BASE/ML	N050634	001	Apr 28,	1989

CEFTIBUTEN DIHYDRATECAPSULE; ORAL  
CEDAX

PERNIX THERAP	EQ 400MG BASE	N050685	002	Dec 20,	1995
FOR SUSPENSION; ORAL					
CEDAX					
+ PERNIX THERAP	EQ 90MG BASE/5ML **	N050686	001	Dec 20,	1995
+ +	EQ 180MG BASE/5ML **	N050686	002	Dec 20,	1995

CEFTIZOXIME SODIUMINJECTABLE; INJECTION  
CEFIZOX

ASTELLAS	EQ 500MG BASE/VIAL	N050560	001	Sep 15,	1983
	EQ 1GM BASE/VIAL	A063294	002	Mar 31,	1994
	EQ 1GM BASE/VIAL	N050560	002	Sep 15,	1983
	EQ 2GM BASE/VIAL	A063294	003	Mar 31,	1994
	EQ 2GM BASE/VIAL	N050560	003	Sep 15,	1983
	EQ 10GM BASE/VIAL	N050560	005	Mar 19,	1993
CEFIZOX IN DEXTROSE 5% IN PLASTIC CONTAINER					
ASTELLAS	EQ 20MG BASE/ML	N050589	001	Oct 03,	1984
	EQ 40MG BASE/ML	N050589	002	Oct 03,	1984
CEFIZOX IN PLASTIC CONTAINER					
ASTELLAS	EQ 20MG BASE/ML	N050589	003	Apr 13,	1995
	EQ 40MG BASE/ML	N050589	004	Apr 13,	1995

CEFTRIAXONE SODIUMINJECTABLE; INJECTION  
CEFTRIAXONE

AGILA SPECLTS	EQ 10GM BASE/VIAL	A091068	001	Jan 07,	2013
AUROBINDO PHARMA LTD	EQ 10GM BASE/VIAL	A065504	001	Jul 31,	2008
FACTA FARMA	EQ 10GM BASE/VIAL	A065269	001	Feb 28,	2007
FRESENIUS KABI USA	EQ 10GM BASE/VIAL	A065252	001	Feb 15,	2006
HOSPIRA INC	EQ 1GM BASE/VIAL	A065231	001	Aug 02,	2005
	EQ 1GM BASE/VIAL	A202563	001	Aug 20,	2012
	EQ 2GM BASE/VIAL	A065231	002	Aug 02,	2005
	EQ 2GM BASE/VIAL	A202563	002	Aug 20,	2012
TEVA	EQ 10GM BASE/VIAL	A065274	001	May 01,	2006
ROCEPHIN					
HOFFMANN LA ROCHE	EQ 250MG BASE/VIAL	A063239	001	Aug 13,	1993
	EQ 500MG BASE/VIAL	A062654	001	Apr 30,	1987
	EQ 500MG BASE/VIAL	A063239	002	Aug 13,	1993
	EQ 1GM BASE/VIAL	A062654	002	Apr 30,	1987
	EQ 1GM BASE/VIAL	A063239	003	Aug 13,	1993
	EQ 2GM BASE/VIAL	A062654	003	Apr 30,	1987
+	EQ 10GM BASE/VIAL	N050585	005	Dec 21,	1984
ROCHE	EQ 250MG BASE/VIAL	A062510	001	Mar 12,	1985
	EQ 500MG BASE/VIAL	A062510	002	Mar 12,	1985
	EQ 1GM BASE/VIAL	A062510	003	Mar 12,	1985

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-73(of 375)

\*\* See List Footnote

CEFTRIAZONE SODIUM

## INJECTABLE; INJECTION

ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER  
 + HOFFMANN LA ROCHE EQ 10MG BASE/ML \*\*  
 + EQ 20MG BASE/ML \*\*  
 + EQ 40MG BASE/ML \*\*

N050624 001 Feb 11, 1987  
 N050624 002 Feb 11, 1987  
 N050624 003 Feb 11, 1987

## INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

## CEFTRIAZONE

AUROBINDO PHARMA LTD EQ 250MG BASE/VIAL  
 EQ 500MG BASE/VIAL  
 EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL  
 BEDFORD EQ 250MG BASE/VIAL  
 EQ 500MG BASE/VIAL  
 EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL  
 CEPHAZONE PHARMA EQ 250MG BASE/VIAL  
 EQ 500MG BASE/VIAL  
 EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL  
 FACTA FARMA EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL  
 FRESENIUS KABI USA EQ 250MG BASE/VIAL  
 EQ 500MG BASE/VIAL  
 EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL  
 TEVA EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL  
 TEVA PHARMS USA EQ 250MG BASE/VIAL  
 EQ 500MG BASE/VIAL  
 EQ 1GM BASE/VIAL  
 EQ 2GM BASE/VIAL

A065505 001 Jul 31, 2008  
 A065505 002 Jul 31, 2008  
 A065505 003 Jul 31, 2008  
 A065505 004 Jul 31, 2008  
 A065465 001 Aug 18, 2008  
 A065465 002 Aug 18, 2008  
 A065465 003 Aug 18, 2008  
 A065465 004 Aug 18, 2008  
 A065294 001 Mar 26, 2007  
 A065294 002 Mar 26, 2007  
 A065294 003 Mar 26, 2007  
 A065294 004 Mar 26, 2007  
 A065268 001 Feb 28, 2007  
 A065268 002 Feb 28, 2007  
 A065245 001 Feb 15, 2006  
 A065245 002 Feb 15, 2006  
 A065245 003 Feb 15, 2006  
 A065245 004 Feb 15, 2006  
 A065262 001 Jun 29, 2006  
 A065262 002 Jun 29, 2006  
 A065227 001 Mar 15, 2007  
 A065227 002 Mar 15, 2007  
 A065227 003 Mar 15, 2007  
 A065227 004 Mar 15, 2007

## ROCEPHIN

+ HOFFMANN LA ROCHE EQ 250MG BASE/VIAL  
 + EQ 500MG BASE/VIAL  
 + EQ 1GM BASE/VIAL  
 + EQ 2GM BASE/VIAL

N050585 001 Dec 21, 1984  
 N050585 002 Dec 21, 1984  
 N050585 003 Dec 21, 1984  
 N050585 004 Dec 21, 1984

CEFTRIAZONE SODIUM; LIDOCAINE

## INJECTABLE; INJECTION

## ROCEPHIN KIT

HOFFMANN LA ROCHE EQ 500MG BASE/VIAL,N/A;N/A,1%  
 EQ 1GM BASE/VIAL,N/A;N/A,1%

N050585 007 May 08, 1996  
 N050585 006 May 08, 1996

CEFUROXIME AXETIL

## FOR SUSPENSION; ORAL

## CEFUROXIME AXETIL

SUN PHARM INDs LTD EQ 125MG BASE/5ML  
 EQ 250MG BASE/5ML

A065323 001 Feb 05, 2008  
 A065323 002 Feb 05, 2008

## TABLET; ORAL

## CEFUROXIME AXETIL

RANBAXY LABS LTD EQ 125MG BASE  
 EQ 250MG BASE  
 EQ 500MG BASE  
 SANDOZ EQ 250MG BASE  
 EQ 500MG BASE  
 SUN PHARM INDs LTD EQ 125MG BASE  
 EQ 250MG BASE  
 EQ 500MG BASE

A065043 003 Feb 15, 2002  
 A065043 002 Feb 15, 2002  
 A065043 001 Feb 15, 2002  
 A065126 001 Oct 28, 2003  
 A065126 002 Oct 28, 2003  
 A065118 001 Apr 25, 2003  
 A065118 002 Apr 25, 2003  
 A065118 003 Apr 25, 2003

CEFUROXIME SODIUM

## INJECTABLE; INJECTION

## CEFUROXIME SODIUM

FRESENIUS KABI USA EQ 1.5GM BASE/VIAL  
 EQ 7.5GM BASE/VIAL  
 TEVA PHARMS EQ 7.5GM BASE/VIAL  
 WATSON LABS INC EQ 1.5GM BASE/VIAL  
 EQ 7.5GM BASE/VIAL

A065001 002 May 30, 2001  
 A065002 001 Sep 28, 1998  
 A064191 001 Apr 16, 1998  
 A064035 002 Feb 26, 1993  
 A064036 001 Feb 26, 1993

## CEFUROXIME SODIUM IN PLASTIC CONTAINER

SAMSON MEDCL EQ 75GM BASE/VIAL  
 EQ 225GM BASE/VIAL

A065251 001 Dec 30, 2009  
 A065251 002 Dec 30, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-74(of 375)

\*\* See List Footnote

CEFUROXIME SODIUM

INJECTABLE; INJECTION

KEFUROX

ACS DOBFAR	EQ 1.5GM BASE/VIAL	A062591 002	Jan 10, 1986
	EQ 7.5GM BASE/VIAL	A062591 003	Dec 17, 1987
LILLY	EQ 1.5GM BASE/VIAL	A062592 002	Jan 10, 1986

KEFUROX IN PLASTIC CONTAINER

LILLY	EQ 1.5GM BASE/VIAL	A062590 002	Jan 10, 1986
-------	--------------------	-------------	--------------

ZINACEF IN PLASTIC CONTAINER

TELIGENT	EQ 15MG BASE/ML	N050643 001	Apr 28, 1989
----------	-----------------	-------------	--------------

INJECTABLE; INTRAMUSCULAR, INTRAVENOUS

CEFUROXIME SODIUM

FRESENIUS KABI USA	EQ 750MG BASE/VIAL	A065001 001	May 30, 2001
TEVA PHARMS	EQ 750MG BASE/VIAL	A064192 002	Apr 16, 1998
	EQ 1.5GM BASE/VIAL	A064192 001	Apr 16, 1998
WATSON LABS INC	EQ 750MG BASE/VIAL	A064035 001	Feb 26, 1993

KEFUROX

ACS DOBFAR	EQ 750MG BASE/VIAL	A062591 001	Jan 10, 1986
------------	--------------------	-------------	--------------

INJECTABLE; INTRAVENOUS

KEFUROX

LILLY	EQ 750MG BASE/VIAL	A062592 001	Jan 10, 1986
-------	--------------------	-------------	--------------

KEFUROX IN PLASTIC CONTAINER

LILLY	EQ 750MG BASE/VIAL	A062590 001	Jan 10, 1986
-------	--------------------	-------------	--------------

CELLULOSE SODIUM PHOSPHATE

POWDER; ORAL

CALCIBIND

MISSION PHARMA	2.5GM/PACKET	N018757 002	Dec 28, 1982
	300GM/BOT	N018757 003	Oct 16, 1984

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

APOTHECON	EQ 250MG BASE	A062973 001	Nov 08, 1988
	EQ 250MG BASE	A063063 001	Sep 29, 1989
	EQ 250MG BASE	A063186 001	Dec 30, 1994
	EQ 500MG BASE	A062974 001	Nov 23, 1988
	EQ 500MG BASE	A063063 002	Sep 29, 1989
	EQ 500MG BASE	A063186 002	Dec 30, 1994

BARR	EQ 250MG BASE	A062773 001	Jun 26, 1987
	EQ 500MG BASE	A062775 001	Apr 22, 1987

FACTA FARMA	EQ 250MG BASE	A062118 001	
	EQ 500MG BASE	A062118 002	

IVAX SUB TEVA PHARMS	EQ 250MG BASE	A061969 001	
	EQ 500MG BASE	A061969 002	

PUREPAC PHARM	EQ 250MG BASE	A062809 001	Apr 22, 1987
	EQ 500MG BASE	A062809 002	Apr 22, 1987

STEVENS J	EQ 250MG BASE	A062870 001	Mar 17, 1988
	EQ 500MG BASE	A062869 001	Mar 17, 1988

SUN PHARM INDs LTD	EQ 250MG BASE	A065007 001	Sep 16, 1999
	EQ 500MG BASE	A065007 002	Sep 16, 1999

TEVA	EQ 250MG BASE	A062760 001	Apr 24, 1987
	EQ 250MG BASE	A062821 001	Feb 05, 1988
	EQ 500MG BASE	A062761 001	Apr 24, 1987

YOSHITOMI	EQ 250MG BASE	A062823 001	Feb 05, 1988
	EQ 500MG BASE	A062872 001	Jun 20, 1988

	EQ 500MG BASE	A062871 001	Jul 05, 1988
--	---------------	-------------	--------------

KEFLEX

+ PRAGMA PHARMS LLC	EQ 333MG BASE **	N050405 004	May 12, 2006
---------------------	------------------	-------------	--------------

FOR SUSPENSION; ORAL

CEPHALEXIN

APOTHECON	EQ 125MG BASE/5ML	A062986 001	Apr 18, 1991
	EQ 250MG BASE/5ML	A062987 001	Jul 25, 1989

BARR	EQ 125MG BASE/5ML	A062778 001	Aug 06, 1987
------	-------------------	-------------	--------------

	EQ 250MG BASE/5ML	A062777 001	Aug 06, 1987
FACTA FARMA	EQ 100MG BASE/ML **	A062117 001	
	EQ 125MG BASE/5ML **	A062117 002	

	EQ 250MG BASE/5ML **	A062117 003	
HIKMA PHARMS	EQ 125MG BASE/5ML	A065444 001	Aug 28, 2009
	EQ 250MG BASE/5ML	A065444 002	Aug 28, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-75(of 375)

\*\* See List Footnote

CEPHALEXIN

FOR SUSPENSION;ORAL

CEPHALEXIN

SUN PHARM INDS LTD	EQ 125MG BASE/5ML EQ 250MG BASE/5ML	A065081 001 Jul 27, 2001 A065081 002 Jul 27, 2001
TEVA	EQ 125MG BASE/5ML EQ 125MG BASE/5ML EQ 250MG BASE/5ML EQ 250MG BASE/5ML	A062767 001 Jun 16, 1987 A062873 001 May 23, 1988 A062768 001 Jun 16, 1987 A062867 001 Apr 15, 1988
VITARINE	EQ 125MG BASE/5ML EQ 250MG BASE/5ML	A062779 001 Dec 22, 1987 A062781 001 Dec 22, 1987
KEFLEX		
+ PRAGMA PHARMS LLC	EQ 100MG BASE/ML **	N050406 003
+ +	EQ 125MG BASE/5ML **	N050406 001
+ +	EQ 250MG BASE/5ML **	N050406 002
TABLET;ORAL		
CEPHALEXIN		
BARR	EQ 250MG BASE EQ 500MG BASE	A062826 001 Aug 17, 1987 A062827 001 Aug 17, 1987
VITARINE	EQ 250MG BASE EQ 500MG BASE EQ 1GM BASE	A062863 001 Aug 11, 1988 A062863 002 Aug 11, 1988 A062863 003 Aug 11, 1988
KEFLET		
LILLY	EQ 250MG BASE EQ 250MG BASE EQ 500MG BASE EQ 500MG BASE EQ 1GM BASE	A062745 001 Dec 01, 1986 N050440 003 Feb 26, 1987 A062745 002 Dec 01, 1986 N050440 001 N050440 002
TABLET, FOR SUSPENSION;ORAL		
PANIXINE DISPERDOSE		
RANBAXY LABS LTD	EQ 125MG BASE EQ 250MG BASE	A065100 002 Sep 11, 2003 A065100 001 Sep 11, 2003

CEPHALEXIN HYDROCHLORIDE

TABLET;ORAL

KEFTAB

LILLY	EQ 250MG BASE EQ 333MG BASE EQ 500MG BASE	N050614 001 Oct 29, 1987 N050614 003 May 16, 1988 N050614 002 Oct 29, 1987
-------	---	--

CEPHALOGLYCIN

CAPSULE;ORAL

KAFOCIN

LILLY	250MG	N050219 001
-------	-------	-------------

CEPHALOTHIN SODIUM

INJECTABLE;INJECTION

CEPHALOTHIN

INT'L MEDICATION	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 4GM BASE/VIAL	A062426 001 May 03, 1985 A062426 002 May 03, 1985 A062426 003 May 03, 1985 A062426 004 May 03, 1985
------------------	--	--

CEPHALOTHIN SODIUM

ABBOTT	EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 2GM BASE/VIAL	A062547 001 Sep 11, 1985 A062548 001 Sep 11, 1985 A062547 002 Sep 11, 1985 A062548 002 Sep 11, 1985
--------	--	--

ABRAXIS PHARM	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL	A062666 002 Jun 10, 1987 A062666 001 Jun 10, 1987
---------------	--------------------------------------	--

BRISTOL	EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 4GM BASE/VIAL	A062464 001 May 07, 1984 A062464 002 May 07, 1984 A062464 003 May 07, 1984
---------	--	--

CEPHALOTHIN SODIUM W/ DEXTROSE IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 20MG BASE/ML EQ 20MG BASE/ML EQ 20MG BASE/ML EQ 40MG BASE/ML EQ 40MG BASE/ML EQ 40MG BASE/ML	A062422 003 Jan 31, 1984 A062422 005 Jul 16, 1991 A062730 001 Mar 05, 1987 A062422 004 Jan 31, 1984 A062422 006 Jul 16, 1991 A062730 002 Mar 05, 1987
-----------------	--	--

**DISCONTINUED DRUG PRODUCT LIST**

6-76(of 375)

\*\* See List Footnote

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION

CEPHALOTHIN SODIUM W/ SODIUM CHLORIDE IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 20MG BASE/ML	A062422 001 Jan 31, 1984
	EQ 40MG BASE/ML	A062422 002 Jan 31, 1984

KEFLIN

LILLY	EQ 1GM BASE/VIAL	N050482 001
	EQ 2GM BASE/VIAL	N050482 002
	EQ 4GM BASE/VIAL	N050482 003
	EQ 20GM BASE/VIAL	N050482 007

KEFLIN IN PLASTIC CONTAINER

LILLY	EQ 1GM BASE/VIAL	A062549 001 Sep 10, 1985
	EQ 2GM BASE/VIAL	A062549 002 Sep 10, 1985

SEFFIN

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	A062435 001 Nov 15, 1983
	EQ 2GM BASE/VIAL	A062435 002 Nov 15, 1983
	EQ 10GM BASE/VIAL	A062435 003 Nov 15, 1983

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEFADYL

APOTHECON	EQ 500MG BASE/VIAL	A062961 001 Sep 20, 1988
	EQ 500MG BASE/VIAL	N050446 005
	EQ 1GM BASE/VIAL	A061769 001
	EQ 1GM BASE/VIAL	A062724 001 Dec 23, 1986
	EQ 1GM BASE/VIAL	A062961 002 Sep 20, 1988
	EQ 1GM BASE/VIAL	N050446 001
	EQ 2GM BASE/VIAL	A061769 002
	EQ 2GM BASE/VIAL	A062724 002 Dec 23, 1986
	EQ 2GM BASE/VIAL	A062961 003 Sep 20, 1988
	EQ 2GM BASE/VIAL	N050446 002
	EQ 4GM BASE/VIAL	A061769 003
	EQ 4GM BASE/VIAL	A062961 004 Sep 20, 1988
	EQ 4GM BASE/VIAL	N050446 003
	EQ 20GM BASE/VIAL	N050446 004

CEPHAPIRIN SODIUM

ABRAXIS PHARM	EQ 500MG BASE/VIAL	A062723 001 Nov 17, 1986
	EQ 1GM BASE/VIAL	A062723 002 Nov 17, 1986
	EQ 2GM BASE/VIAL	A062723 003 Nov 17, 1986
	EQ 4GM BASE/VIAL	A062723 004 Nov 17, 1986
	EQ 20GM BASE/VIAL	A062723 005 Nov 17, 1986
WEST-WARD PHARMS INT	EQ 500MG BASE/VIAL	A062720 001 Jul 02, 1987
	EQ 1GM BASE/VIAL	A062720 002 Jul 02, 1987
	EQ 2GM BASE/VIAL	A062720 003 Jul 02, 1987
	EQ 20GM BASE/VIAL	A062720 004 Jul 02, 1987

CEPHRADINE

CAPSULE; ORAL

ANSPOR

GLAXOSMITHKLINE	250MG	A061859 001
	500MG	A061859 002

CEPHRADINE

BARR	250MG	A062850 001 Apr 22, 1988
	500MG	A062851 001 Apr 22, 1988
IVAX SUB TEVA PHARMS	250MG	A062762 001 Mar 06, 1987
	500MG	A062762 002 Mar 06, 1987
TEVA	250MG	A062683 001 Jan 09, 1987
	500MG	A062683 002 Jan 09, 1987
VITARINE	250MG	A062813 001 Feb 25, 1988
	500MG	A062813 002 Feb 25, 1988

VELOSEF

APOTHECON	250MG	A061764 001
	500MG	A061764 002

VELOSEF '250'

ERSANA	250MG	N050548 001
--------	-------	-------------

VELOSEF '500'

ERSANA	500MG	N050548 002
--------	-------	-------------

FOR SUSPENSION; ORAL

ANSPOR

GLAXOSMITHKLINE	125MG/5ML	A061866 001
-----------------	-----------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-77(of 375)

\*\* See List Footnote

CEPHRADINEFOR SUSPENSION;ORAL  
ANSPOR

CEPHRADINE	250MG/5ML	A061866 002
BARR	125MG/5ML	A062858 001 May 19, 1988
	250MG/5ML	A062859 001 May 19, 1988
TEVA	125MG/5ML	A062693 001 Jan 09, 1987
	250MG/5ML	A062693 002 Jan 09, 1987
VELOSEF '125'		
APOTHECON	125MG/5ML	A061763 001
VELOSEF '250'		
APOTHECON	250MG/5ML	A061763 002
INJECTABLE; INJECTION		
VELOSEF		
APOTHECON	250MG/VIAL	A061976 001
	500MG/VIAL	A061976 002
	1GM/VIAL	A061976 004
	2GM/VIAL	A061976 003
	4GM/VIAL	A061976 005

TABLET;ORAL  
VELOSEF

BRISTOL MYERS SQUIBB 1GM N050530 001

CERIVASTATIN SODIUMTABLET;ORAL  
BAYCOL

BAYER PHARMS	0.05MG	N020740 001 Jun 26, 1997
	0.1MG	N020740 002 Jun 26, 1997
	0.2MG	N020740 003 Jun 26, 1997
	0.3MG	N020740 004 Jun 26, 1997
	0.4MG	N020740 005 May 24, 1999
	0.8MG	N020740 006 Jul 24, 2000

CERULETIDE DIETHYLAMINEINJECTABLE; INJECTION  
TYMTRAN

PHARMACIA AND UPJOHN 0.02MG/ML N018296 001

CETIRIZINE HYDROCHLORIDE

SYRUP;ORAL

CETIRIZINE HYDROCHLORIDE		
ACTAVIS MID ATLANTIC	5MG/5ML	A078617 001 Feb 02, 2010
APOTEX INC	5MG/5ML	A078412 001 Jun 18, 2008
AUROBINDO PHARMA LTD	5MG/5ML	A090751 001 Dec 16, 2009
RANBAXY LABS LTD	5MG/5ML	A077472 001 Jun 18, 2008
WOCKHARDT	5MG/5ML	A078757 001 Aug 28, 2009
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY		
ACTAVIS MID ATLANTIC	5MG/5ML	A090378 002 May 09, 2008
APOTEX INC	5MG/5ML	A090188 002 Apr 22, 2008
CYPRESS PHARM	5MG/5ML	A090300 001 Oct 10, 2008
RANBAXY LABS LTD	5MG/5ML	A090183 002 Apr 24, 2008

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

ACTAVIS MID ATLANTIC	5MG/5ML	A090378 001 May 09, 2008
APOTEX INC	5MG/5ML	A090188 001 Apr 22, 2008
CYPRESS PHARM	5MG/5ML	A090300 002 Oct 10, 2008
RANBAXY LABS LTD	5MG/5ML	A090183 001 Apr 24, 2008

ZYRTEC

J AND J CONSUMER INC 5MG/5ML \*\* N020346 001 Sep 27, 1996

TABLET;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY		
ACTAVIS ELIZABETH	5MG	A078615 003 Dec 28, 2007
	10MG	A078615 004 Dec 28, 2007

TABLET, CHEWABLE;ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY		
SUN PHARM INDS INC	5MG	A077631 004 Jan 11, 2008
	10MG	A077631 003 Jan 11, 2008

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

SUN PHARM INDS INC	5MG	A077631 001 Jan 11, 2008
	10MG	A077631 002 Jan 11, 2008

**DISCONTINUED DRUG PRODUCT LIST**

6-78(of 375)

\*\* See List Footnote

CETIRIZINE HYDROCHLORIDE

TABLET, CHEWABLE;ORAL		
CHILDREN'S ZYRTEC ALLERGY		
+ J AND J CONSUMER INC 5MG **	N021621 003	Nov 16, 2007
+ 10MG **	N021621 004	Nov 16, 2007
CHILDREN'S ZYRTEC HIVES RELIEF		
+ J AND J CONSUMER INC 5MG **	N021621 005	Nov 16, 2007
+ 10MG **	N021621 006	Nov 16, 2007

CETRORELIX

INJECTABLE;INJECTION		
CETROTIDE		
EMD SERONO INC EQ 3MG BASE/ML	N021197 002	Aug 11, 2000

CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION;INTRATRACHEAL		
EXOSURF NEONATAL		
GLAXOSMITHKLINE 12MG/VIAL;108MG/VIAL;8MG/VIAL	N020044 001	Aug 02, 1990

CHENODIOL

TABLET;ORAL		
CHENIX		
+ LEADIANT BIOSCI INC 250MG **	N018513 002	Jul 28, 1983

CHLOPHEDIANOL HYDROCHLORIDE

SYRUP;ORAL		
ULO		
3M 25MG/5ML	N012126 001	

CHLORAMPHENICOL

CREAM;TOPICAL		
CHLOROMYCETIN		
PARKER DAVIS 1%	N050183 001	
FOR SOLUTION;OPHTHALMIC		
CHLOROMYCETIN		
PARKADEALE 25MG/VIAL	N050143 001	
INJECTABLE;INJECTION		
CHLOROMYCETIN		
PARKE DAVIS 250MG/ML	N050153 001	
OINTMENT;OPHTHALMIC		
CHLORAMPHENICOL		
ALTANA 1%	A060133 001	
CHLOROFAIR		
PHARMAFAIR 1%	A062439 001	Apr 21, 1983
CHLOROMYCETIN		
PARKADEALE 1%	N050156 001	
CHLOROPTIC S.O.P.		
ALLERGAN 1%	A061187 001	
ECONOCHLOR		
ALCON 1%	A061648 001	
SOLUTION/DROPS;OPHTHALMIC		
CHLORAMPHENICOL		
AKORN 0.5%	A062042 001	
ALCON 0.5%	A062628 001	Sep 25, 1985
CHLOROFAIR		
PHARMAFAIR 0.5%	A062437 001	Apr 14, 1983
CHLOROPTIC		
ALLERGAN 0.5%	N050091 001	
ECONOCHLOR		
ALCON 0.5%	A061645 001	
OPHTHOCHLOR		
PARKADEALE 0.5%	A061220 001	
OPTOMYCIN		
OPTOPICS 0.5%	A062171 001	Mar 31, 1982
SOLUTION/DROPS;OTIC		
CHLOROMYCETIN		
PARKADEALE 0.5%	N050205 001	

**DISCONTINUED DRUG PRODUCT LIST**

6-79(of 375)

\*\* See List Footnote

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL

ELKINS SINK EQ 1GM BASE/VIAL

A062406 001 Nov 09, 1982

CHLORAMPHENICOL SODIUM SUCCINATE

GRUPPO LEPETIT EQ 1GM BASE/VIAL

A062278 001

CHLOROMYCETIN

+ PARKEDALE

EQ 1GM BASE/VIAL

N050155 001

MYCHEL-S

ANGUS

EQ 1GM BASE/VIAL

A060132 001

CHLORAMPHENICOL; DESOXYRIBONUCLEASE; FIBRINOLYSIN

OINTMENT; TOPICAL

ELASE-CHLOROMYCETIN

PARKE DAVIS 10MG/GM; 666 UNITS/GM; 1 UNITS/GM

N050294 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE

FOR SUSPENSION; OPHTHALMIC

CHLOROMYCETIN HYDROCORTISONE

PARKADEL 12.5MG/VIAL; 25MG/VIAL

N050202 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

OPHTHOCORT

PARKADEL 10MG/GM; 5MG/GM; 10,000 UNITS/GM

N050201 002

CHLORAMPHENICOL; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

CHLOROMYXIN

PARKE DAVIS 1%; 10,000 UNITS/GM

N050203 002

CHLORAMPHENICOL; PREDNISOLONE

OINTMENT; OPHTHALMIC

CHLOROPTIC-P S.O.P.

ALLERGAN 1%; 0.5%

A061188 001

CHLORDIAZEPOXIDE

CAPSULE, EXTENDED RELEASE; ORAL

LIBRELEASE

VALEANT PHARM INTL 30MG

N017813 001 Sep 12, 1983

TABLET; ORAL

LIBRITABS

VALEANT PHARM INTL 5MG

A085482 001

10MG

A085481 001

25MG

A085488 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

A-POXIDE

ABBOTT 5MG

A085447 001

5MG

A085517 001

10MG

A085447 002

10MG

A085518 001

25MG

A085447 003

25MG

A085513 001

CHLORDIAZACHEL

RACHELLE 5MG

A085086 001

10MG

A084639 001

25MG

A085087 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

ASCOT 5MG

A087525 001 Jan 07, 1982

10MG

A087524 001 Jan 07, 1982

25MG

A087512 001 Jan 07, 1982

FERRANTE 5MG

A085118 001

10MG

A085119 001

25MG

A085120 001

HALSEY 5MG

A085340 001

10MG

A085339 001

25MG

A084685 001

IMPAX LABS 5MG

A086213 001

10MG

A085113 001

25MG

A086212 001

IVAX SUB TEVA PHARMS 5MG

A083741 001

**DISCONTINUED DRUG PRODUCT LIST**

6-80(of 375)

\*\* See List Footnote

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

	10MG	A083742 001
	25MG	A083570 001
LEDERLE	5MG	A086892 001
	5MG	A087234 001
	10MG	A086876 001
	10MG	A087037 001
	25MG	A086893 001
	25MG	A087231 001
MAST MM	10MG	A086217 001
MYLAN	5MG	A084886 001
	10MG	A084601 001
	25MG	A084887 001
PARKE DAVIS	5MG	A085163 001
	10MG	A084598 001
	25MG	A085164 001
PIONEER PHARMS	10MG	A089533 001 Jul 15, 1988
	25MG	A089558 001 Jul 15, 1988
PUREPAC PHARM	5MG	A085155 001
	10MG	A084939 002
	25MG	A085144 001
ROXANE	5MG	A084706 001
	10MG	A084700 001
	25MG	A084705 001
SUPERPHARM	5MG	A088987 001 Apr 25, 1985
	10MG	A088986 001 Apr 25, 1985
	25MG	A088988 001 Apr 25, 1985
TEVA	5MG	A088705 001 Jan 18, 1985
	10MG	A088706 001 Jan 18, 1985
	25MG	A086494 001
	25MG	A088707 001 Jan 18, 1985
UPSHER-SMITH LABS	5MG	A084678 001
	5MG	A084919 001
	10MG	A084041 001
	10MG	A084920 001
	25MG	A084679 002
	25MG	A084823 001
USL PHARMA	5MG	A084644 001
	10MG	A084623 001
	25MG	A084645 001
VANGARD	5MG	A088129 001 Mar 28, 1983
	10MG	A088010 001 Mar 28, 1983
	25MG	A088130 001 Mar 28, 1983
WATSON LABS	5MG	A086383 001
	10MG	A086294 001
	25MG	A086382 001
WEST WARD	5MG	A085014 001
	10MG	A085000 001
	25MG	A085294 001
LIBRIUM		
VALEANT PHARM INTL	5MG	N012249 002
	10MG	N012249 001
	25MG	N012249 003
LYGEN		
ALRA	5MG	A085107 001
	10MG	A085009 001
	25MG	A085108 001

## INJECTABLE; INJECTION

LIBRIUM

VALEANT PHARMS LLC 100MG/AMP

N012301 001

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL

MENRIUM 10-4

ROCHE

10MG; 0.4MG

N014740 006

MENRIUM 5-2

ROCHE

5MG; 0.2MG

N014740 002

**DISCONTINUED DRUG PRODUCT LIST**

6-81(of 375)

\*\* See List Footnote

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET;ORAL		
MENRUM 5-4		
ROCHE	5MG; 0.4MG	N014740 004

CHLORHEXIDINE GLUCONATE

SOLUTION;DENTAL		
CHLORHEXIDINE GLUCONATE		
APOTEX INC	0.12%	A075561 001 Nov 14, 2000
SOLUTION;TOPICAL		
EXIDINE		
XTTRIUM	2.5%	N019421 001 Dec 17, 1985
MICRODERM		
J AND J	4%	A072255 001 Apr 15, 1991
PREVACARE R		
J AND J	0.5%	A072292 001 Jan 28, 1992
STERI-STAT		
MATRIX MEDCL	4%	A070104 001 Jul 24, 1986
SPONGE;TOPICAL		
CHLORHEXIDINE GLUCONATE		
KENDALL IL	4%	N019490 001 Mar 27, 1987
E-Z SCRUB		
BECTON DICKINSON	4%	A073416 001 Mar 14, 2000
HIBICLENS		
+ MOLNLYCKE HLTH	4% **	N018423 001
MICRODERM		
J AND J	4%	A072295 001 Feb 28, 1991
PHARMASEAL SCRUB CARE		
CAREFUSION 2200	4%	N019793 001 Dec 02, 1988

CHLORMERODRIN HG-197

INJECTABLE; INJECTION		
CHLORMERODRIN HG 197		
BRACCO	0.6-1.4mCi/ML	N017269 001

CHLORMEZANONE

TABLET;ORAL		
TRANCOPAL		
SANOFI AVENTIS US	100MG	N011467 003
	200MG	N011467 005

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION		
NESACAIN-E-MPF		
FRESENIUS KABI USA	2%	N009435 003
	3%	N009435 004

CHLOROQUINE HYDROCHLORIDE

INJECTABLE; INJECTION		
ARALEN HYDROCHLORIDE		
SANOFI AVENTIS US	EQ 40MG BASE/ML	N006002 002

CHLOROQUINE PHOSPHATE

TABLET;ORAL		
ARALEN		
+ SANOFI AVENTIS US	EQ 300MG BASE	N006002 001
CHLOROQUINE PHOSPHATE		
IMPAX LABS	EQ 150MG BASE	A080880 001
	EQ 300MG BASE	A040516 001 Aug 29, 2003
MD PHARM	EQ 150MG BASE	A087228 001
PUREPAC PHARM	EQ 150MG BASE	A080886 001
TEVA	EQ 150MG BASE	A087504 001 Jan 13, 1982
WATSON LABS	EQ 150MG BASE	A087979 001 Dec 21, 1982
	EQ 300MG BASE	A088030 001 Dec 21, 1982

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET;ORAL		
ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE		
SANOFI AVENTIS US	EQ 300MG BASE;EQ 45MG BASE	N014860 002

**DISCONTINUED DRUG PRODUCT LIST**

6-82(of 375)

\*\* See List Footnote

CHLOROTHIAZIDE

TABLET;ORAL

## CHLOROTHIAZIDE

ABC HOLDING	250MG	A085569 001
HIKMA INTL PHARMS	250MG	A086028 001 Jul 14, 1982
	500MG	A087736 001 Jul 14, 1982
LEDERLE	250MG	A086940 001
	500MG	A086938 001
SANDOZ	250MG	A085485 001
WATSON LABS	250MG	A085165 001
	250MG	A085173 001
	500MG	A086795 001 Aug 15, 1983
	500MG	A084026 001 Sep 01, 1982
	500MG	A086796 001 Aug 15, 1983
DIURIL		
+ OAK PHARMS AKORN	250MG **	N011145 004
+	500MG **	N011145 002

CHLOROTHIAZIDE; METHYLDOPA

TABLET;ORAL

## ALDOCLOR-150

MERCK	150MG;250MG	N016016 001
-------	-------------	-------------

## ALDOCLOR-250

MERCK	250MG;250MG	N016016 002
-------	-------------	-------------

## METHYLDOPA AND CHLOROTHIAZIDE

PAR PHARM	150MG;250MG	A070783 001 Nov 06, 1987
	250MG;250MG	A070654 001 Nov 06, 1987

CHLOROTHIAZIDE; RESERPINE

TABLET;ORAL

## CHLOROTHIAZIDE AND RESERPINE

HIKMA PHARMS	250MG;0.125MG	A088557 001 Dec 22, 1983
	500MG;0.125MG	A088365 001 Dec 22, 1983

## CHLOROTHIAZIDE W/ RESERPINE

WATSON LABS	250MG;0.125MG	A084853 001
	500MG;0.125MG	A088151 001 Jun 09, 1983

## CHLOROTHIAZIDE-RESERPINE

MYLAN	250MG;0.125MG	A087744 001 May 06, 1982
	500MG;0.125MG	A087745 001 May 06, 1982

## DIUPRES-250

MERCK	250MG;0.125MG	N011635 003 Aug 26, 1987
-------	---------------	--------------------------

## DIUPRES-500

MERCK	500MG;0.125MG	N011635 006 Aug 26, 1987
-------	---------------	--------------------------

CHLOROTRIANISENE

CAPSULE;ORAL

## CHLOROTRIANISENE

BANNER PHARMACAPS	12MG	A084652 001
TACE		
SANOFI AVENTIS US	12MG	N008102 004
	25MG	N011444 001
	72MG	N016235 001

CHLOROXINE

SHAMPOO;TOPICAL

## CAPITROL

WESTWOOD SQUIBB	2%	N017594 001
-----------------	----	-------------

CHLORPHENESIN CARBAMATE

TABLET;ORAL

## MAOLATE

PAMLAB LLC	400MG	N014217 002
------------	-------	-------------

CHLORPHENIRAMINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL

## CHLORPHENIRAMINE MALEATE

AUROLIFE PHARMA LLC	12MG	A070797 001 Aug 12, 1988
TELDRIN		
GLAXOSMITHKLINE	8MG	N017369 001
	12MG	N017369 002

**DISCONTINUED DRUG PRODUCT LIST**

6-83(of 375)

\*\* See List Footnote

CHLORPHENIRAMINE MALEATE

INJECTABLE; INJECTION

CHLOR-TRIMETON

SCHERING PLOUGH

10MG/ML

N008826 001

100MG/ML

N008794 001

CHLORPHENIRAMINE MALEATE

BEL MAR

10MG/ML

A080821 001

ELKINS SINK

10MG/ML

A080797 001

WATSON LABS

10MG/ML

A083593 001

10MG/ML

A086096 001

100MG/ML

A086095 001

PYRIDAMAL 100

BEL MAR

100MG/ML

A083733 001

SYRUP; ORAL

CHLOR-TRIMETON

SCHERING

2MG/5ML

N006921 006

CHLORPHENIRAMINE MALEATE

PHARM ASSOC

2MG/5ML

A087520 001 Feb 10, 1982

TABLET; ORAL

ANTAGONATE

BAYER PHARMS

4MG

A083381 001

CHLOR-TRIMETON

SCHERING

4MG

N006921 002

CHLORPHENIRAMINE MALEATE

ANABOLIC

4MG

A083078 001

AUROLIFE PHARMA LLC

4MG

A080961 001

BELL PHARMA

4MG

A083062 001

ELKINS SINK

4MG

A080938 001

IMPAX LABS

4MG

A080809 001

IVAX SUB TEVA PHARMS

4MG

A080779 001

KV PHARM

4MG

A087164 001

LEDERLE

4MG

A086941 001

NEWTRON PHARMS

4MG

A086519 001

PANRAY

4MG

A083243 001

PHARMAVITE

4MG

A085104 001

PHARMERAL

4MG

A083753 001

PIONEER PHARMS

4MG

A088556 001 Jul 13, 1984

PUREPAC PHARM

4MG

A086306 001

PVT FORM

4MG

A080786 001

ROXANE

4MG

A080626 001

SUN PHARM INDUSTRIES

4MG

A080700 001

VITARINE

4MG

A085837 001

WATSON LABS

4MG

A080696 001

4MG

A080791 001

4MG

A085139 001

WEST WARD

4MG

A083787 001

KLOROMIN

HALSEY

4MG

A083629 001

PHENETRON

LANNETT

4MG

A080846 001

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

BAYER HEALTHCARE LLC 8MG

N007638 001

EFIDAC 24 CHLORPHENIRAMINE MALEATE

ALZA 16MG

N019746 002 Nov 18, 1994

CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE

TABLET, EXTENDED RELEASE; ORAL

CODEINE PHOSPHATE AND CHLORPHENIRAMINE MALEATE

SPRIASO LLC 8MG;54.3MG

N206323 001 Jun 22, 2015

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE

TRIS PHARMA INC 4MG/5ML;5MG/5ML

A206438 001 Jan 27, 2015

**DISCONTINUED DRUG PRODUCT LIST**

6-84(of 375)

\*\* See List Footnote

CHLORPHENIRAMINE MALEATE; HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE

TRIS PHARMA INC 4MG/5ML;5MG/5ML;60MG/5ML A203838 001 Nov 26, 2014

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

COLD CAPSULE IV

GRAHAM DM 12MG;75MG

N018793 001 Apr 25, 1985

COLD CAPSULE V

GRAHAM DM 8MG;75MG

N018794 001 Apr 23, 1985

TABLET, EXTENDED RELEASE;ORAL

TRIAMINIC-12

NOVARTIS 12MG;75MG

N018115 001

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

CODIMAL-L.A. 12

SCHWARZ PHARMA 12MG;120MG

N018935 001 Apr 15, 1985

ISOCLOR

FISONS 8MG;120MG

N018747 001 Mar 06, 1986

PSEUDOEPHEDRINE HYDROCHLORIDE AND CHLORPHENIRAMINE MALEATE

CENT PHARMS 8MG;120MG

N019428 001 Aug 02, 1988

GRAHAM DM 8MG;120MG

N018844 001 Mar 20, 1985

12MG;120MG

N018843 001 Mar 18, 1985

KV PHARM 12MG;120MG

A071455 001 Mar 01, 1989

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL

CHLOR-TRIMETON

+ BAYER HEALTHCARE LLC 8MG;120MG

N018397 001

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL

CODEPREX

UCB INC EQ 4MG MALEATE/5ML;EQ 20MG BASE/5ML

N021369 001 Jun 21, 2004

PENNTUSS

FISONS EQ 4MG MALEATE/5ML;EQ 10MG BASE/5ML

N018928 001 Aug 14, 1985

CHLORPHENTERMINE HYDROCHLORIDE

TABLET;ORAL

PRE-SATE

PARKE DAVIS EQ 65MG BASE

N014696 001

CHLORPROMAZINE

SUPPOSITORY;RECTAL

THORAZINE

+ GLAXOSMITHKLINE 25MG \*\*  
+ 100MG \*\*

N009149 024

N009149 033

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

THORAZINE

GLAXOSMITHKLINE 30MG  
75MG  
150MG  
200MG  
300MG

N011120 016

N011120 017

N011120 018

N011120 019

N011120 020

CONCENTRATE;ORAL

CHLORPROMAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC 100MG/ML  
PHARM ASSOC 30MG/ML  
100MG/ML  
WOCKHARDT 30MG/ML  
100MG/ML

A086863 001

A040231 001 Dec 30, 1999

A040224 001 Jan 26, 1999

A087032 001 Jul 08, 1982

A087053 001

CHLORPROMAZINE HYDROCHLORIDE INTENSOL

CYCLE PHARMS LTD 30MG/ML  
100MG/ML

A088157 001 Apr 27, 1983

A088158 001 Apr 27, 1983

SONAZINE

SANDOZ 30MG/ML  
100MG/ML

A080983 004

A080983 005

**DISCONTINUED DRUG PRODUCT LIST**

6-85(of 375)

\*\* See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

THORAZINE

+ GLAXOSMITHKLINE	30MG/ML **	N009149 032
+ GLAXOSMITHKLINE	100MG/ML **	N009149 043

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

ABRAXIS PHARM	25MG/ML	A084911 001
MARSAM PHARMS LLC	25MG/ML	A089563 001 Apr 15, 1988
WATSON LABS	25MG/ML	A080365 001
	25MG/ML	A085591 001
WYETH AYERST	25MG/ML	A080370 001

THORAZINE

+ GLAXOSMITHKLINE	25MG/ML **	N009149 011
-------------------	------------	-------------

SYRUP; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	A086712 001
SONAZINE	10MG/5ML	A083040 001
SANDOZ	10MG/5ML	A083040 001
THORAZINE	10MG/5ML **	N009149 022

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ABBOTT	10MG	A084414 001
	25MG	A084415 001
	50MG	A084411 001
	100MG	A084412 001
	200MG	A084413 001
CYCLE PHARMS LTD	10MG	A085331 001
	25MG	A085331 002
	50MG	A085331 003
	100MG	A085331 004
	200MG	A085331 005
IVAX SUB TEVA PHARMS	10MG	A083549 001
	25MG	A083549 002
	50MG	A083549 003
	100MG	A083574 001
	200MG	A083575 001
KV PHARM	10MG	A085750 002 Jan 04, 1982
	25MG	A085751 001
	50MG	A085484 001
	100MG	A085752 001
	200MG	A085748 002 Jan 04, 1982
LEDERLE	10MG	A084803 001
	25MG	A084801 001
	50MG	A084800 001
	100MG	A084789 001
	200MG	A084802 001
PUREPAC PHARM	10MG	A080403 004
	25MG	A080403 001
	50MG	A080403 002
	100MG	A080403 003
	200MG	A080403 005
PVT FORM	25MG	A080340 001
	50MG	A080340 002
	200MG	A080340 003
SANDOZ	10MG **	A080439 001
	25MG **	A080439 002
	50MG **	A080439 003
	100MG **	A080439 004
	200MG **	A080439 005
VANGARD	10MG	A088038 001 Aug 16, 1982
	25MG	A087645 001
	50MG	A087646 001
WATSON LABS	10MG	A085959 001
	25MG	A085956 001
	50MG	A085960 001
	100MG	A085957 001
	200MG	A085958 001

**DISCONTINUED DRUG PRODUCT LIST**

6-86(of 375)

\*\* See List Footnote

CHLORPROMAZINE HYDROCHLORIDE

TABLET;ORAL

CHLORPROMAZINE HYDROCHLORIDE

WEST WARD	10MG	A087783 001	Sep 16, 1982
	25MG	A087865 001	Sep 16, 1982
	50MG	A087878 001	Sep 15, 1982
	100MG	A087884 001	Sep 15, 1982
	200MG	A087880 001	Sep 16, 1982
PROMAPAR			
PARKE DAVIS	10MG	A086886 001	
	25MG	A084423 001	
	50MG	A086887 001	
	100MG	A086888 001	
	200MG	A086885 001	
THORAZINE			
GLAXOSMITHKLINE	10MG **	N009149 002	
	25MG **	N009149 007	
	50MG **	N009149 013	
	100MG **	N009149 018	
	200MG **	N009149 020	

CHLORPROPAMIDE

TABLET;ORAL

CHLORPROPAMIDE

ANI PHARMS INC	100MG	A088768 001	Oct 11, 1984
	100MG	A088812 001	Oct 19, 1984
	100MG	A088840 001	Oct 25, 1984
	100MG	A088918 001	Oct 16, 1984
	100MG	A089446 001	Nov 17, 1986
	250MG	A087353 001	
	250MG	A088813 001	Oct 19, 1984
	250MG	A088919 001	Oct 16, 1984
	250MG	A089447 001	Nov 17, 1986
AUROLIFE PHARMA LLC	100MG	A088725 001	Aug 31, 1984
	250MG	A088726 001	Aug 31, 1984
DAVA PHARMS INC	100MG	A089561 001	Sep 04, 1987
	250MG	A089562 001	Sep 04, 1987
HALSEY	100MG	A089321 001	Jan 16, 1986
	250MG	A088662 001	Jan 09, 1986
PAR PHARM	100MG	A088175 001	Feb 27, 1984
	250MG	A088176 001	Feb 27, 1984
SANDOZ	250MG	A084669 001	
SUPERPHARM	100MG	A088694 001	Sep 17, 1984
	250MG	A088695 001	Sep 17, 1984
USL PHARMA	100MG	A088708 001	Aug 30, 1984
	250MG	A088709 001	Aug 30, 1984
WATSON LABS	100MG	A086865 001	Sep 24, 1984
	100MG	A088608 001	Apr 12, 1984
	250MG	A086866 001	
	250MG	A088568 001	Apr 12, 1984
WATSON LABS TEVA	100MG	A088852 001	Sep 26, 1984
	250MG	A088826 001	Sep 26, 1984

CHLORPROTHIXENE

CONCENTRATE;ORAL

TARACTAN

ROCHE	100MG/5ML	N016149 002
-------	-----------	-------------

INJECTABLE;INJECTION

TARACTAN

ROCHE	12.5MG/ML	N012487 001
-------	-----------	-------------

TABLET;ORAL

TARACTAN

ROCHE	10MG	N012486 005
	25MG	N012486 004
	50MG	N012486 003
	100MG	N012486 001

**DISCONTINUED DRUG PRODUCT LIST**

6-87(of 375)

\*\* See List Footnote

CHLORTETRACYCLINE HYDROCHLORIDE

OINTMENT;OPHTHALMIC

AUREOMYCIN

LEDERLE

1%

N050404 001

CHLORTHALIDONE

TABLET;ORAL

CHLORTHALIDONE

ABBOTT

25MG

A087364 001

50MG

A087384 001

ASCOT

25MG

A087698 001 Oct 20, 1982

50MG

A087699 001 Oct 20, 1982

BARR LABS INC

25MG

A088902 001 Sep 19, 1985

50MG

A088903 001 Sep 19, 1985

DAVA PHARMS INC

25MG

A087451 001

50MG

A087450 001

G AND W LABS INC

50MG

A088651 001 May 30, 1985

IVAX PHARMS

25MG

A087555 001

25MG

A088164 001 Jan 09, 1984

50MG

A087176 001

KV PHARM

25MG

A087947 001 Feb 27, 1984

50MG

A087311 001

MUTUAL PHARM

25MG

A087312 001

25MG

A087292 001

50MG

A089738 001 Sep 19, 1988

50MG

A087293 001

PIONEER PHARMS

50MG

A089739 001 Sep 19, 1988

PUREPAC PHARM

25MG

A089591 001 Jul 21, 1988

50MG

A088139 001 Jul 16, 1986

SANDOZ

25MG

A088140 001 Aug 11, 1983

50MG

A087380 001

50MG

A087118 001

SUPERPHARM

25MG

A087381 001

50MG

A087473 001 Feb 09, 1983

USL PHARMA

25MG

A087247 001 Feb 09, 1983

50MG

A089051 001 Jun 01, 1987

VANGARD

25MG

A089052 001 Jun 01, 1987

50MG

A088012 001 Jul 14, 1982

WARNER CHILCOTT

25MG

A088073 001 Mar 25, 1983

50MG

A087515 001 Jan 24, 1983

WATSON LABS

25MG

A087516 001 Feb 09, 1983

25MG

A087050 001

25MG

A087100 001

25MG

A087296 001

50MG

A087706 001

50MG

A087029 001

50MG

A087082 001

50MG

A087521 001

50MG

A087689 001

HYGROTON

+ SANOFI AVENTIS US 25MG \*\*

N012283 004

+ 50MG \*\*

N012283 003

THALITONE

CASPER PHARMA LLC

15MG \*\*

N019574 001 Dec 20, 1988

25MG

N019574 002 Feb 12, 1992

MONARCH PHARMS

25MG

A088051 001 Nov 12, 1982

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE AND CHLORTHALIDONE

PAR PHARM

15MG;0.1MG

A071179 001 Dec 16, 1987

15MG;0.2MG

A071178 001 Dec 16, 1987

15MG;0.3MG

A071142 001 Dec 16, 1987

COMBIPRES

+ BOEHRINGER INGELHEIM 15MG;0.1MG \*\*

N017503 001

+ 15MG;0.2MG \*\*

N017503 002

+ 15MG;0.3MG \*\*

N017503 003 Apr 10, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-88(of 375)

\*\* See List Footnote

CHLORTHALIDONE; METOPROLOL TARTRATE

CAPSULE;ORAL

LOPRESSIDONE

NOVARTIS

25MG;100MG

25MG;200MG

N019451 001 Dec 31, 1987

N019451 002 Dec 31, 1987

CHLORTHALIDONE; RESERPINE

TABLET;ORAL

DEMI-REGROTOM

SANOFI AVENTIS US

25MG;0.125MG

N015103 002

REGROTOM

SANOFI AVENTIS US

50MG;0.25MG

N015103 001

CHLORZOXAZONE

TABLET;ORAL

CHLORZOXAZONE

ACTAVIS ELIZABETH

250MG

A088928 001 May 08, 1987

500MG

A040113 001 Sep 29, 1995

AUROLIFE PHARMA LLC

250MG

A089852 001 May 04, 1988

500MG

A089853 001 May 04, 1988

OHM LABS

250MG

A081298 001 Dec 29, 1993

500MG

A081299 001 Dec 29, 1993

PAR PHARM

250MG

A087981 001 Sep 20, 1983

PIONEER PHARMS

250MG

A089592 001 Jan 06, 1989

500MG

A089948 001 Jan 06, 1989

SUN PHARM INDUSTRIES

500MG

A089970 001 Sep 27, 1990

WATSON LABS

250MG

A086901 001

250MG

A086948 001 Aug 09, 1982

500MG

A040137 001 Aug 09, 1996

500MG

A081019 001 Jul 29, 1991

500MG

A081040 001 Aug 22, 1989

PARAFLEX

+ ORTHO MCNEIL PHARM 250MG \*\*

N011300 003

PARAFON FORTE DSC

+ JANSEN R AND D 500MG \*\*

N011529 002 Jun 15, 1987

STRIFON FORTE DSC

FERNDALE LABS 500MG

A081008 001 Dec 23, 1988

CHOLESTYRAMINE

BAR, CHEWABLE;ORAL

CHOLYBAR

PARKE DAVIS

EQ 4GM RESIN/BAR

A071621 001 May 26, 1988

EQ 4GM RESIN/BAR

A071739 001 May 26, 1988

POWDER;ORAL

CHOLESTYRAMINE

ANI PHARMS INC

EQ 4GM RESIN/PACKET

A074554 001 Oct 02, 1996

EQ 4GM RESIN/SCOOPFUL

A074554 002 Oct 02, 1996

IVAX SUB TEVA PHARMS

EQ 4GM RESIN/PACKET

A074771 001 Jul 09, 1997

EQ 4GM RESIN/SCOOPFUL

A074771 002 Jul 09, 1997

TEVA

EQ 4GM RESIN/PACKET

A074347 001 May 28, 1998

EQ 4GM RESIN/SCOOPFUL

A074347 002 May 28, 1998

CHOLESTYRAMINE LIGHT

TEVA

EQ 4GM RESIN/PACKET

A074348 001 May 28, 1998

EQ 4GM RESIN/SCOOPFUL

A074348 002 May 28, 1998

TEVA PHARMS

EQ 4GM RESIN/PACKET

A074555 001 Sep 30, 1998

EQ 4GM RESIN/SCOOPFUL

A074555 002 Sep 30, 1998

LOCHOLEST

SANDOZ

EQ 4GM RESIN/PACKET

A074561 001 Aug 15, 1996

EQ 4GM RESIN/SCOOPFUL

A074561 002 Aug 15, 1996

LOCHOLEST LIGHT

SANDOZ

EQ 4GM RESIN/PACKET

A074562 001 Aug 15, 1996

EQ 4GM RESIN/SCOOPFUL

A074562 002 Aug 15, 1996

QUESTRAN

+ BRISTOL MYERS

EQ 4GM RESIN/PACKET \*\*

N016640 001

+

EQ 4GM RESIN/SCOOPFUL \*\*

N016640 003

QUESTRAN LIGHT

+ BRISTOL MYERS

EQ 4GM RESIN/PACKET \*\*

N019669 001 Dec 05, 1988

+

EQ 4GM RESIN/SCOOPFUL \*\*

N019669 003 Dec 05, 1988

**DISCONTINUED DRUG PRODUCT LIST**

6-89(of 375)

\*\* See List Footnote

CHOLESTYRAMINETABLET;ORAL  
QUESTRANAPOTHECON  
EQ 800MG RESIN  
EQ 1GM RESINA073403 002 Dec 27, 1999  
A073403 001 Apr 28, 1994CHORIOGONADOTROPIN ALFAINJECTABLE;INJECTION  
OVIDREL

EMD SERONO 0.25MG/VIAL

N021149 001 Sep 20, 2000

CHROMIC CHLORIDEINJECTABLE;INJECTION  
CHROMIC CHLORIDE

ABRAXIS PHARM EQ 0.004MG CHROMIUM/ML

N019271 001 May 05, 1987

CHROMIC PHOSPHATE P-32INJECTABLE;INJECTION  
PHOSPHOCOL P32

MALLINKRODT NUCLEAR 5mCi/ML

N017084 001

CHYMOPAPAIN

INJECTABLE;INJECTION

CHYMODIACTIN

CHART MEDCL 4,000 UNITS/VIAL  
+ 10,000 UNITS/VIAL \*\*N018663 002 Aug 21, 1984  
N018663 001 Nov 10, 1982

DISCASE

ABBOTT 12,500 UNITS/VIAL

N018625 001 Jan 18, 1984

CHYMOTRYPSIN

FOR SOLUTION;OPHTHALMIC

ALPHA CHYMAR

SOLA BARNES HIND 750 UNITS/VIAL

N011837 001

CATARASE

CIBA 300 UNITS/VIAL

N016938 001

NOVARTIS 150 UNITS/VIAL

N018121 001

ZOLYSE

ALCON 750 UNITS/VIAL

N011903 001

CICLOPIROX

SOLUTION;TOPICAL

CICLOPIROX

MYLAN PHARMS INC 8%  
TEVA PHARMS 8%A078567 001 Sep 18, 2007  
A078079 001 Sep 18, 2007CIDOFOVIR

INJECTABLE;INJECTION

VISTIDE

+ GILEAD SCIENCES INC EQ 75MG BASE/ML \*\*

N020638 001 Jun 26, 1996

CILASTATIN SODIUM; IMIPENEM

INJECTABLE;INJECTION

PRIMAXIN

MERCK EQ 250MG BASE/VIAL;250MG/VIAL  
EQ 500MG BASE/VIAL;500MG/VIALA062756 001 Jan 08, 1987  
A062756 002 Jan 08, 1987

POWDER;INTRAMUSCULAR

PRIMAXIN

MERCK EQ 500MG BASE/VIAL;500MG/VIAL  
EQ 750MG BASE/VIAL;750MG/VIALN050630 001 Dec 14, 1990  
N050630 002 Dec 14, 1990CILOSTAZOL

TABLET;ORAL

CILOSTAZOL

ACTAVIS ELIZABETH 100MG  
EPIC PHARMA LLC 50MG  
FRONTIDA BIOPHARM 100MG  
IVAX SUB TEVA PHARMS 100MG  
MYLAN 50MG  
MYLAN PHARMS INC 50MG  
PLIVA HRVATSKA DOO 50MGA077028 002 Nov 26, 2004  
A077150 001 Mar 11, 2005  
A077022 001 Nov 23, 2004  
A077208 002 Mar 29, 2006  
A077208 001 Mar 29, 2006  
A077020 002 Mar 01, 2005  
A077323 002 Apr 20, 2006  
A077323 001 Apr 20, 2006  
A077019 001 Nov 23, 2004  
A077019 002 Nov 23, 2004  
A077898 001 Oct 29, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-90(of 375)

\*\* See List Footnote

**CILOSTAZOL**

TABLET;ORAL  
CILOSTAZOL

PLETAL	100MG	A077898 002 Oct 29, 2007
+ OTSUKA	50MG **	N020863 001 Jan 15, 1999
+	100MG **	N020863 002 Jan 15, 1999

**CIMETIDINE**

SUSPENSION;ORAL

TAGAMET HB 200	200MG/20ML	N020951 001 Jul 09, 1999
----------------	------------	--------------------------

TABLET;ORAL  
CIMETIDINE

CHARTWELL MOLECULES	200MG	A074281 001 May 17, 1994
	300MG	A074281 002 May 17, 1994
	400MG	A074281 003 May 17, 1994
	800MG	A074329 001 May 17, 1994
CONTRACT PHARMACAL	200MG	A074961 001 Jun 19, 1998
	200MG	A074963 001 Jun 19, 1998
CYCLE PHARMS LTD	300MG	A074361 001 Dec 23, 1994
	400MG	A074361 002 Dec 23, 1994
	800MG	A074371 001 Dec 23, 1994
DAVA PHARMS INC	300MG	A074340 001 Jun 23, 1995
	400MG	A074340 002 Jun 23, 1995
	800MG	A074339 001 Jun 23, 1995
FOSUN PHARMA	200MG	A074100 001 Jan 31, 1995
	300MG	A074100 002 Jan 31, 1995
	400MG	A074100 003 Jan 31, 1995
	800MG	A074100 004 Jan 31, 1995
IVAX SUB TEVA PHARMS	200MG	A074401 001 May 30, 1995
	200MG	A074424 001 Jul 28, 1995
	300MG	A074401 002 May 30, 1995
	300MG	A074424 002 Jul 28, 1995
	400MG	A074401 003 May 30, 1995
	400MG	A074424 003 Jul 28, 1995
	800MG	A074402 001 May 30, 1995
	800MG	A074424 004 Jul 28, 1995
PERRIGO	100MG	A074972 001 Jun 19, 1998
PLIVA	200MG	A074568 001 Feb 27, 1997
	300MG	A074568 002 Feb 27, 1997
	400MG	A074568 003 Feb 27, 1997
SANDOZ INC	100MG	A075122 001 Jun 19, 1998
	200MG	A074250 001 Jun 29, 1995
	200MG	A075122 002 Jun 19, 1998
	300MG	A074250 002 Jun 29, 1995
	400MG	A074250 003 Jun 29, 1995
	800MG	A074250 004 Jun 29, 1995
TEVA	200MG	A074365 001 Feb 28, 1995
	300MG	A074365 002 Feb 28, 1995
	400MG	A074365 003 Feb 28, 1995
	800MG	A074365 004 Feb 28, 1995
UPSHER-SMITH LABS	200MG	A074506 001 Jan 24, 1996
	300MG	A074506 002 Jan 24, 1996
	400MG	A074506 003 Jan 24, 1996
	800MG	A074506 004 Jan 24, 1996
WATSON LABS INC	200MG	A074349 001 Aug 30, 1996
	300MG	A074349 002 Aug 30, 1996
	400MG	A074349 003 Aug 30, 1996
	800MG	A074316 001 Feb 28, 1996
WATSON LABS TEVA	200MG	A075425 001 Jul 29, 1999
TAGAMET		
GLAXOSMITHKLINE	200MG **	N017920 002
	300MG **	N017920 003
	400MG **	N017920 004 Dec 14, 1983
	800MG **	N017920 005 Apr 30, 1986
TAGAMET HB		
+ MEDTECH PRODUCTS	100MG **	N020238 001 Jun 19, 1995

**DISCONTINUED DRUG PRODUCT LIST**

6-91(of 375)

\*\* See List Footnote

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

HOSPIRA	EQ 300MG BASE/2ML	A074296 001	Mar 28, 1997
	EQ 300MG BASE/2ML	A074344 001	Jan 31, 1995
	EQ 300MG BASE/2ML	A074345 001	Jan 31, 1995
	EQ 300MG BASE/2ML	A074412 001	Mar 28, 1997
	EQ 300MG BASE/2ML	A074422 001	Jan 31, 1995
LUITPOLD	EQ 300MG BASE/2ML	A074353 001	Dec 20, 1994
TEVA PARENTERAL	EQ 300MG BASE/2ML	A074252 001	Nov 26, 1997
VINTAGE PHARMS LLC	EQ 300MG BASE/2ML	A074005 001	Aug 31, 1994
CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
HOSPIRA	EQ 6MG BASE/ML	A074269 001	Dec 27, 1994
	EQ 90MG BASE/100ML	A074468 005	Dec 29, 1994
	EQ 120MG BASE/100ML	A074468 006	Dec 29, 1994
	EQ 180MG BASE/100ML	A074468 003	Dec 29, 1994
	EQ 240MG BASE/100ML	A074468 004	Dec 29, 1994
	EQ 360MG BASE/100ML	A074468 001	Dec 29, 1994
	EQ 480MG BASE/100ML	A074468 002	Dec 29, 1994
TAGAMET			
GLAXOSMITHKLINE	EQ 300MG BASE/2ML **	N017939 002	
TAGAMET HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
+ GLAXOSMITHKLINE	EQ 6MG BASE/ML **	N019434 001	Oct 31, 1985
SOLUTION; ORAL			
CIMETIDINE HYDROCHLORIDE			
ANI PHARMS INC	EQ 300MG BASE/5ML	A074859 001	Jul 09, 1998
	EQ 300MG BASE/5ML	A075110 001	Jun 18, 1998
APOTEX INC	EQ 300MG BASE/5ML	A075560 001	Mar 15, 2000
CYCLE PHARMS LTD	EQ 300MG BASE/5ML	A074541 001	Aug 05, 1997
G AND W LABS INC	EQ 300MG BASE/5ML	A074176 001	Jun 01, 1994
VINTAGE PHARMS LLC	EQ 300MG BASE/5ML	A074251 001	Dec 22, 1994
TAGAMET			
GLAXOSMITHKLINE	EQ 300MG BASE/5ML **	N017924 001	

CINOXACIN

CAPSULE; ORAL

CINOBAC

LILLY	250MG	N018067 001
	500MG	N018067 002
CINOXACIN		
TEVA	250MG	A073005 001
	500MG	A073006 001

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

+ BAYER HLTHCARE	400MG/40ML (10MG/ML)	N019847 001	Dec 26, 1990
+	200MG/20ML (10MG/ML)	N019847 002	Dec 26, 1990
	1200MG/120ML (10MG/ML)	N019847 003	Dec 26, 1990

CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER

+ BAYER HLTHCARE	200MG/100ML	N019857 001	Dec 26, 1990
+	400MG/200ML	N019857 002	Dec 26, 1990

CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

BAYER PHARMS	200MG/100ML	N019858 001	Dec 26, 1990
--------------	-------------	-------------	--------------

CIPROFLOXACIN

BEDFORD LABS	200MG/20ML (10MG/ML)	A076992 001	Aug 28, 2006
	400MG/40ML (10MG/ML)	A076992 002	Aug 28, 2006
	1200MG/120ML (10MG/ML)	A076993 001	Aug 28, 2006

FRESENIUS KABI USA	200MG/20ML (10MG/ML)	A076484 001	Aug 28, 2006
	400MG/40ML (10MG/ML)	A076484 002	Aug 28, 2006

TEVA PHARMS USA	200MG/20ML (10MG/ML)	A077782 001	Aug 28, 2006
	400MG/40ML (10MG/ML)	A077782 002	Aug 28, 2006

CIPROFLOXACIN IN DEXTROSE 5%

HIKMA FARMACEUTICA	200MG/100ML	A076757 001	Apr 21, 2008
--------------------	-------------	-------------	--------------

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	200MG/100ML	A077888 001	Mar 18, 2008
	400MG/200ML	A077888 002	Mar 18, 2008

BEDFORD	200MG/100ML	A078114 001	Mar 18, 2008
	400MG/200ML	A078114 002	Mar 18, 2008

TEVA PHARMS	200MG/100ML	A077138 001	Mar 18, 2008
-------------	-------------	-------------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-92(of 375)

\*\* See List Footnote

**CIPROFLOXACIN**

INJECTABLE; INJECTION

CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER  
400MG/200ML

A077138 002 Mar 18, 2008

**CIPROFLOXACIN HYDROCHLORIDE**

SOLUTION/DROPS; OPHTHALMIC

CIPROFLOXACIN HYDROCHLORIDE

AMRING PHARMS EQ 0.3% BASE  
APOTEX INC EQ 0.3% BASEA078598 001 Jan 16, 2008  
A075928 001 Jun 09, 2004

TABLET; ORAL

CIPROFLOXACIN HYDROCHLORIDE

BARR EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA074124 001 Jun 09, 2004  
A074124 002 Jun 09, 2004  
A074124 003 Jun 09, 2004IDT AUSTRALIA LTD EQ 100MG BASE  
EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA075939 001 Mar 03, 2005  
A075939 002 Jun 09, 2004  
A075939 003 Jun 09, 2004  
A075939 004 Jun 09, 2004MYLAN EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA075685 002 Jun 09, 2004  
A075685 003 Jun 09, 2004NOSTRUM LABS EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA076138 001 Jun 09, 2004  
A076138 002 Jun 09, 2004  
A076138 003 Jun 09, 2004PLIVA EQ 100MG BASE  
EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA076426 001 Jun 15, 2005  
A076426 002 Jun 15, 2005  
A076426 003 Jun 15, 2005  
A076426 004 Jun 15, 2005SANDOZ EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA076593 002 Jun 09, 2004  
A076593 003 Jun 09, 2004TEVA EQ 250MG BASE  
EQ 500MG BASE  
EQ 750MG BASEA076136 001 Jun 09, 2004  
A076136 002 Jun 09, 2004  
A076136 003 Jun 09, 2004

TABLET, EXTENDED RELEASE; ORAL

PROQUIN XR

DEPOMED INC EQ 500MG BASE

N021744 001 May 19, 2005

**CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE**

TABLET, EXTENDED RELEASE; ORAL

CIPRO XR

BAYER HLTHCARE 212.6MG; EQ 287.5MG BASE \*\*  
425.2MG; EQ 574.9MG BASE \*\*N021473 001 Dec 13, 2002  
N021473 002 Aug 28, 2003

CIPROFLOXACIN EXTENDED RELEASE

ACTAVIS LABS FL INC 212.6MG; EQ 287.5MG BASE  
425.2MG; EQ 574.9MG BASE  
DR REDDYS LABS LTD 212.6MG; EQ 287.5MG BASE  
SANDOZ 212.6MG; EQ 287.5MG BASEA077417 001 Nov 30, 2010  
A077809 001 Nov 30, 2010  
A077701 002 Oct 31, 2007  
A078712 001 Dec 11, 2007**CISAPRIDE MONOHYDRATE**

SUSPENSION; ORAL

PROPULSID

JANSSEN PHARMS EQ 1MG BASE/ML

N020398 001 Sep 15, 1995

TABLET; ORAL

PROPULSID

JANSSEN PHARMS EQ 10MG BASE  
EQ 20MG BASEN020210 001 Jul 29, 1993  
N020210 002 Dec 23, 1993

TABLET, ORALLY DISINTEGRATING; ORAL

PROPULSID QUICKSOLV

JANSSEN PHARMA EQ 20MG BASE

N020767 001 Nov 07, 1997

**CISPLATIN**

INJECTABLE; INJECTION

CISPLATIN

BEDFORD 10MG/VIAL  
50MG/VIAL  
TEVA PHARMS USA 1MG/ML  
PLATINOL + HQ SPCLT PHARMA 10MG/VIAL  
+ 50MG/VIALA074713 001 Nov 14, 2000  
A074713 002 Nov 14, 2000  
A074814 001 May 16, 2000  
N018057 001  
N018057 002

**DISCONTINUED DRUG PRODUCT LIST**

6-93(of 375)

\*\* See List Footnote

CISPLATIN

INJECTABLE; INJECTION

PLATINOL-AQ

+ HQ SPCLT PHARMA

0.5MG/ML

N018057 003 Jul 18, 1984

CITALOPRAM HYDROBROMIDE

CAPSULE; ORAL

CITALOPRAM HYDROBROMIDE

MYLAN PHARMS INC

EQ 10MG BASE

A077668 001 Feb 28, 2007

EQ 20MG BASE

A077668 002 Feb 28, 2007

EQ 40MG BASE

A077668 003 Feb 28, 2007

SOLUTION; ORAL

CELEXA

+ FOREST LABS

EQ 10MG BASE/5ML \*\*

N021046 001 Dec 22, 1999

CITALOPRAM HYDROBROMIDE

APOTEX INC

EQ 10MG BASE/5ML

A077601 001 Nov 15, 2005

TABLET; ORAL

CELEXA

FOREST LABS

EQ 60MG BASE

N020822 004 Jul 17, 1998

CITALOPRAM HYDROBROMIDE

ACTAVIS ELIZABETH

EQ 10MG BASE

A077033 001 Oct 28, 2004

EQ 20MG BASE

A077033 002 Oct 28, 2004

EQ 40MG BASE

A077033 003 Oct 28, 2004

EPIC PHARMA LLC

EQ 10MG BASE

A077036 001 Oct 28, 2004

EQ 20MG BASE

A077036 002 Oct 28, 2004

EQ 40MG BASE

A077036 003 Oct 28, 2004

MYLAN

EQ 10MG BASE

A077039 001 Feb 03, 2005

EQ 20MG BASE

A077039 002 Feb 03, 2005

EQ 40MG BASE

A077039 003 Feb 03, 2005

MYLAN PHARMS INC

EQ 10MG BASE

A077037 001 Nov 05, 2004

EQ 20MG BASE

A077037 002 Nov 05, 2004

EQ 40MG BASE

A077037 003 Nov 05, 2004

NATCO PHARMA LTD

EQ 20MG BASE

A077141 002 Apr 10, 2008

EQ 40MG BASE

A077141 001 Apr 10, 2008

ROXANE

EQ 10MG BASE

A077041 001 Nov 23, 2004

EQ 20MG BASE

A077041 002 Nov 23, 2004

EQ 40MG BASE

A077041 003 Nov 23, 2004

SANDOZ

EQ 10MG BASE

A077035 001 Oct 28, 2004

EQ 10MG BASE

A077040 001 Aug 17, 2005

EQ 20MG BASE

A077035 002 Oct 28, 2004

EQ 20MG BASE

A077040 002 Aug 17, 2005

EQ 40MG BASE

A077035 003 Oct 28, 2004

EQ 40MG BASE

A077040 003 Aug 17, 2005

SUN PHARM INDUSTRIES

EQ 10MG BASE

A077052 001 Jul 03, 2006

EQ 20MG BASE

A077052 002 Jul 03, 2006

EQ 40MG BASE

A077052 003 Jul 03, 2006

TARO

EQ 10MG BASE

A077278 001 Mar 22, 2006

EQ 20MG BASE

A077278 002 Mar 22, 2006

EQ 40MG BASE

A077278 003 Mar 22, 2006

TEVA PHARMS

EQ 10MG BASE

A077213 001 Mar 31, 2006

EQ 20MG BASE

A077213 002 Mar 31, 2006

EQ 40MG BASE

A077213 003 Mar 31, 2006

WATSON LABS

EQ 10MG BASE

A077034 001 Jun 30, 2005

EQ 20MG BASE

A077034 002 Jun 30, 2005

EQ 40MG BASE

A077034 003 Jun 30, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

CITALOPRAM HYDROBROMIDE

BIOVAIL LABS INTL

EQ 10MG BASE

N021763 001 Dec 20, 2005

EQ 20MG BASE

N021763 002 Dec 20, 2005

EQ 40MG BASE

N021763 003 Dec 20, 2005

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

SOLUTION; IRRIGATION

IRRIGATING SOLUTION G IN PLASTIC CONTAINER

BAXTER HLTHCARE 3.24GM/100ML;380MG/100ML;430MG/100ML

N018519 001 Jun 22, 1982

UROLOGIC G IN PLASTIC CONTAINER

HOSPIRA

3.24GM/100ML;380MG/100ML;430MG/100ML

N018904 001 May 27, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-94(of 375)

\*\* See List Footnote

CLADRIBINE

INJECTABLE; INJECTION

LEUSTATIN

+ JANSSEN PHARMS

1MG/ML \*\*

N020229 001 Feb 26, 1993

CLARITHROMYCIN

FOR SUSPENSION; ORAL

BIAXIN

ABBVIE

187MG/5ML

N050698 003 Sep 30, 1998

TABLET; ORAL

CLARITHROMYCIN

IVAX SUB TEVA PHARMS 250MG

A065137 001 May 31, 2005

500MG

A065137 002 May 31, 2005

MYLAN 250MG

A065195 001 Mar 11, 2005

500MG

A065195 002 Mar 11, 2005

TABLET, EXTENDED RELEASE; ORAL

BIAXIN XL

+ ABBVIE

500MG \*\*

N050775 001 Mar 03, 2000

CLARITHROMYCIN

IDT AUSTRALIA LTD 500MG

A065250 001 Aug 25, 2005

RANBAXY 1GM

A065210 001 Jan 26, 2005

CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TIMENTIN

GLAXOSMITHKLINE

EQ 100MG BASE/VIAL; EQ 3GM BASE/VIAL

A062691 001 Dec 19, 1986

EQ 100MG BASE/VIAL; EQ 3GM BASE/VIAL

N050590 001 Apr 01, 1985

EQ 200MG BASE/VIAL; EQ 3GM BASE/VIAL

N050590 002 Apr 01, 1985

EQ 1GM BASE/VIAL; EQ 30GM BASE/VIAL

N050590 003 Aug 18, 1987

TIMENTIN IN PLASTIC CONTAINER

GLAXOSMITHKLINE

EQ 100MG BASE/100ML; EQ 3GM BASE/100ML

N050658 001 Dec 15, 1989

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

ACTAVIS MID ATLANTIC EQ 0.5MG BASE/5ML

A074075 001 Oct 31, 1993

APOTEX INC

EQ 0.5MG BASE/5ML

A075703 001 Nov 27, 2000

SILARX

EQ 0.5MG BASE/5ML

A074884 001 Dec 17, 1997

TEVA PHARMS

EQ 0.5MG BASE/5ML

A073095 001 Apr 21, 1992

TAVIST

+ NOVARTIS

EQ 0.5MG BASE/5ML \*\*

N018675 001 Jun 28, 1985

TABLET; ORAL

CLEMASTINE FUMARATE

ANI PHARMS INC

1.34MG

A073282 001 Jan 31, 1992

1.34MG

A073282 002 Dec 03, 1992

SANDOZ

2.68MG

A073459 001 Oct 31, 1993

TAVIST

NOVARTIS

2.68MG

N017661 001

TAVIST-1

NOVARTIS

1.34MG

N017661 002

NOVARTIS

1.34MG

N017661 003 Aug 21, 1992

CLIDINIUM BROMIDE

CAPSULE; ORAL

QUARZAN

ROCHE

2.5MG

N010355 001

5MG

N010355 002

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 75MG BASE

A061809 001

EQ 150MG BASE

A061809 002

CLINDAMYCIN HYDROCHLORIDE

MYLAN PHARMS INC

EQ 75MG BASE

A091225 001 May 31, 2011

EQ 150MG BASE

A091225 002 May 31, 2011

EQ 300MG BASE

A091225 003 May 31, 2011

TEVA

EQ 75MG BASE

A063027 001 Sep 20, 1989

WATSON LABS

EQ 75MG BASE

A063082 001 Jul 31, 1991

**DISCONTINUED DRUG PRODUCT LIST**

6-95(of 375)

\*\* See List Footnote

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION;ORAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 75MG BASE/5ML \*\*

A061827 001

CLINDAMYCIN PHOSPHATE

CREAM;VAGINAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 2% BASE

N050680 001 Aug 11, 1992

INJECTABLE;INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN EQ 150MG BASE/ML

A061839 001

CLINDAMYCIN PHOSPHATE

ABRAXIS PHARM EQ 150MG BASE/ML

A062747 001 Jun 03, 1988

BEDFORD EQ 150MG BASE/ML

A063163 001 Jun 30, 1994

BRISTOL MYERS SQUIBB EQ 150MG BASE/ML

A062908 001 Feb 01, 1989

IGI LABS INC EQ 150MG BASE/ML

A062928 001 Feb 13, 1989

LOCH EQ 150MG BASE/ML

A062905 001 May 09, 1988

MARSAM PHARMS LLC EQ 150MG BASE/ML

A062913 001 Oct 20, 1988

SOLOPAK EQ 150MG BASE/ML

A062819 001 Mar 15, 1988

EQ 150MG BASE/ML

A062852 001 Mar 17, 1988

TEVA PARENTERAL EQ 150MG BASE/ML

A063041 001 Dec 29, 1989

EQ 150MG BASE/ML

A063282 001 May 29, 1992

WATSON LABS EQ 150MG BASE/ML

A062900 001 Jun 08, 1988

EQ 150MG BASE/ML

A063079 001 Mar 05, 1990

WEST-WARD PHARMS INT EQ 150MG BASE/ML

A062806 001 Oct 15, 1987

EQ 150MG BASE/ML

A062953 001 Apr 21, 1988

EQ 150MG BASE/ML

A063068 001 Aug 28, 1989

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

ABRAXIS PHARM EQ 12MG BASE/ML

N050636 001 Dec 22, 1989

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBVIE EQ 6MG BASE/ML

A065027 001 Jun 29, 2001

EQ 12MG BASE/ML

A065027 002 Jun 29, 2001

EQ 18MG BASE/ML

A065027 003 Jun 29, 2001

BAXTER HLTHCARE EQ 6MG BASE/ML

N050648 001 Dec 29, 1989

EQ 12MG BASE/ML

N050648 002 Dec 29, 1989

EQ 900MG BASE/100ML

N050648 003 Dec 29, 1989

SOLUTION;TOPICAL

CLEOCIN T

PHARMACIA AND UPJOHN EQ 1% BASE

A062363 001 Feb 08, 1982

CLINDAMYCIN PHOSPHATE

BOCA PHARMA LLC EQ 1% BASE

A062944 001 Jan 11, 1989

NOVAST LABS LTD EQ 1% BASE

A064108 001 Sep 27, 1996

VINTAGE PHARMS EQ 1% BASE

A062930 001 Jun 28, 1989

CLIQUINOL; NYSTATIN

OINTMENT;TOPICAL

NYSTAFORM

BAYER PHARMS 10MG/GM;100,000 UNITS/GM

N050235 001

CLOBAZAM

TABLET;ORAL

ONFI

LUNDBECK PHARMS LLC 5MG

N202067 001 Oct 21, 2011

CLOBETASOL PROPIONATE

CREAM;TOPICAL

CLOBETASOL PROPIONATE

TEVA PHARMS USA 0.05%

A074087 001 Feb 16, 1994

CLOBETASOL PROPIONATE (EMOLLIENT)

NOVAST LABS LTD 0.05%

A075733 001 Aug 22, 2001

TEMOVATE

+ FOUGERA PHARMS 0.05% \*\*

N019322 001 Dec 27, 1985

TEMOVATE E

+ FOUGERA PHARMS 0.05% \*\*

N020340 001 Jun 17, 1994

GEL;TOPICAL

TEMOVATE

+ FOUGERA PHARMS 0.05% \*\*

N020337 001 Apr 29, 1994

OINTMENT;TOPICAL

CLOBETASOL PROPIONATE

ACTAVIS MID ATLANTIC 0.05%

A074128 001 Aug 03, 1994

**DISCONTINUED DRUG PRODUCT LIST**

6-96(of 375)

\*\* See List Footnote

CLOBETASOL PROPIONATE

OINTMENT;TOPICAL

TEMOVATE

+ FOUGERA PHARMS

0.05% \*\*

N019323 001 Dec 27, 1985

SOLUTION;TOPICAL

TEMOVATE

+ FOUGERA PHARMS

0.05% \*\*

N019966 001 Feb 22, 1990

CLOFAZIMINE

CAPSULE;ORAL

LAMPRENE

+ NOVARTIS

50MG

100MG

N019500 002 Dec 15, 1986

N019500 001 Dec 15, 1986

CLOFIBRATE

CAPSULE;ORAL

ATROMID-S

WYETH AYERST

500MG

N016099 002

CLOFIBRATE

BANNER PHARMACAPS

500MG

A073396 001 Mar 20, 1992

SANDOZ

500MG

A072191 001 May 02, 1988

TEVA

500MG

A072600 001 Jul 25, 1991

USL PHARMA

500MG

A070531 001 Jun 16, 1986

WATSON LABS

500MG

A071603 001 Sep 18, 1987

CLOMIPHENE CITRATE

TABLET;ORAL

MILOPHENE

MILEX

50MG

A072196 001 Dec 20, 1988

SEROPHENE

EMD SERONO

50MG

N018361 001 Mar 22, 1982

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

CLOMIPRAMINE HYDROCHLORIDE

TEVA

25MG

A074849 001 Apr 04, 1997

50MG

A074849 002 Apr 04, 1997

75MG

A074849 003 Apr 04, 1997

WATSON LABS

25MG

A074600 001 Nov 27, 1996

25MG

A074751 001 Sep 30, 1998

50MG

A074600 002 Nov 27, 1996

50MG

A074751 002 Sep 30, 1998

75MG

A074600 003 Nov 27, 1996

75MG

A074751 003 Sep 30, 1998

CLONAZEPAM

TABLET;ORAL

CLONAZEPAM

APOTEX INC

0.5MG

A075468 001 Oct 06, 2000

1MG

A075468 002 Oct 06, 2000

2MG

A075468 003 Oct 06, 2000

MYLAN PHARMS INC

0.5MG

A074940 001 Oct 30, 1997

1MG

A074940 002 Oct 30, 1997

2MG

A074940 003 Oct 30, 1997

SANDOZ

0.5MG

A074925 001 Sep 30, 1997

1MG

A074925 002 Sep 30, 1997

2MG

A074925 003 Sep 30, 1997

TEVA

0.5MG

A074920 001 Aug 04, 1998

1MG

A074920 002 Aug 04, 1998

2MG

A074920 003 Aug 04, 1998

KLOONOPIN

ROCHE

0.125MG

N017533 005 Apr 09, 1997

0.25MG

N017533 006 Apr 09, 1997

TABLET, ORALLY DISINTEGRATING;ORAL

KLONOPIN RAPIDLY DISINTEGRATING

+ ROCHE

0.125MG \*\*

N020813 001 Dec 23, 1997

+

0.25MG \*\*

N020813 002 Dec 23, 1997

+

0.5MG \*\*

N020813 003 Dec 23, 1997

+

1MG \*\*

N020813 004 Dec 23, 1997

+

2MG \*\*

N020813 005 Dec 23, 1997

**DISCONTINUED DRUG PRODUCT LIST**

6-97(of 375)

\*\* See List Footnote

CLONIDINESUSPENSION, EXTENDED RELEASE;ORAL  
CLONIDINE

TRIS PHARMA INC	EQ 0.09MG BASE/ML	N022499 001 Dec 03, 2009
TABLET, EXTENDED RELEASE;ORAL		
CLONIDINE		
TRIS PHARMA INC	EQ 0.17MG BASE	N022500 001 Dec 03, 2009
	EQ 0.26MG BASE	N022500 002 Dec 03, 2009

CLONIDINE HYDROCHLORIDE

TABLET;ORAL

CLONIDINE HYDROCHLORIDE

AM THERAP	0.1MG	A070881 001 Jul 08, 1986
	0.2MG	A070882 001 Jul 08, 1986
	0.3MG	A070883 001 Jul 08, 1986
AUROLIFE PHARMA LLC	0.1MG	A070887 001 Aug 31, 1988
	0.2MG	A070886 001 Aug 31, 1988
	0.3MG	A071294 001 Aug 31, 1988
CHARTWELL MOLECULES	0.3MG	A071785 001 Apr 05, 1988
DURAMED PHARMS BARR	0.1MG	A071103 001 Aug 14, 1986
	0.2MG	A071102 001 Aug 14, 1986
	0.3MG	A071101 001 Aug 14, 1986
INTERPHARM	0.1MG	A071252 001 Oct 01, 1986
	0.2MG	A071253 001 Oct 01, 1986
	0.3MG	A071254 001 Oct 01, 1986
PAR PHARM	0.1MG	A070461 001 Jul 08, 1986
	0.2MG	A070460 001 Jul 08, 1986
	0.3MG	A070459 001 Jul 08, 1986
TEVA	0.1MG	A070747 001 Jul 08, 1986
	0.2MG	A070702 001 Jul 08, 1986
	0.3MG	A070659 001 Jul 08, 1986
WARNER CHILCOTT	0.1MG	A072138 001 Jun 13, 1988
	0.2MG	A072139 001 Jun 13, 1988
	0.3MG	A072140 001 Jun 13, 1988
WATSON LABS	0.1MG	A070395 001 Mar 23, 1987
	0.1MG	A070965 001 Jul 08, 1986
	0.2MG	A070396 001 Mar 23, 1987
	0.2MG	A070964 001 Jul 08, 1986
	0.3MG	A070397 001 Mar 23, 1987
	0.3MG	A070963 001 Jul 08, 1986

TABLET, EXTENDED RELEASE;ORAL

CLONIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH	0.2MG	A202792 002 May 15, 2015
	0.2MG	A203320 002 May 15, 2015
ANCHEN PHARMS	0.1MG	A202983 001 Apr 02, 2014
	0.2MG	A202983 002 Apr 02, 2014

CLONIDINE HYDROCHLORIDE

ANCHEN PHARMS 0.2MG

A202984 002 Sep 30, 2013

JENLOGA

+ CONCORDIA PHARMS INC 0.1MG \*\*  
+ 0.2MG \*\*

N022331 001 Sep 30, 2009

N022331 002 May 25, 2010

KAPVAY

+ CONCORDIA PHARMS INC 0.2MG \*\*

N022331 004 Sep 28, 2010

CLOPIDOGREL BISULFATE

TABLET;ORAL

CLOPIDOGREL BISULFATE

ACTAVIS TOTOWA EQ 75MG BASE

A090307 001 May 28, 2013

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	A071777 001 Jul 14, 1987
	7.5MG	A071778 001 Jul 14, 1987
	15MG	A071779 001 Jul 14, 1987
AM THERAP	3.75MG	A071429 001 Jun 23, 1987
	7.5MG	A071430 001 Jun 23, 1987
	15MG	A071431 001 Jun 23, 1987
AUROLIFE PHARMA LLC	15MG	A072112 001 Aug 26, 1988
DAVA PHARMS INC	3.75MG	A071742 001 Dec 14, 1987
	7.5MG	A071743 001 Dec 14, 1987

**DISCONTINUED DRUG PRODUCT LIST**

6-98(of 375)

\*\* See List Footnote

CLORAZEPATE DIPOTASSIUM

CAPSULE;ORAL

CLORAZEPATE DIPOTASSIUM

	15MG	A071744 001	Dec 14, 1987
GD SEARLE LLC	3.75MG	A071727 001	Dec 18, 1987
	7.5MG	A071728 001	Dec 18, 1987
	15MG	A071729 001	Dec 18, 1987
MYLAN	3.75MG	A071509 001	Oct 19, 1987
	7.5MG	A071510 001	Oct 19, 1987
	15MG	A071511 001	Oct 19, 1987
PUREPAC PHARM	3.75MG	A071924 001	Apr 25, 1988
	7.5MG	A071925 001	Apr 25, 1988
	15MG	A071926 001	Apr 25, 1988
QUANTUM PHARMICS	3.75MG	A071549 001	Sep 12, 1988
	7.5MG	A071550 001	Sep 12, 1988
	15MG	A071522 001	Sep 12, 1988
USL PHARMA	3.75MG	A071242 001	Jun 23, 1987
	7.5MG	A071243 001	Jun 23, 1987
	15MG	A071244 001	Jun 23, 1987
WARNER CHILCOTT	3.75MG	A071774 001	Mar 01, 1988
	7.5MG	A071775 001	Mar 01, 1988
	15MG	A071776 001	Mar 01, 1988
WATSON LABS	3.75MG	A071878 001	Mar 15, 1988
	7.5MG	A071879 001	Mar 15, 1988
	15MG	A071860 001	Mar 15, 1988
TRANXENE			
RECORDATI RARE	3.75MG	N017105 001	
	7.5MG	N017105 002	
	15MG	N017105 003	

## TABLET;ORAL

CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	A071780 001	Jun 26, 1987
	7.5MG	A071781 001	Jun 26, 1987
	15MG	A071782 001	Jun 26, 1987
AM THERAP	3.75MG	A071747 001	Jun 23, 1987
	7.5MG	A071748 001	Jun 23, 1987
	15MG	A071749 001	Jun 23, 1987
AUROLIFE PHARMA LLC	3.75MG	A072512 001	May 11, 1990
	7.5MG	A072513 001	May 11, 1990
	15MG	A072514 001	May 11, 1990
LEDERLE	3.75MG	A072013 001	Dec 15, 1987
	7.5MG	A072014 001	Dec 15, 1987
	15MG	A072015 001	Dec 15, 1987
PUREPAC PHARM	3.75MG	A072330 001	Aug 08, 1988
	7.5MG	A072331 001	Aug 08, 1988
	15MG	A072332 001	Aug 08, 1988
QUANTUM PHARMICS	3.75MG	A071730 001	Oct 26, 1987
	7.5MG	A071731 001	Oct 26, 1987
	15MG	A071702 001	Oct 26, 1987
SUN PHARM INDS LTD	3.75MG	A076911 001	Sep 29, 2004
	7.5MG	A076911 002	Sep 29, 2004
	15MG	A076911 003	Sep 29, 2004
WARNER CHILCOTT	3.75MG	A071828 001	Mar 03, 1988
	7.5MG	A071829 001	Mar 03, 1988
	15MG	A071830 001	Mar 03, 1988
WATSON LABS	3.75MG	A071852 001	Feb 09, 1988
	7.5MG	A071853 001	Feb 09, 1988
	15MG	A071854 001	Feb 09, 1988
TRANXENE			
RECORDATI RARE	3.75MG	N017105 006	
TRANXENE SD			
RECORDATI RARE	11.25MG	N017105 005	
	22.5MG	N017105 004	

**DISCONTINUED DRUG PRODUCT LIST**

6-99(of 375)

\*\* See List Footnote

CLOTRIMAZOLE

CREAM;TOPICAL			
LOTRIMIN			
SCHERING PLOUGH	1%		N017619 001
MYCELEX			
BAYER HEALTHCARE LLC	1%		N018183 001
CREAM;VAGINAL			
GYNE-LOTRIMIN			
+ BAYER HEALTHCARE LLC	1%		N018052 002 Nov 30, 1990
GYNE-LOTRIMIN 3			
+ BAYER HEALTHCARE LLC	2%		N020574 001 Nov 24, 1998
CREAM, TABLET;TOPICAL, VAGINAL			
GYNE-LOTRIMIN 3 COMBINATION PACK			
+ BAYER HEALTHCARE LLC	1%,200MG		N020526 002 Jul 29, 1996
GYNE-LOTRIMIN COMBINATION PACK			
+ BAYER HEALTHCARE LLC	1%,100MG		N020289 002 Apr 26, 1993
LOTION;TOPICAL			
LOTRIMIN			
SCHERING	1%		N018813 001 Feb 17, 1984
SOLUTION;TOPICAL			
LOTRIMIN			
+ SCHERING PLOUGH	1%		N017613 001
MYCELEX			
+ BAYER HLTHCARE	1% **		N018181 001
TABLET;VAGINAL			
GYNE-LOTRIMIN			
+ BAYER HEALTHCARE LLC	100MG		N017717 002 Nov 30, 1990
GYNE-LOTRIMIN 3			
+ BAYER HEALTHCARE LLC	200MG		N020525 001 Jul 29, 1996
GYNIX			
TEVA PHARMS	100MG		A073249 001 Feb 13, 1998
MYCELEX-G			
BAYER PHARMS	500MG		N019069 001 Apr 19, 1985
TROCHE/LOZENGE;ORAL			
MYCELEX			
+ BAYER HLTHCARE	10MG **		N018713 001 Jun 17, 1983

CLOXACILLIN SODIUM

CAPSULE;ORAL			
CLOXACILLIN SODIUM			
APOTHECON	EQ 250MG BASE		A061452 001
	EQ 500MG BASE		A061452 002
TEVA	EQ 250MG BASE		A062240 001
	EQ 500MG BASE		A062240 002
CLOXAPEN			
GLAXOSMITHKLINE	EQ 250MG BASE		A061806 001
	EQ 250MG BASE		A062233 001
	EQ 500MG BASE		A061806 002
	EQ 500MG BASE		A062233 002
FOR SOLUTION;ORAL			
CLOXACILLIN SODIUM			
TEVA	EQ 125MG BASE/5ML		A062268 001
	EQ 125MG BASE/5ML		A062978 001 Apr 06, 1989
TEGOPEN			
APOTHECON	EQ 125MG BASE/5ML		A061453 001
	EQ 125MG BASE/5ML		N050192 001

CLOZAPINE

TABLET;ORAL			
CLOZAPINE			
PAR PHARM	25MG		A075162 001 Apr 26, 2005
	100MG		A075162 002 Apr 26, 2005
SANDOZ	25MG		A074546 001 Aug 30, 1996
	100MG		A074546 002 Aug 30, 1996
TABLET, ORALLY DISINTEGRATING;ORAL			
FAZACLO ODT			
JAZZ PHARMS III	50MG		N021590 003 Jun 03, 2005

**DISCONTINUED DRUG PRODUCT LIST**

6-100(of 375)

\*\* See List Footnote

COBALT CHLORIDE CO-57; CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A;N/A

RUBRATOPE-57 KIT

BRACCO

N/A;N/A;N/A;N/A

N016089 001

COBALT CHLORIDE CO-60; CYANOCOBALAMIN; CYANOCOBALAMIN CO-60; INTRINSIC FACTOR

N/A;N/A

RUBRATOPE-60 KIT

BRACCO

N/A;N/A;N/A;N/A

N016090 001

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN VC W/ CODEINE

+ ANI PHARMS

10MG/5ML;5MG/5ML;6.25MG/5ML \*\*

N008306 005 Apr 02, 1984

PHERAZINE VC W/ CODEINE

HALSEY

10MG/5ML;5MG/5ML;6.25MG/5ML

A088870 001 Mar 02, 1987

PROMETHAZINE VC W/ CODEINE

CENCI

10MG/5ML;5MG/5ML;6.25MG/5ML

A088816 001 Nov 22, 1985

WOCKHARDT

10MG/5ML;5MG/5ML;6.25MG/5ML

A088896 001 Jan 04, 1985

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN W/ CODEINE

+ ANI PHARMS

10MG/5ML;6.25MG/5ML \*\*

N008306 004 Apr 02, 1984

PHERAZINE W/ CODEINE

HALSEY

10MG/5ML;6.25MG/5ML

A088739 001 Dec 23, 1988

PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE

PHARM ASSOC

10MG/5ML;6.25MG/5ML

A089647 001 Dec 22, 1988

PROMETHAZINE W/ CODEINE

CENCI

10MG/5ML;6.25MG/5ML

A088814 001 Nov 22, 1985

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIFED W/ CODEINE

GLAXOSMITHKLINE

10MG/5ML;30MG/5ML;1.25MG/5ML

N012575 003 Apr 04, 1984

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES W/ CODEINE

CENCI

10MG/5ML;30MG/5ML;1.25MG/5ML

A089018 001 Jul 23, 1986

TRIPROLIDINE HCL, PSEUDOEPHEDRINE HCL AND CODEINE PHOSPHATE

WOCKHARDT

10MG/5ML;30MG/5ML;1.25MG/5ML

A088833 001 Nov 16, 1984

CODEINE SULFATE

SOLUTION;ORAL

CODEINE SULFATE

WEST-WARD PHARMS INT 30MG/5ML

N202245 001 Jun 30, 2011

COLCHICINE; PROBENECID

TABLET;ORAL

COLBENEMID

+ MERCK

0.5MG;500MG \*\*

N012383 001

PROBEN-C

WATSON LABS

0.5MG;500MG

A085552 001

PROBENECID AND COLCHICINE

ANI PHARMS INC

0.5MG;500MG

A083734 001

BEECHAM

0.5MG;500MG

A084321 001

IMPAK LABS

0.5MG;500MG

A083720 002

SANDOZ

0.5MG;500MG

A086130 001

PROBENECID W/ COLCHICINE

LEDERLE

0.5MG;500MG

A086954 001

WATSON LABS

0.5MG;500MG

A083221 001

COLESEVELAM HYDROCHLORIDE

CAPSULE;ORAL

WELCHOL

DAIICHI SANKYO

375MG

N021141 001 May 26, 2000

COLISTIN SULFATE

SUSPENSION;ORAL

COLY-MYCIN S

PARKE DAVIS

EQ 25MG BASE/5ML

N050355 001

**DISCONTINUED DRUG PRODUCT LIST**

6-101(of 375)

\*\* See List Footnote

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; IV (INFUSION)

VAPRISOL

CUMBERLAND PHARMS 20MG/4ML (5MG/ML)

N021697 001 Dec 29, 2005

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE

CU-7

GD SEARLE LLC 89MG

N017408 001

TATUM-T

GD SEARLE LLC 120MG

N018205 001

CORTICOTROPIN

INJECTABLE; INJECTION

ACTH

PARKEDALE 25 UNITS/VIAL  
40 UNITS/VIALN008317 002  
N008317 004

ACTHAR

SANOFI AVENTIS US 25 UNITS/VIAL  
40 UNITS/VIALN007504 002  
N007504 003

CORTICOTROPIN

ORGANICS LAGRANGE 40 UNITS/ML  
80 UNITS/MLN010831 001  
N010831 002

WATSON LABS 40 UNITS/VIAL

A088772 001 Nov 21, 1984

H.P. ACTHAR GEL

MALLINCKRODT ARD 40 UNITS/ML

N008372 006

PURIFIED CORTROPHIN GEL

ANI PHARMS INC 40 UNITS/ML  
80 UNITS/MLN008975 001  
N008975 002CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION

CORTROPHIN-ZINC

ANI PHARMS INC 40 UNITS/ML

N009854 001

CORTISONE ACETATE

INJECTABLE; INJECTION

CORTISONE ACETATE

PHARMACIA AND UPJOHN 25MG/ML  
WATSON LABS 25MG/ML  
25MG/ML  
50MG/ML  
50MG/MLN008126 002  
A083147 003  
A085677 001  
A083147 004  
A085677 002

CORTONE

MERCK 25MG/ML  
50MG/MLN007110 002  
N007110 003

TABLET; ORAL

CORTISONE ACETATE

BARR 25MG  
ELKINS SINK 25MG  
EVERYLIFE 25MG  
HEATHER 25MG  
IMPAX LABS 25MG  
INWOOD LABS 25MG  
IVAX SUB TEVA PHARMS 25MG  
25MG  
LANNETT 25MG  
PANRAY 5MG  
25MG  
PHARMACIA AND UPJOHN 5MG  
10MG  
25MG  
PUREPAC PHARM 25MG  
VITARINE 25MG  
WATSON LABS 25MG  
WHITEWORTH TOWN PLSN 25MGA083471 001  
A080836 001  
A084246 001  
A085736 001  
N009458 001  
A080731 001  
A080630 001  
A083536 001  
A080694 001  
N008284 002  
N008284 001  
N008126 003  
N008126 004  
N008126 001  
A080493 001  
A080333 001  
A085884 001  
A080341 001

CORTONE

+ MERCK 25MG \*\*

N007750 003

**DISCONTINUED DRUG PRODUCT LIST**

6-102(of 375)

\*\* See List Footnote

COSYNTROPIN

SOLUTION; INTRAVENOUS  
 COSYNTROPIN  
 SANDOZ INC 0.25MG/ML (0.25MG/ML) N022028 001 Feb 21, 2008

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION  
 INTAL  
 KING PHARMS LLC 0.8MG/INH N018887 001 Dec 05, 1985

CAPSULE; INHALATION  
 INTAL  
 + SANOFI AVENTIS US 20MG \*\* N016990 001

CAPSULE; ORAL  
 GASTROCROM  
 UCB INC 100MG N019188 001 Dec 22, 1989

CONCENTRATE; ORAL  
 CROMOLYN SODIUM  
 GENERA PHARMS 100MG/5ML A090954 001 Dec 18, 2009

SOLUTION; INHALATION  
 CROMOLYN SODIUM  
 ACTAVIS MID ATLANTIC 10MG/ML A075067 001 Jul 19, 1999  
 APOTEX INC 10MG/ML A075333 001 Apr 30, 2002  
 BAUSCH AND LOMB 10MG/ML A075585 001 Dec 21, 2000  
 PHARMASCIENCE INC 10MG/ML A075437 001 Apr 21, 2000  
 ROXANE 10MG/ML A075175 001 Sep 30, 1999  
 WATSON LABS 10MG/ML A076469 001 Jun 17, 2005

INTAL  
 + KING PHARMS LLC 10MG/ML \*\* N018596 001 May 28, 1982

SOLUTION/DROPS; OPHTHALMIC  
 CROLOM  
 BAUSCH AND LOMB 4% A074443 001 Jan 30, 1995

CROMOLYN SODIUM  
 APOTEX INC 4% A075615 001 Jan 26, 2001

CROMOPTIC  
 KING PHARMS 4% A075088 001 Apr 27, 1999

OPTICROM  
 + ALLERGAN 4% \*\* N018155 001 Oct 03, 1984

SPRAY, METERED; NASAL  
 CROMOLYN SODIUM  
 ACTAVIS MID ATLANTIC 5.2MG/SPRAY A074800 001 Jul 26, 2001  
 HH AND P 5.2MG/SPRAY A077976 001 Sep 07, 2007

NASALCROM  
 + BLACKSMITH BRANDS 5.2MG/SPRAY \*\* N020463 001 Jan 03, 1997

CRYPTENAMINE ACETATES

INJECTABLE; INJECTION  
 UNITENSEN  
 MEDPOINTE PHARM HLC 260CSR UNIT/ML N008814 001

CRYPTENAMINE TANNATES

TABLET; ORAL  
 UNITENSEN  
 MEDPOINTE PHARM HLC 260CSR UNIT N009217 001

CUPRIC SULFATE

INJECTABLE; INJECTION  
 CUPRIC SULFATE  
 ABRAXIS PHARM EQ 0.4MG COPPER/ML N019350 001 May 05, 1987

CYANOCOBALAMIN

GEL, METERED; NASAL  
 NASCOBAL  
 PAR PHARM 0.5MG/INH N019722 001 Nov 05, 1996

INJECTABLE; INJECTION  
 BERUBIGEN  
 PHARMACIA AND UPJOHN 1MG/ML N006798 001

BETALIN 12  
 LILLY 0.1MG/ML A080855 001  
 1MG/ML A080855 002

COBAVITE  
 WATSON LABS 0.1MG/ML A083013 001  
 1MG/ML A083064 001

**DISCONTINUED DRUG PRODUCT LIST**

6-103(of 375)

\*\* See List Footnote

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMIN

ABRAXIS PHARM	0.03MG/ML 0.1MG/ML 1MG/ML	A080510 003 A080510 001 A080510 002
AKORN	1MG/ML	A087969 001 Nov 10, 1983
DELL LABS	0.03MG/ML 0.1MG/ML 1MG/ML	A080689 001 A080689 002 A080689 003
FRESENIUS KABI USA	0.1MG/ML	A080557 002
LUITPOLD	0.03MG/ML	A080668 001
LYPHOMED	1MG/ML	A083075 001
MYLAN INSTITUTIONAL	1MG/ML	A040451 001 Sep 23, 2003
SANOFI AVENTIS US	1MG/ML	A080564 001
SOLOPAK	1MG/ML	A087551 001 Feb 29, 1984
WARNER CHILCOTT	1MG/ML	N007085 002
WATSON LABS	0.1MG/ML 0.1MG/ML 1MG/ML 1MG/ML	A080573 002 A083120 001 A080573 001 A083120 002
WYETH AYERST	0.1MG/ML 1MG/ML	A080554 001 A080554 002
DODEX		
ACCORD HLTHCARE	1MG/ML	A083022 001
REDISOL		
MERCK	1MG/ML	N006668 010
RUBIVITE		
BEL MAR	0.03MG/ML 0.05MG/ML 0.1MG/ML 0.12MG/ML 1MG/ML	N010791 004 N010791 001 N010791 002 N010791 005 N010791 003
RUBRAMIN PC		
BRISTOL MYERS SQUIBB	0.1MG/ML	N006799 002
+	1MG/ML **	N006799 004
+	1MG/ML **	N006799 010 Apr 28, 1988
RVITONE		
SAVAGE LABS	1MG/ML	A080570 002
VI-TWEL		
BAYER HLTHCARE	1MG/ML	N007012 002
SPRAY, METERED;NASAL		
CALOMIST		
PAR PHARM	25MCG/SPRAY	N022102 001 Jul 27, 2007
TABLET;ORAL		
CYANOCOBALAMIN		
WEST WARD	1MG	A084264 001

CYANOCOBALAMIN CO-57

CAPSULE;ORAL

RUBRATOPE-57

BRACCO

0.5-1uCi

N016089 002

CYANOCOBALAMIN CO-60

CAPSULE;ORAL

RUBRATOPE-60

BRACCO

0.5-1uCi

N016090 002

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; CYANOCOBALAMIN CO-58

N/A;N/A

DICOPAC KIT

GE HEALTHCARE

N/A;N/A;N/A

N017406 001

CYANOCOBALAMIN; CYANOCOBALAMIN CO-57; INTRINSIC FACTOR

N/A;N/A

CYANOCOBALAMIN CO 57 SCHILLING TEST KIT

MALLINCKRODT

0.1MG;0.5uCi;60MG

N016635 001

**DISCONTINUED DRUG PRODUCT LIST**

6-104(of 375)

\*\* See List Footnote

CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR

ARMOUR PHARM

0.5MG/ML; 2.3MG/ML; 1MG/ML

N011208 001

CYCLACILLIN

FOR SUSPENSION; ORAL

CYCLAPEN-W

WYETH AYERST

125MG/5ML

N050508 001

250MG/5ML

N050508 002

500MG/5ML

N050508 003

TABLET; ORAL

CYCLACILLIN

TEVA

250MG

A062895 001 Aug 04, 1988

500MG

A062895 002 Aug 04, 1988

CYCLAPEN-W

WYETH AYERST

250MG

N050509 001

500MG

N050509 002

CYCLIZINE LACTATE

INJECTABLE; INJECTION

MAREZINE

GLAXOSMITHKLINE

50MG/ML

N009495 001

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

TWI PHARMS INC

15MG

A091281 001 Jan 31, 2013

30MG

A091281 002 Jan 31, 2013

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

SANDOZ

10MG

A073683 001 Feb 26, 1993

UPSHER-SMITH LABS

5MG

A072854 002 Feb 03, 2006

10MG

A072854 001 Nov 19, 1991

WATSON LABS

10MG

A073143 001 Nov 27, 1991

10MG

A074436 001 Nov 30, 1994

FLEXERIL

+ JANSSEN RES AND DEV

5MG \*\*

N017821 001

+

10MG \*\*

N017821 002

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

AKORN

1%

A085555 001

AKPENTOLATE

AKORN

2%

A040165 001 Jan 13, 1997

CYCLOPENTOLATE HYDROCHLORIDE

ALCON PHARMS LTD

1%

A089162 001 Jan 24, 1991

SOLA BARNES HIND

1%

A084150 001

1%

A084863 001

PENTOLAIR

PHARMAFAIR

0.5%

A088643 001 Feb 09, 1987

1%

A088150 001 Feb 25, 1983

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

BAXTER HLTHCARE

100MG/VIAL

A088371 001 Jul 03, 1986

200MG/VIAL

A088372 001 Jul 03, 1986

500MG/VIAL

A088373 001 Jul 03, 1986

1GM/VIAL

A088374 001 Sep 24, 1986

CYTOXAN

+ BAXTER HLTHCARE

100MG/VIAL \*\*

N012142 001

+

200MG/VIAL \*\*

N012142 002

CYTOXAN (LYOPHILIZED)

+ BAXTER HLTHCARE

500MG/VIAL

N012142 003

+

500MG/VIAL \*\*

N012142 008 Jan 04, 1984

+

1GM/VIAL

N012142 004 Aug 30, 1982

+

1GM/VIAL \*\*

N012142 010 Sep 24, 1985

+

2GM/VIAL

N012142 005 Aug 30, 1982

+

2GM/VIAL \*\*

N012142 009 Dec 10, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-105(of 375)

\*\* See List Footnote

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

LYOPHILIZED CYTOXAN

+ BAXTER HLTHCARE	100MG/VIAL **
+	200MG/VIAL **

NEOSAR

BEDFORD	100MG/VIAL
	200MG/VIAL
	500MG/VIAL
	1GM/VIAL
	2GM/VIAL
TEVA PARENTERAL	100MG/VIAL
	200MG/VIAL
	500MG/VIAL
	1GM/VIAL
	2GM/VIAL

N012142 006	Dec 05, 1985
N012142 007	Dec 10, 1985
A087442 001	Feb 16, 1982
A087442 002	Feb 16, 1982
A087442 003	Feb 16, 1982
A087442 004	Jul 08, 1983
A087442 005	Mar 30, 1989
A040015 001	Apr 29, 1993
A040015 002	Apr 29, 1993
A040015 003	Apr 29, 1993
A040015 004	Apr 29, 1993
A040015 005	Apr 29, 1993

TABLET; ORAL

CYCLOPHOSPHAMIDE

ROXANE	25MG
	50MG

A040032 001	Aug 17, 1999
A040032 002	Aug 17, 1999

CYTOXAN

+ BAXTER HLTHCARE	25MG **
+	50MG **

N012141 002	
N012141 001	

CYCLOSPORINE

CAPSULE; ORAL

NEORAL

+ NOVARTIS	50MG **
------------	---------

N050715 003	Jul 14, 1995
-------------	--------------

SOLUTION; ORAL

CYCLOSPORINE

APOTEX INC	100MG/ML
------------	----------

A065167 001	Jan 05, 2005
-------------	--------------

CYCLOTHIAZIDE

TABLET; ORAL

ANHYDRON

LILLY	2MG
FLUIDIL	
PHARMACIA AND UPJOHN	2MG

N013157 002	
N018173 001	

CYCRIMINE HYDROCHLORIDE

TABLET; ORAL

PAGITANE

LILLY	1.25MG
	2.5MG

N008951 001	
N008951 002	

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	2MG/5ML **
HALSEY	2MG/5ML
MORTON GROVE	2MG/5ML
NASKA	2MG/5ML

A086833 001	
A089199 001	Jul 03, 1986
A087001 001	Nov 04, 1982
A089021 001	Dec 21, 1987

PERIACTIN

+ MERCK	2MG/5ML **
---------	------------

N013220 002	
-------------	--

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

AM THERAP	4MG
ASCOT	4MG
CHARTWELL RX	4MG
DURAMED PHARMS BARR	4MG
HALSEY	4MG
KV PHARM	4MG
MD PHARM	4MG
MYLAN	4MG
PIONEER PHARMS	4MG
PLIVA	4MG
SANDOZ	4MG
SUPERPHARM	4MG
VITARINE	4MG
WATSON LABS	4MG
	4MG

A088798 001	Feb 15, 1985
A087685 001	Oct 25, 1982
A088212 001	May 26, 1983
A088232 001	Oct 25, 1983
A089057 001	Jul 03, 1986
A086737 001	
A087566 001	Nov 10, 1982
A086678 001	
A087839 001	Feb 08, 1984
A088205 001	Jul 26, 1983
A086808 001	
A087405 001	
A087284 001	
A085245 001	
A086165 001	

**DISCONTINUED DRUG PRODUCT LIST**

6-106(of 375)

\*\* See List Footnote

CYPROHEPTADINE HYDROCHLORIDE

TABLET;ORAL

CYPROHEPTADINE HYDROCHLORIDE

4MG

A086580 001

PERIACTIN

+ MERCK

4MG \*\*

N012649 001

CYSTEINE HYDROCHLORIDE

INJECTABLE;INJECTION

CYSTEINE HYDROCHLORIDE

+ HOSPIRA

7.25% \*\*

N019523 001 Oct 22, 1986

CYTARABINE

INJECTABLE;INJECTION

CYTARABINE

+ TEVA PARENTERAL

100MG/VIAL \*\*

N016793 001

+

500MG/VIAL \*\*

N016793 002

+

1GM/VIAL \*\*

N016793 003 Dec 21, 1987

+

2GM/VIAL \*\*

N016793 004 Dec 21, 1987

CYTOSAR-U

TEVA PHARMS USA

100MG/VIAL

A075206 001 Dec 30, 1998

500MG/VIAL

A075206 002 Dec 30, 1998

1GM/VIAL

A075206 004 Dec 30, 1998

2GM/VIAL

A075206 003 Dec 30, 1998

DACARBAZINE

INJECTABLE;INJECTION

DACARBAZINE

ABRAXIS PHARM

100MG/VIAL

A070962 001 Aug 28, 1986

200MG/VIAL

A070990 001 Aug 28, 1986

DTIC-DOME

+ BAYER HLTHCARE

100MG/VIAL

N017575 001

+

200MG/VIAL

N017575 002

DACTINOMYCIN

INJECTABLE;INJECTION

DACTINOMYCIN

WEST-WARD PHARMS INT 0.5MG/VIAL

A090304 001 Mar 16, 2010

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE;IV (INFUSION)

SYNERCID

KING PHARMS

420MG/VIAL;180MG/VIAL

N050748 002 Aug 24, 2000

DALTEPARIN SODIUM

INJECTABLE;INJECTION

FRAGMIN

PFIZER INC

7,500 IU/0.75ML

N020287 008 Apr 04, 2002

INJECTABLE;SUBCUTANEOUS

FRAGMIN

PFIZER INC

10,000IU/0.4ML (25,000IU/ML)

N020287 002 May 01, 2007

95,000IU/9.5ML (10,000IU/ML)

N020287 007 Apr 04, 2002

DANAPAROID SODIUM

INJECTABLE;INJECTION

ORGARAN

ASPEN GLOBAL INC

750 UNITS/0.6ML

N020430 001 Dec 24, 1996

DANAZOL

CAPSULE;ORAL

DANAZOL

AM THERAP

200MG

A071569 001 Dec 30, 1987

DANOCRINE

SANOFI AVENTIS US

50MG

N017557 003

100MG

N017557 004

200MG

N017557 002

DAPIPRAZOLE HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC

DAPIPRAZOLE HYDROCHLORIDE

+ FERA PHARMS

0.5% \*\*

N019849 001 Dec 31, 1990

**DISCONTINUED DRUG PRODUCT LIST**

6-107(of 375)

\*\* See List Footnote

DAPTOMYCIN

POWDER;IV (INFUSION)

CUBICIN

CUBIST PHARMS LLC

250MG/VIAL

N021572 001 Sep 12, 2003

DARUNAVIR ETHANOLATE

TABLET;ORAL

PREZISTA

+ JANSSEN PRODS

EQ 300MG BASE \*\*

N021976 001 Jun 23, 2006

+

EQ 400MG BASE \*\*

N021976 003 Oct 21, 2008

DAUNORUBICIN CITRATE

INJECTABLE, LIPOSOMAL; INJECTION

DAUNOXOME

GALEN (UK)

EQ 2MG BASE/ML

N050704 002 Apr 08, 1996

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

SANOFI AVENTIS US

EQ 20MG BASE/VIAL

A061876 001

WYETH AYERST

EQ 20MG BASE/VIAL

N050484 001

DAUNORUBICIN HYDROCHLORIDE

TEVA PARENTERAL

EQ 20MG BASE/VIAL

A064212 001 Jun 23, 1998

EQ 50MG BASE/VIAL

A064212 002 May 03, 1999

DECAMETHONIUM BROMIDE

INJECTABLE; INJECTION

SYNCURINE

GLAXOSMITHKLINE

1MG/ML

N006931 002

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

WATSON LABS

500MG/VIAL

A076806 001 Mar 31, 2006

2GM/VIAL

A076806 002 Mar 31, 2006

DEMECARIUM BROMIDE

SOLUTION/DROPS; OPHTHALMIC

HUMORSOL

MERCK

0.125%

N011860 002

0.25%

N011860 001

DEMECLOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DECLOMYCIN

LEDERLE

150MG

N050262 001

SYRUP; ORAL

DECLOMYCIN

LEDERLE

75MG/5ML

N050257 001

TABLET; ORAL

DECLOMYCIN

COREPHARMA

75MG

N050261 001

150MG

N050261 002

300MG

N050261 003

DEMECLOCYCLINE HYDROCHLORIDE

IMPAK LABS

150MG

A065094 001 Mar 22, 2004

300MG

A065094 002 Mar 22, 2004

DESERPIDINE

TABLET; ORAL

HARMONYL

ABBVIE

0.1MG

N010796 001

0.25MG

N010796 002

DESERPIDINE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ORETICYL 25

ABBVIE

0.125MG; 25MG

N012148 001

ORETICYL 50

ABBVIE

0.125MG; 50MG

N012148 003

ORETICYL FORTE

ABBVIE

0.25MG; 25MG

N012148 002

**DISCONTINUED DRUG PRODUCT LIST**

6-108(of 375)

\*\* See List Footnote

DESERPIDINE; METHYCLOTHIAZIDE

TABLET;ORAL

ENDURONYL

ABBOTT

0.25MG;5MG

N012775 001

ENDURONYL FORTE

ABBOTT

0.5MG;5MG

N012775 002

METHYCLOTHIAZIDE AND DESERPIDINE

WATSON LABS

0.25MG;5MG

A088486 001 Aug 10, 1984

0.5MG;5MG

A088452 001 Aug 10, 1984

DESIPRAMINE HYDROCHLORIDE

CAPSULE;ORAL

PERTOFRANE

SANOFI AVENTIS US

25MG

N013621 001

50MG

N013621 002

TABLET;ORAL

DESIPRAMINE HYDROCHLORIDE

ANI PHARMS INC

100MG

A071803 001 May 29, 1997

USL PHARMA

25MG

A071864 001 Sep 09, 1987

50MG

A071865 001 Sep 09, 1987

75MG

A071866 001 Sep 09, 1987

100MG

A071867 001 Sep 09, 1987

DESLANOSIDE

INJECTABLE;INJECTION

CEDILANID-D

NOVARTIS

0.2MG/ML

N009282 002

DESMOPRESSIN ACETATE

INJECTABLE;INJECTION

DDAVP

FERRING PHARMS INC

0.015MG/ML

N018938 002 Apr 25, 1995

DESMOPRESSIN ACETATE

BEDFORD

0.004MG/ML

A074575 001 Feb 18, 2000

HOSPIRA

0.004MG/ML

A075220 001 Aug 28, 2000

TEVA PHARMS USA

0.004MG/ML

A074888 001 Oct 15, 1997

DESMOPRESSIN ACETATE PRESERVATIVE FREE

BEDFORD

0.004MG/ML

A074574 001 Feb 18, 2000

SOLUTION;NASAL

CONCENTRAID

FERRING

0.01%

N019776 001 Dec 26, 1990

SPRAY, METERED;NASAL

DDAVP

+ FERRING PHARMS INC

0.01MG/SPRAY \*\*

N017922 002 Feb 06, 1989

STIMATE

FERRING PHARMS INC

0.15MG/SPRAY

N020355 001 Mar 07, 1994

TABLET;ORAL

DESMOPRESSIN ACETATE

FERRING

0.1MG

N021795 001 May 08, 2008

0.2MG

N021795 002 May 08, 2008

DESOGESTREL; ETHINYLMESTRADIOL

TABLET;ORAL-21

DESOGEN

ORGANON USA INC

0.15MG;0.03MG

N020071 001 Dec 10, 1992

DESOGESTREL AND ETHINYLMESTRADIOL

DURAMED PHARMS BARR

0.15MG;0.03MG

A075256 001 Aug 12, 1999

ORTHO-CEPT

JANSSEN PHARMS

0.15MG;0.03MG

N020301 001 Dec 14, 1992

TABLET;ORAL-28

MIRCETTE

+ TEVA BRANDED PHARM

0.15MG,N/A;0.02MG,0.01MG \*\*

N020713 001 Apr 22, 1998

ORTHO-CEPT

JANSSEN PHARMS

0.15MG;0.03MG

N020301 002 Dec 14, 1992

DESOXIMETASONE

CREAM;TOPICAL

TOPICORT

+ TARO

0.25% \*\*

N017856 001

TOPICORT LP

+ TARO

0.05% \*\*

N018309 001

**DISCONTINUED DRUG PRODUCT LIST**

6-109(of 375)

\*\* See List Footnote

DESOXIMETASONE

GEL;TOPICAL			
TOPICORT			
+ TARO	0.05% **		N018586 001 Mar 29, 1982
OINTMENT;TOPICAL			
DESOXIMETASONE			
ALTANA	0.25%		A073440 001 Apr 01, 1998
TOPICORT			
+ TARO	0.25% **		N018763 001 Sep 30, 1983

DESOXYCORTICOSTERONE ACETATE

INJECTABLE;INJECTION			
DOCA			
ORGANON USA INC	5MG/ML		N001104 001
PELLET;IMPLANTATION			
PERCORTEN			
NOVARTIS	125MG		N005151 001

DESOXYCORTICOSTERONE PIVALATE

INJECTABLE;INJECTION			
PERCORTEN			
NOVARTIS	25MG/ML		N008822 001

DESVENLAFAKINE FUMARATE

TABLET, EXTENDED RELEASE;ORAL			
DESVENLAFAKINE			
TEVA PHARMS USA	EQ 50MG BASE		N205208 001 Oct 11, 2013
	EQ 100MG BASE		N205208 002 Oct 11, 2013

DEXAMETHASONE

AEROSOL;TOPICAL			
AEROSEB-DEX			
ALLERGAN HERBERT	0.01% **		A083296 002
DECASPRAY			
+ MERCK	0.04% **		N012731 002
ELIXIR;ORAL			
DECADRON			
MERCK	0.5MG/5ML		N012376 002
DEXAMETHASONE			
ALPHARMA US PHARMS	0.5MG/5ML		A088997 001 Oct 10, 1986
HEXDROL			
ORGANON USA INC	0.5MG/5ML		N012674 001
GEL;TOPICAL			
DECADERM			
MERCK	0.1%		N013538 001
SUSPENSION/DROPS;OPHTHALMIC			
DEXAMETHASONE			
WATSON LABS	0.1%		A089170 001 May 09, 1989
TABLET;ORAL			
DECADRON			
+ MERCK	0.25MG **		N011664 004
+	0.5MG **		N011664 001
+	0.75MG **		N011664 002
+	1.5MG **		N011664 003
+	4MG **		N011664 005
+	6MG **		N011664 006 Jul 30, 1982
DEXAMETHASONE			
IDT AUSTRALIA LTD	0.75MG		A080399 001
IMPAX LABS	0.75MG		A085376 001
PAR PHARM	0.25MG		A088149 001 Apr 28, 1983
PHOENIX LABS NY	0.75MG		A083806 001
PVT FORM	0.75MG		A083420 001
ROXANE	0.25MG		A084614 001
SUN PHARM INDUSTRIES	0.25MG		A084013 001
	0.25MG		A084764 001
	0.5MG		A084084 001
	0.75MG		A084766 001
	0.75MG		A084081 001
	1.5MG		A084765 001
	1.5MG		A084086 001
	1.5MG		A084763 001

**DISCONTINUED DRUG PRODUCT LIST**

6-110(of 375)

\*\* See List Footnote

DEXAMETHASONE

TABLET;ORAL

DEXAMETHASONE

UPSHER SMITH	0.75MG 1.5MG	A087534 001 A087533 001
WATSON LABS	0.25MG 0.5MG 0.75MG 0.75MG 0.75MG 1.5MG 1.5MG	A085455 001 A085458 001 A080968 001 A084457 001 A085818 001 A085456 001 A085840 001
WHITEWORTH TOWN PLSN	0.75MG	A084327 001
DEXONE 0.5		
SOLVAY	0.5MG	A084991 001
DEXONE 0.75		
SOLVAY	0.75MG	A084993 001
DEXONE 1.5		
SOLVAY	1.5MG	A084990 001
DEXONE 4		
SOLVAY	4MG	A084992 001
HEXDROL		
ASPEN GLOBAL INC	0.5MG 0.75MG 1.5MG 4MG	N012675 004 N012675 007 N012675 009 N012675 010

DEXAMETHASONE ACETATE

INJECTABLE;INJECTION

DECADRON-LA

+ MERCK	EQ 8MG BASE/ML **	N016675 001
---------	-------------------	-------------

DEXAMETHASONE ACETATE

WATSON LABS	EQ 8MG BASE/ML	A084315 001
WATSON LABS TEVA	EQ 16MG BASE/ML	A087711 001 May 24, 1982

DEXAMETHASONE SODIUM PHOSPHATE

AEROSOL;NASAL

DEXACORT

UCB INC	EQ 0.1MG PHOSPHATE/INH	N014242 001
---------	------------------------	-------------

AEROSOL, METERED;INHALATION

DEXACORT

UCB INC	EQ 0.1MG PHOSPHATE/INH	N013413 001
---------	------------------------	-------------

CREAM;TOPICAL

DECADRON

MERCK	EQ 0.1% PHOSPHATE	N011983 002
-------	-------------------	-------------

INJECTABLE;INJECTION

DECADRON

+ MERCK	EQ 4MG PHOSPHATE/ML **	N012071 002
---------	------------------------	-------------

+	EQ 24MG PHOSPHATE/ML **	N012071 004
---	-------------------------	-------------

DEXACEN-4

CENT PHARMS	EQ 4MG PHOSPHATE/ML	A084342 001
-------------	---------------------	-------------

DEXAMETHASONE

ABRAXIS PHARM	EQ 4MG PHOSPHATE/ML	A088448 001 Jan 25, 1984
---------------	---------------------	--------------------------

FRESENIUS KABI USA	EQ 10MG PHOSPHATE/ML	A088469 001 Jan 25, 1984
--------------------	----------------------	--------------------------

DEXAMETHASONE SODIUM PHOSPHATE

AKORN	EQ 4MG PHOSPHATE/ML	A084493 001
-------	---------------------	-------------

BEL MAR	EQ 4MG PHOSPHATE/ML	A084752 001
---------	---------------------	-------------

DELL LABS	EQ 4MG PHOSPHATE/ML	A083161 001
-----------	---------------------	-------------

INTL MEDICATION	EQ 20MG PHOSPHATE/ML	A088522 001 Feb 17, 1984
-----------------	----------------------	--------------------------

LYPHOMED	EQ 4MG PHOSPHATE/ML	A087065 001
----------	---------------------	-------------

TEVA PARENTERAL	EQ 4MG PHOSPHATE/ML	A081125 001 Aug 31, 1990
-----------------	---------------------	--------------------------

EQ 10MG PHOSPHATE/ML	A081126 001 Aug 31, 1990	
----------------------	--------------------------	--

EQ 24MG PHOSPHATE/ML	A083702 001	
----------------------	-------------	--

EQ 4MG PHOSPHATE/ML	A084355 001	
---------------------	-------------	--

EQ 4MG PHOSPHATE/ML	A089169 001 Apr 09, 1986	
---------------------	--------------------------	--

EQ 10MG PHOSPHATE/ML	A087668 001 Jul 01, 1982	
----------------------	--------------------------	--

EQ 24MG PHOSPHATE/ML	A085606 001	
----------------------	-------------	--

EQ 4MG PHOSPHATE/ML	A085641 001	
---------------------	-------------	--

HEXDROL

+ ORGANON USA INC	EQ 4MG PHOSPHATE/ML **	N014694 002
-------------------	------------------------	-------------

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-111(of 375)

\*\* See List Footnote

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HEXDROL			
+	EQ 10MG PHOSPHATE/ML **	N014694 003	
	EQ 20MG PHOSPHATE/ML	N014694 004	
OINTMENT; OPHTHALMIC			
DECADRON			
MERCK	EQ 0.05% PHOSPHATE	N011977 001	
DEXAIR			
PHARMAFAIR	EQ 0.05% PHOSPHATE	A088071 001	Dec 28, 1982
MAXIDEX			
ALCON	EQ 0.05% PHOSPHATE	A083342 001	
SOLUTION/DROPS; OPHTHALMIC			
DEXAIR			
PHARMAFAIR	EQ 0.1% PHOSPHATE	A088433 001	Dec 15, 1983
DEXAMETHASONE SODIUM PHOSPHATE			
SOLA BARNES HIND	EQ 0.1% PHOSPHATE	A084170 001	
	EQ 0.1% PHOSPHATE	A084173 001	
SOLUTION/DROPS; OPHTHALMIC, OTIC			
DECADRON			
MERCK	EQ 0.1% PHOSPHATE	N011984 001	
SOLUTION/DROPS; OTIC			
DEXAMETHASONE SODIUM PHOSPHATE			
AKORN	EQ 0.1% PHOSPHATE	A084855 001	

DEXAMETHASONE SODIUM PHOSPHATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DECADRON W/ XYLOCAINE		
MERCK	EQ 4MG PHOSPHATE/ML; 10MG/ML	N013334 002

DEXAMETHASONE SODIUM PHOSPHATE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC		
NEODECADRON		
MERCK	EQ 0.05% PHOSPHATE; EQ 3.5MG BASE/GM	N050324 001
SOLUTION/DROPS; OPHTHALMIC		
NEODECADRON		
MERCK	EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML	N050322 001
NEOMYCIN SULFATE AND DEXAMETHASONE SODIUM PHOSPHATE		
BAUSCH AND LOMB	EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML	A064055 001
NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE		Oct 30, 1995
ALCON PHARMS LTD	EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML	A062714 001
PHARMAFAIR	EQ 0.1% PHOSPHATE; EQ 3.5MG BASE/ML	A062539 001
		Jul 21, 1986

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC		
DEXACIDIN		
NOVARTIS	0.1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	A062566 001
DEXASPORIN		Feb 22, 1985
PHARMAFAIR	0.1%; EQ 3.5MG BASE/GM; 10,000 UNITS/GM	A062411 001
SUSPENSION/DROPS; OPHTHALMIC		May 16, 1983
DEXACIDIN		
NOVARTIS	0.1%; EQ 3.5MG BASE/ML; 10,000 UNITS/ML	A062544 001
DEXASPORIN		Oct 29, 1984
PHARMAFAIR	0.1%; EQ 3.5MG BASE/ML; 10,000 UNITS/ML	A062428 001
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE		May 18, 1983
ALCON PHARMS LTD	0.1%; EQ 3.5MG BASE/ML; 10,000 UNITS/ML	A062721 001
		Nov 17, 1986

DEXBROMPHENIRAMINE MALEATE

SYRUP; ORAL		
DISOMER		
SCHERING	2MG/5ML	N011814 002
TABLET; ORAL		
DISOMER		
SCHERING	2MG	N011814 001

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET; ORAL		
DISOPHROL		
SCHERING	2MG; 60MG	N012394 002

**DISCONTINUED DRUG PRODUCT LIST**

6-112(of 375)

\*\* See List Footnote

**DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE**

TABLET, EXTENDED RELEASE;ORAL

BROMPHERIL

COPLEY PHARM 6MG;120MG

A089116 001 Jan 22, 1987

DISOBROM

SANDOZ 6MG;120MG

A070770 001 Sep 30, 1991

DISOPHROL

SCHERING PLOUGH 6MG;120MG

N013483 004 Sep 13, 1982

DRIXORAL

+ SCHERING PLOUGH 6MG;120MG \*\*

N013483 003 Sep 13, 1982

RESPORAL

PIONEER PHARMS 6MG;120MG

A089139 001 Jun 16, 1988

**DEXCHLORPHENIRAMINE MALEATE**

SYRUP;ORAL

POLARAMINE

SCHERING 2MG/5ML

A086837 001 Jul 19, 1982

TABLET;ORAL

DEXCHLORPHENIRAMINE MALEATE

ANI PHARMS INC 2MG

A088682 001 Jan 17, 1986

POLARAMINE

SCHERING 2MG

A086835 001

**DEXLANSOPRAZOLE**

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

DEXILANT SOLUTAB

+ TAKEDA PHARMS USA 30MG

N208056 001 Jan 26, 2016

**DEXTROAMPHETAMINE SULFATE**

CAPSULE;ORAL

DEXAMPEX

TEVA 15MG

A085355 001

CAPSULE, EXTENDED RELEASE;ORAL

DEXTROAMPHETAMINE SULFATE

ABLE 5MG

A076814 001 Aug 25, 2004

10MG

A076814 002 Aug 25, 2004

15MG

A076814 003 Aug 25, 2004

ELIXIR;ORAL

DEXEDRINE

GLAXOSMITHKLINE 5MG/5ML \*\*

A083902 001

TABLET;ORAL

DEXAMPEX

TEVA 5MG

A083735 001

10MG

A083735 002

DEXEDRINE

GLAXOSMITHKLINE 5MG

A084935 001

DEXTROAMPHETAMINE SULFATE

EPIC PHARMA LLC 5MG

A090652 001 Mar 07, 2014

10MG

A090652 002 Mar 07, 2014

HALSEY 10MG

A083930 001

IDT AUSTRALIA LTD 5MG

A085370 001

LANNETT 5MG

A083903 001

10MG

A083903 003

15MG

A085652 001

MAST MM 5MG

A086521 001

NESHER PHARMS 5MG

A040365 001 Oct 31, 2002

10MG

A040367 001 Oct 31, 2002

PUREPAC PHARM 5MG

A084125 001

SANDOZ 10MG

A085371 001

VINTAGE PHARMS LLC 5MG

A040299 001 May 13, 1999

VITARINE 5MG

A084986 001

10MG

A085892 001

DEXTROSTAT

SHIRE 5MG \*\*

A084051 001

10MG \*\*

A084051 002

FERNDEX

FERNDALE LABS 5MG

A084001 001

**DISCONTINUED DRUG PRODUCT LIST**

6-113(of 375)

\*\* See List Footnote

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL			
PHERAZINE DM			
HALSEY	15MG/5ML; 6.25MG/5ML	A088913	001 Mar 02, 1987
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE			
AMNEAL PHARMS	15MG/5ML; 6.25MG/5ML	A090575	001 Feb 08, 2011
+ ANI PHARMS	15MG/5ML; 6.25MG/5ML **	N011265	002 Apr 02, 1984
TRIS PHARMA INC	15MG/5ML; 6.25MG/5ML	A091687	001 Jun 28, 2012

DEXTROSE

INJECTABLE; INJECTION			
DEXTROSE 10% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML	N018046	001
MILES	10GM/100ML	N018504	001
DEXTROSE 2.5% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML	N018358	001
	2.5GM/100ML	N019626	001 Feb 02, 1988
DEXTROSE 38.5% IN PLASTIC CONTAINER			
ABBOTT	38.5GM/100ML	N018923	001 Sep 19, 1984
DEXTROSE 5% IN PLASTIC CONTAINER			
DHL	5GM/100ML	N019971	001 Sep 28, 1995
DEXTROSE 50% IN PLASTIC CONTAINER			
ICU MEDICAL INC	50GM/100ML	N019894	001 Dec 26, 1989
DEXTROSE 60%			
B BRAUN	60GM/100ML	N017995	002 Sep 22, 1982
DEXTROSE 60% IN PLASTIC CONTAINER			
B BRAUN	60GM/100ML	N017995	001
BAXTER HLTHCARE	60GM/100ML	N020047	002 Jul 02, 1991
HOSPIRA	60GM/100ML	N019346	001 Jan 25, 1985
DEXTROSE 7.7% IN PLASTIC CONTAINER			
B BRAUN	7.7GM/100ML	N019626	003 Feb 02, 1988

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION			
ISOLYTE P W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 31MG/100ML; 130MG/100ML; 26MG/100ML; 320MG/100ML	N019025	001 Dec 27, 1984

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

INJECTABLE; INJECTION			
IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA	5GM/100ML; 53MG/100ML; 100MG/100ML; 100MG/100ML; 180MG/100ML; 280MG/100ML; 16MG/100ML	N019515	001 May 08, 1986

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION			
ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/100ML; 140MG/100ML	N019844	001 Jun 10, 1993
ISOLYTE H W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 30MG/100ML; 97MG/100ML; 220MG/100ML; 140MG/100ML	N018273	001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION			
ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML	N019843	001 Aug 09, 1993
ISOLYTE S W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML; 30MG/100ML; 37MG/100ML; 370MG/100ML; 530MG/100ML; 500MG/100ML	N018274	001
PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML; 30MG/100ML; 37MG/100ML; 368MG/100ML; 526MG/100ML; 502MG/100ML	N017451	001

**DISCONTINUED DRUG PRODUCT LIST**

6-114(of 375)

\*\* See List Footnote

DEXTROSE; POTASSIUM CHLORIDE

## INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;37MG/100ML	N019699 001 Sep 29, 1989
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;75MG/100ML 5GM/100ML;75MG/100ML	N018744 001 Nov 09, 1982 N019699 002 Sep 29, 1989
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;110MG/100ML	N019699 003 Sep 29, 1989
POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;150MG/100ML	N018744 002 Nov 09, 1982
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;220MG/100ML 5GM/100ML;220MG/100ML	N018744 003 Nov 09, 1982 N019699 005 Sep 29, 1989
POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;300MG/100ML	N018744 004 Nov 09, 1982

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM LACTATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, MONOBASIC ANHYDROUS

## INJECTABLE; INJECTION

IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER HOSPIRA	5GM/100ML;111MG/100ML;256MG/100ML;146MG /100ML;207MG/100ML	N019514 001 May 08, 1986
---	---	--------------------------

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

## INJECTABLE; INJECTION

ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG /100ML;91MG/100ML	N019870 001 Jun 10, 1993
ISOLYTE M W/ DEXTROSE 5% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG /100ML;91MG/100ML	N018270 001

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

## INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO. 75 IN PLASTIC CONTAINER BAXTER HLTHCARE	5GM/100ML;205MG/100ML;100MG/100ML;120MG /100ML;220MG/100ML	N018840 001 Jun 29, 1983
--	---	--------------------------

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

## INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.075% B BRAUN	5GM/100ML;75MG/100ML;200MG/100ML	N018268 009
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;150MG/100ML;200MG/100ML	N018268 004
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;220MG/100ML;200MG/100ML	N018268 005
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;300MG/100ML;200MG/100ML	N018268 006
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;75MG/100ML;330MG/100ML	N018268 011 Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;150MG/100ML;330MG/100ML	N018268 012 Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;220MG/100ML;330MG/100ML	N018268 013 Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.30% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;300MG/100ML;330MG/100ML	N018268 014 Jan 18, 1986
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.075% B BRAUN	5GM/100ML;75MG/100ML;450MG/100ML	N018268 010
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;150MG/100ML;450MG/100ML	N018268 001
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;220MG/100ML;450MG/100ML	N018268 002
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;300MG/100ML;450MG/100ML	N018268 003
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 15MEO IN PLASTIC CONTAINER BAXTER HLTHCARE	5GM/100ML;224MG/100ML;450MG/100ML	N018008 003
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEO (K) IN PLASTIC CONTAINER BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N018008 001
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 5MEO IN PLASTIC CONTAINER BAXTER HLTHCARE	5GM/100ML;75MG/100ML;450MG/100ML	N018008 002

**DISCONTINUED DRUG PRODUCT LIST**

6-115(of 375)

\*\* See List Footnote

DEXTROSE; SODIUM CHLORIDE

## INJECTABLE; INJECTION

DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER B BRAUN	10GM/100ML;200MG/100ML	N018386 001
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER B BRAUN	10GM/100ML;450MG/100ML	N018229 001
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER B BRAUN	10GM/100ML;900MG/100ML	N018047 001
BAXTER HLTHCARE	10GM/100ML;900MG/100ML	N016696 001
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER B BRAUN	2.5GM/100ML;450MG/100ML	N018030 001
HOSPIRA	2.5GM/100ML;450MG/100ML	N018096 001
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER B BRAUN	2.5GM/100ML;900MG/100ML	N018376 001
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER ABBOTT	3.3GM/100ML;300MG/100ML	N018055 001
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;110MG/100ML	N018030 005
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;200MG/100ML	N018030 004
MILES	5GM/100ML;200MG/100ML	N018399 001
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER ABBOTT	5GM/100ML;225MG/100ML	N019482 001 Oct 04, 1985
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER ABBOTT	5GM/100ML;300MG/100ML	N019486 001 Oct 04, 1985
MILES	5GM/100ML;300MG/100ML	N018501 001
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER B BRAUN	5GM/100ML;330MG/100ML	N018030 003
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER ABBOTT	5GM/100ML;450MG/100ML	N019484 001 Oct 04, 1985
B BRAUN	5GM/100ML;450MG/100ML	N018030 002
MILES	5GM/100ML;450MG/100ML	N018400 001
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER ABBOTT	5GM/100ML;900MG/100ML	N019483 001 Oct 04, 1985
B BRAUN	5GM/100ML;900MG/100ML	N018026 001
MILES	5GM/100ML;900MG/100ML	N018500 001

DEXTROTHYROXINE SODIUM

## TABLET; ORAL

## CHOLOXIN

ABBVIE	1MG	N012302 005
	2MG	N012302 002
	4MG	N012302 004
	6MG	N012302 006

DEZOCINE

## INJECTABLE; INJECTION

## DALGAN

ASTRAZENECA	5MG/ML	N019082 001 Dec 29, 1989
	10MG/ML	N019082 002 Dec 29, 1989
	15MG/ML	N019082 003 Dec 29, 1989

DIATRIZOATE MEGLUMINE

## INJECTABLE; INJECTION

## ANGIOVIST 282

BAYER HLTHCARE	60%	A087726 001 Sep 23, 1982
----------------	-----	--------------------------

## CARDIOGRAFIN

BRACCO	85%	N011620 002
--------	-----	-------------

## DIATRIZOATE MEGLUMINE

BRACCO	76%	N010040 017
--------	-----	-------------

## HYPAQUE

GE HEALTHCARE	30%	N016403 002
	60%	N016403 001

## RENO-60

BRACCO	60%	N010040 016
--------	-----	-------------

## RENO-DIP

BRACCO	30%	N010040 012
--------	-----	-------------

## UROVIST MEGLUMINE DIU/CT

BAYER HLTHCARE	30%	A087739 001 Sep 23, 1982
----------------	-----	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-116(of 375)

\*\* See List Footnote

DIATRIZOATE MEGLUMINE

SOLUTION;URETERAL

RENO-30

BRACCO 30%

N010040 021

UROVIST CYSTO

BAYER HLTHCARE 30%

A087729 001 Sep 23, 1982

UROVIST CYSTO PEDIATRIC

BAYER HLTHCARE 30%

A087731 001 Sep 23, 1982

SOLUTION;URETHRAL

HYPAQUE-CYSTO

GE HEALTHCARE 30%

N016403 003

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

ANGIOVIST 292

BAYER HLTHCARE 52%;8%

A087724 001 Sep 23, 1982

ANGIOVIST 370

BAYER HLTHCARE 66%;10%

A087723 001 Sep 23, 1982

DIATRIZOATE-60

INTL MEDICATION 52%;8%

A088166 001 Jun 17, 1983

HYPAQUE-76

GE HEALTHCARE 66%;10%

A086505 001

HYPAQUE-M, 75%

GE HEALTHCARE 50%;25%

N010220 003

HYPAQUE-M, 90%

GE HEALTHCARE 60%;30%

N010220 002

MD-60

MALLINCKRODT 52%;8%

A087074 001

MD-76

MALLINCKRODT 66%;10%

A087073 001

MD-76R

+ LIEBEL-FLARSHEIM 66%;10%

N019292 001 Sep 29, 1989

RENOCAL-76

BRACCO 66%;10%

A089347 001 Jun 01, 1988

RENOGRAFIN-60

BRACCO 52%;8%

N010040 006

RENOVIST

BRACCO 34.3%;35%

N010040 020

RENOVIST II

BRACCO 28.5%;29.1%

N010040 019

SOLUTION;ORAL, RECTAL

GASTROVIST

BAYER HLTHCARE 66%;10%

A087728 001 Sep 23, 1982

DIATRIZOATE SODIUM

FOR SOLUTION;ORAL, RECTAL

HYPAQUE

GE HEALTHCARE 100%

N011386 001

INJECTABLE; INJECTION

HYPAQUE

GE HEALTHCARE 25%

N009561 003

50%

N009561 001

MD-50

MALLINCKRODT 50%

A087075 001

UROVIST SODIUM 300

BAYER HLTHCARE 50%

A087725 001 Sep 23, 1982

SOLUTION;ORAL, RECTAL

HYPAQUE

GE HEALTHCARE 40%

N011386 003

SOLUTION;URETERAL

HYPAQUE SODIUM 20%

GE HEALTHCARE 20%

N009561 002

DIAZEPAM

CAPSULE, EXTENDED RELEASE;ORAL

VALRELEASE

ROCHE 15MG

N018179 001

GEL;RECTAL

DIASTAT

+ VALEANT PHARMS NORTH 5MG/ML (5MG/ML) \*\*

N020648 002 Jul 29, 1997

+ 10MG/2ML (5MG/ML) \*\*

N020648 003 Jul 29, 1997

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-117(of 375)

\*\* See List Footnote

**DIAZEPAM**

GEL;RECTAL

DIASTAT

+	15MG/3ML (5MG/ML) **
+	20MG/4ML (5MG/ML) **

N020648 004	Jul 29, 1997
N020648 005	Jul 29, 1997

INJECTABLE;INJECTION

DIAZEPAM

ABRAXIS PHARM	5MG/ML	A070662 001	Jun 25, 1986
HOSPIRA	5MG/ML	A071584 001	Oct 13, 1987
MARSAM PHARMS LLC	5MG/ML	A072371 001	Jan 29, 1993
PARENTA PHARMS	5MG/ML	A076815 001	Apr 15, 2004
US ARMY	5MG/ML **	N020124 001	Dec 05, 1990
WARNER CHILCOTT	5MG/ML	A071613 001	Oct 22, 1987
	5MG/ML	A071614 001	Oct 22, 1987
WATSON LABS	5MG/ML	A070296 001	Feb 12, 1986
	5MG/ML	A070911 001	Aug 28, 1986
	5MG/ML	A070912 001	Aug 28, 1986
	5MG/ML	A070930 001	Dec 01, 1986
WATSON LABS INC	5MG/ML	A072370 001	Jan 29, 1993
	5MG/ML	A072397 001	Jan 29, 1993
WEST-WARD PHARMS INT	5MG/ML	A070311 001	Dec 16, 1985
	5MG/ML	A070312 001	Dec 16, 1985
	5MG/ML	A070313 001	Dec 16, 1985
	5MG/ML	A071308 001	Jul 17, 1987
	5MG/ML	A071309 001	Jul 17, 1987
	5MG/ML	A071310 001	Jul 17, 1987

DIZAC

PHARMACIA AND UPJOHN	5MG/ML **	N019287 001	Jun 18, 1993
----------------------	-----------	-------------	--------------

VALIUM

+ ROCHE	5MG/ML **	N016087 001
---------	-----------	-------------

TABLET;ORAL

DIAZEPAM

ACTAVIS ELIZABETH	2MG	A070781 001	Mar 19, 1986
	5MG	A070706 001	Mar 19, 1986
	10MG	A070707 001	Mar 19, 1986
DAVA PHARMS INC	10MG	A070228 001	Sep 26, 1985
DURAMED PHARMS BARR	2MG	A070894 001	Aug 27, 1986
	5MG	A070895 001	Aug 27, 1986
	10MG	A070896 001	Aug 27, 1986
FERNDALE LABS	2MG	A070903 001	Apr 01, 1987
	5MG	A070904 001	Apr 01, 1987
	10MG	A070905 001	Apr 01, 1987
HALSEY	2MG	A070987 001	Aug 15, 1986
	5MG	A070996 001	Aug 15, 1986
	10MG	A070956 001	Aug 15, 1986
IVAX SUB TEVA PHARMS	2MG	A070360 001	Sep 04, 1985
	5MG	A070361 001	Sep 04, 1985
	10MG	A070362 001	Sep 04, 1985
MARTEC USA LLC	10MG	A072402 001	Apr 25, 1989
PIONEER PHARMS	2MG	A070787 001	Aug 02, 1988
	5MG	A070788 001	Aug 02, 1988
	10MG	A070776 001	Aug 02, 1988
ROXANE	2MG	A070356 001	Jun 17, 1986
	5MG	A070357 001	Jun 17, 1986
	10MG	A070358 001	Jun 17, 1986
TEVA PHARMS	5MG	A070153 001	Nov 01, 1985
UPSHER-SMITH LABS	2MG	A070302 001	Dec 20, 1985
	5MG	A070303 001	Dec 20, 1985
	10MG	A070304 001	Dec 20, 1985
VIRTUS PHARMS	2MG	A070462 001	Feb 25, 1986
	5MG	A070463 001	Feb 25, 1986
	10MG	A070464 001	Feb 25, 1986
WARNER CHILCOTT	2MG	A070209 001	Sep 04, 1985
	5MG	A070210 001	Sep 04, 1985
	10MG	A070222 001	Sep 04, 1985
WATSON LABS	2MG	A070456 001	Nov 01, 1985
	5MG	A070457 001	Nov 01, 1985
	10MG	A070458 001	Nov 01, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-118(of 375)

\*\* See List Footnote

DIAZEPAMTABLET;ORAL  
Q-PAM

QUANTUM PHARMICS	2MG	A070423 001 Dec 12, 1985
	2MG	A072431 001 Apr 29, 1988
	5MG	A070424 001 Dec 12, 1985
	5MG	A072432 001 Apr 29, 1988
	10MG	A070425 001 Dec 12, 1985
	10MG	A072433 001 Apr 29, 1988

DIAZOXIDECAPSULE;ORAL  
PROGLYCEM

TEVA BRANDED PHARM	50MG	N017425 001
	100MG	N017425 002
INJECTABLE; INJECTION		
DIAZOXIDE		
ABRAXIS PHARM		
HYPERSTAT		
SCHERING		

ABRAXIS PHARM	15MG/ML	A071519 001 Aug 26, 1987
HYPERSTAT		
SCHERING		

DIBUCAINE HYDROCHLORIDEINJECTABLE; INJECTION  
HEAVY SOLUTION NUPERCAINE  
NOVARTIS

2.5MG/ML N006203 001

DICHLORPHENAMIDETABLET;ORAL  
DARANIDE  
+ STRONGBRIDGE US

50MG \*\* N011366 001

DICLOFENAC POTASSIUMTABLET;ORAL  
CATAFLAM

+ NOVARTIS	25MG **	N020142 001 Nov 24, 1993
	50MG **	N020142 002 Nov 24, 1993
DICLOFENAC POTASSIUM		
SANDOZ		
SUN PHARM INDUSTRIES		
WATSON LABS TEVA		

DICLOFENAC SODIUMSOLUTION;TOPICAL  
PENNSAID

+ NUVO PHARMS INC	1.5% **	N020947 001 Nov 04, 2009
SOLUTION/DROPS;OPHTHALMIC		

DICLOFENAC SODIUM

APOTEX INC	0.1%	A077600 001 Nov 13, 2008
FALCON PHARMS	0.1%	N020809 001 May 04, 1998

TABLET, DELAYED RELEASE;ORAL

DICLOFENAC SODIUM

ALLIED PHARMA INC	50MG	A074986 001 Feb 26, 1999
	75MG	A074986 002 Feb 26, 1999
PLIVA		
50MG		
ROXANE		
25MG		
50MG		
75MG		
TEVA		
50MG		
75MG		
TEVA PHARMS		
25MG		
50MG		
75MG		

VOLTAREN

+ NOVARTIS	25MG **	N019201 001 Jul 28, 1988
+	50MG **	N019201 002 Jul 28, 1988
+	75MG **	N019201 003 Jul 28, 1988

TABLET, EXTENDED RELEASE;ORAL

DICLOFENAC SODIUM

ACTAVIS ELIZABETH	100MG	A075910 001 Jan 07, 2002
-------------------	-------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-119(of 375)

\*\* See List Footnote

DICLOFENAC SODIUM

TABLET, EXTENDED RELEASE;ORAL

VOLTAREN-XR

+ NOVARTIS

100MG \*\*

N020254 001 Mar 08, 1996

DICLOxacillin sodium

CAPSULE;ORAL

DYCILL

GLAXOSMITHKLINE

EQ 250MG BASE

A060254 002

EQ 250MG BASE

A062238 001

EQ 500MG BASE

A060254 003

EQ 500MG BASE

A062238 002

PATHOCIL

WYETH AYERST

EQ 250MG BASE

N050011 002

EQ 500MG BASE

N050011 003 Mar 28, 1983

FOR SUSPENSION;ORAL

DICLOxacillin sodium

APOTHECON

EQ 62.5MG BASE/5ML

A061455 001

DYNAPEN

APOTHECON

EQ 62.5MG BASE/5ML

N050337 002

PATHOCIL

WYETH AYERST

EQ 62.5MG BASE/5ML

N050092 001

DICUMAROL

CAPSULE;ORAL

DICUMAROL

LILLY

25MG

N005509 003

50MG

N005509 001

TABLET;ORAL

DICUMAROL

ABBVIE

25MG

N005545 003

50MG

N005545 004

100MG

N005545 005

DICYCLOMINE HYDROCHLORIDE

CAPSULE;ORAL

DICYCLOMINE HYDROCHLORIDE

HIKMA PHARMS

10MG

A040204 001 Feb 28, 1997

PIONEER PHARMS

10MG

A089361 001 Jan 10, 1989

SUN PHARM INDUSTRIES

10MG

A084505 001 Oct 21, 1986

WATSON LABS

10MG

A083179 001 Feb 12, 1986

INJECTABLE;INJECTION

DICYCLOMINE HYDROCHLORIDE

WATSON LABS

10MG/ML

A080614 001 Feb 11, 1986

SYRUP;ORAL

BENTYL

+ APTALIS PHARMA US

10MG/5ML \*\*

N007961 002 Oct 15, 1984

DICYCLOMINE HYDROCHLORIDE

ALPHARMA US PHARMS

10MG/5ML

A084479 001

TABLET;ORAL

DICYCLOMINE HYDROCHLORIDE

HIKMA PHARMS

20MG

A040161 001 Oct 01, 1996

PIONEER PHARMS

20MG

A088585 001 Aug 20, 1986

SUN PHARM INDUSTRIES

20MG

A084600 001 Jul 29, 1985

WATSON LABS

20MG

A084361 001 Feb 06, 1986

DIDANOSINE

CAPSULE, DELAYED REL PELLETS;ORAL

DIDANOSINE

MYLAN PHARMS INC

125MG

A090788 001 Apr 08, 2010

200MG

A090788 002 Apr 08, 2010

250MG

A090788 003 Apr 08, 2010

400MG

A090788 004 Apr 08, 2010

FOR SOLUTION;ORAL

DIDANOSINE

AUROBINDO PHARMA

10MG/ML

A078112 001 Mar 08, 2007

VIDEX

BRISTOL MYERS SQUIBB

100MG/PACKET

N020155 003 Oct 09, 1991

167MG/PACKET

N020155 004 Oct 09, 1991

250MG/PACKET

N020155 005 Oct 09, 1991

375MG/PACKET

N020155 006 Oct 09, 1991

**DISCONTINUED DRUG PRODUCT LIST**

6-120(of 375)

\*\* See List Footnote

**DIDANOSINE**

TABLET, CHEWABLE;ORAL

VIDEX

+	BRISTOL MYERS SQUIBB	25MG **	N020154 002	Oct 09, 1991
+		50MG **	N020154 003	Oct 09, 1991
+		100MG **	N020154 004	Oct 09, 1991
+		150MG **	N020154 005	Oct 09, 1991
+		200MG **	N020154 006	Oct 28, 1999

TABLET, FOR SUSPENSION;ORAL

DIDANOSINE

AUROBINDO	100MG	A077275 001	Aug 14, 2012
	150MG	A077275 002	Aug 14, 2012
	200MG	A077275 003	Aug 14, 2012

**DIENESTROL**

CREAM;VAGINAL

DIENESTROL

ORTHO MCNEIL PHARM	0.01%	N006110 005
DV		
SANOFI AVENTIS US	0.01%	A083518 001
ESTRAGUARD		
SOLVAY	0.01%	A084436 001

SUPPOSITORY;VAGINAL

DV

SANOFI AVENTIS US	0.7MG	A083517 001
-------------------	-------	-------------

**DIETHYLCARBAMAZINE CITRATE**

TABLET;ORAL

HETRAZAN

LEDERLE	50MG	N006459 001
---------	------	-------------

**DIETHYLPROPION HYDROCHLORIDE**

TABLET;ORAL

DIETHYLPROPION HYDROCHLORIDE

CHARTWELL RX	25MG	A088267 001	Aug 25, 1983
	25MG	A088268 001	Aug 25, 1983
EPIC PHARMA LLC	25MG	A040828 001	Nov 05, 2008
SANDOZ	25MG	A085916 001	
TEVA	25MG	A088642 001	Sep 20, 1984
UCB INC	25MG	A085544 001	
WATSON LABS	25MG	A085741 001	
TENUATE			
SANOFI AVENTIS US	25MG	N017668 001	
TEPANIL			
3M	25MG	N011673 001	

TABLET, EXTENDED RELEASE;ORAL

TENUATE

SANOFI AVENTIS US	75MG	N017669 001
-------------------	------	-------------

TEPANIL TEN-TAB

3M	75MG	N017956 001
----	------	-------------

**DIETHYLSTILBESTROL**

INJECTABLE;INJECTION

STILBESTROL

BRISTOL MYERS SQUIBB	0.2MG/ML	N004056 003	
	0.5MG/ML	N004056 004	
	1MG/ML	N004056 005	
	5MG/ML	N004056 006	

SUPPOSITORY;VAGINAL

DIETHYLSTILBESTROL

LILLY	0.1MG	N004040 001	
	0.5MG	N004040 002	

STILBESTROL

BRISTOL MYERS SQUIBB	0.1MG	N004056 001	
	0.5MG	N004056 002	

TABLET;ORAL

DIETHYLSTILBESTROL

LILLY	0.1MG	N004041 002	
	0.5MG	N004041 003	
	1MG	N004041 004	
	5MG	N004041 005	

**DISCONTINUED DRUG PRODUCT LIST**

6-121(of 375)

\*\* See List Footnote

DIETHYLSILBESTROL

TABLET;ORAL

STILBESTROL

TABLICAPS

0.5MG

1MG

5MG

A083004 001

A083002 001

A083006 001

STILBETIN

BRISTOL MYERS SQUIBB

0.1MG

0.25MG

0.5MG

1MG

5MG

N004056 007

N004056 017

N004056 008

N004056 009

N004056 010

TABLET, DELAYED RELEASE;ORAL

DIETHYLSILBESTROL

LILLY

0.1MG

0.25MG

0.5MG

1MG

5MG

N004039 002

N004039 005

N004039 003

N004039 004

N004039 006

STILBESTROL

TABLICAPS

0.5MG

1MG

5MG

A083003 001

A083005 001

A083007 001

STILBETIN

BRISTOL MYERS SQUIBB

0.1MG

0.5MG

1MG

5MG

N004056 011

N004056 012

N004056 013

N004056 014

DIETHYLSILBESTROL DIPHOSPHATE

INJECTABLE; INJECTION

STILPHOSTROL

BAYER PHARMS

250MG/5ML

N010010 001

TABLET;ORAL

STILPHOSTROL

BAYER PHARMS

50MG

N010010 002

DIFLORASONE DIACETATE

CREAM;TOPICAL

DIFLORASONE DIACETATE

FOUGERA PHARMS

0.05%

A075187 001 Mar 30, 1998

FLORONE

PHARMACIA AND UPJOHN

0.05% \*\*

N017741 001

FLORONE E

PHARMACIA AND UPJOHN

0.05%

N019259 001 Aug 28, 1985

PSORCON

+ TARO PHARMS NORTH

0.05% \*\*

N020205 001 Nov 20, 1992

OINTMENT;TOPICAL

PSORCON

+ PHARMACIA AND UPJOHN

0.05%

N019260 001 Aug 28, 1985

PSORCON E

PHARMACIA AND UPJOHN

0.05%

N017994 001

DIFLUNISAL

TABLET;ORAL

DIFLUNISAL

ALLIED PHARMA INC

250MG

A073562 001 Nov 27, 1992

500MG

A073563 001 Nov 27, 1992

IDT AUSTRALIA LTD

500MG

A074604 001 Jun 10, 1996

PUREPAC PHARM

250MG

A074285 001 May 07, 1996

500MG

A074285 002 May 07, 1996

TEVA

250MG

A073679 001 Jul 31, 1992

WATSON LABS

250MG

A074400 001 Jul 17, 1997

500MG

A074400 002 Jul 17, 1997

DOLOBID

+ MERCK

250MG \*\*

N018445 001 Apr 19, 1982

+

500MG \*\*

N018445 002 Apr 19, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-122(of 375)

\*\* See List Footnote

**DIGITOXIN**

INJECTABLE; INJECTION

CRYSTODIGIN

LILLY

0.2MG/ML

A084100 005

**DIGOXIN**

CAPSULE; ORAL

LANOXICAPS

GLAXOSMITHKLINE LLC  
0.05MG  
0.1MG  
0.15MG  
0.2MGN018118 002 Jul 26, 1982  
N018118 003 Jul 26, 1982  
N018118 004 Sep 24, 1984  
N018118 001 Jul 26, 1982

INJECTABLE; INJECTION

DIGOXIN

ABRAXIS PHARM  
HOSPIRA  
WYETH AYERST  
DIGOXIN PEDIATRIC  
HOSPIRA  
0.25MG/ML  
0.25MG/ML  
0.25MG/ML  
0.25MG/ML  
0.1MG/MLA083217 001  
A040093 001 May 16, 1996  
A040206 001 Aug 28, 1998  
A084386 001  
A040092 001 Apr 25, 1996

TABLET; ORAL

LANOXIN

CONCORDIA PHARMS INC  
0.375MG  
0.5MGN020405 005 Sep 30, 1997  
N020405 006 Sep 30, 1997**DIHYDROERGOTAMINE MESYLATE; HEPARIN SODIUM; LIDOCAINE HYDROCHLORIDE**

INJECTABLE; INJECTION

EMBOLEX

NOVARTIS  
0.5MG/0.5ML; 2,500  
UNITS/0.5ML; 5.33MG/0.5ML  
0.5MG/0.7ML; 5,000  
UNITS/0.7ML; 7.46MG/0.7MLN018885 001 Nov 30, 1984  
N018885 002 Nov 30, 1984**DILTIAZEM HYDROCHLORIDE**

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM SR

+ BIOVAIL  
+  
+  
+  
120MG \*\*  
180MG \*\*  
180MG \*\*N019471 001 Jan 23, 1989  
N019471 002 Jan 23, 1989  
N019471 003 Jan 23, 1989  
N019471 004 Jan 23, 1989

DILACOR XR

+ ALLERGAN SALES LLC  
+  
+  
120MG \*\*  
180MG \*\*  
240MG \*\*N020092 001 May 29, 1992  
N020092 002 May 29, 1992  
N020092 003 May 29, 1992

DILT-CD

APOTEX  
120MG  
180MG  
240MG  
300MGA076151 001 May 20, 2004  
A076151 002 May 20, 2004  
A076151 003 May 20, 2004  
A076151 004 May 20, 2004

DILTIAZEM HYDROCHLORIDE

ACTAVIS LABS FL INC  
120MG  
180MG  
240MGA074852 001 Oct 10, 1997  
A074852 002 Oct 10, 1997  
A074852 003 Oct 10, 1997BIOVAIL  
60MG  
90MG  
120MG  
120MG  
180MG  
240MG  
300MG  
360MG  
420MGA074845 001 Sep 15, 1999  
A074845 002 Sep 15, 1999  
A074845 003 Sep 15, 1999  
N020939 001 Jan 28, 2000  
N020939 002 Jan 28, 2000  
N020939 003 Jan 28, 2000  
N020939 004 Jan 28, 2000  
N020939 005 Sep 14, 2001  
N020939 006 Sep 14, 2001NESHER PHARMS  
120MG  
180MG  
240MG  
300MG  
360MG  
420MGA076563 002 Sep 12, 2006  
A076563 003 Sep 12, 2006  
A076563 004 Sep 12, 2006  
A076563 005 Sep 12, 2006  
A076563 006 Sep 12, 2006  
A076563 001 Sep 12, 2006TEVA  
60MG  
90MG  
120MGA074079 001 Nov 30, 1993  
A074079 002 Nov 30, 1993  
A074079 003 Nov 30, 1993

**DISCONTINUED DRUG PRODUCT LIST**

6-123(of 375)

\*\* See List Footnote

DILTIAZEM HYDROCHLORIDE

INJECTABLE; INJECTION

CARDIZEM

BIOVAIL	100MG/VIAL	N020792 001	Sep 05, 1997
+ BIOVAIL LABS INTL	5MG/ML **	N020027 001	Oct 24, 1991
+	25MG/VIAL **	N020027 003	Aug 18, 1995

DILTIAZEM HYDROCHLORIDE

HOSPIRA	5MG/ML	A075004 001	Feb 16, 2000
	5MG/ML	A075106 001	Apr 29, 1999
MYLAN LABS LTD	5MG/ML	A075375 001	Sep 30, 1999
TEVA PHARMS USA	5MG/ML	A074894 001	Aug 26, 1997

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

APOTHECON	30MG	A074051 001	Mar 31, 1993
	60MG	A074051 002	Mar 31, 1993
	90MG	A074051 003	Mar 31, 1993
	120MG	A074051 004	Mar 31, 1993
CHARTWELL MOLECULES	30MG	A074093 001	Nov 05, 1992
	60MG	A074093 002	Nov 05, 1992
	90MG	A074093 003	Nov 05, 1992
	120MG	A074093 004	Nov 05, 1992
IVAX SUB TEVA PHARMS	30MG	A074168 001	Mar 03, 1995
	60MG	A074168 002	Mar 03, 1995
	90MG	A074168 003	Mar 03, 1995
	120MG	A074168 004	Mar 03, 1995
TEVA	30MG	A074084 001	Feb 25, 1994
	60MG	A074084 002	Feb 25, 1994
TEVA PHARMS	30MG	A074067 001	Nov 05, 1992
	60MG	A074067 002	Nov 05, 1992
	90MG	A074067 003	Nov 05, 1992
	120MG	A074067 004	Nov 05, 1992

DILTIAZEM MALATE

TABLET, EXTENDED RELEASE; ORAL

TIAMATE

MERCK	EQ 120MG HCL	N020506 001	Oct 04, 1996
	EQ 180MG HCL	N020506 002	Oct 04, 1996
	EQ 240MG HCL	N020506 003	Oct 04, 1996

DILTIAZEM MALATE; ENALAPRIL MALEATE

TABLET, EXTENDED RELEASE; ORAL

TECZEM

BIOVAIL	EQ 180MG HCL; 5MG	N020507 001	Oct 04, 1996
---------	-------------------	-------------	--------------

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

BAXTER HLTHCARE	50MG/ML	A084767 001	
WATSON LABS	50MG/ML	A083531 001	
WATSON LABS TEVA	50MG/ML	A080615 001	
WYETH AYERST	50MG/ML	A084316 001	

LIQUID; ORAL

DIMENHYDRINATE

ALRA	12.5MG/4ML	A080715 001	
------	------------	-------------	--

TABLET; ORAL

DIMENHYDRINATE

HEATHER	50MG	A080841 001	
NEXGEN PHARMA INC	50MG	A085985 001	
WATSON LABS	50MG	A085166 001	

DIMYRISTOYL LECITHIN; PERFLUXANE

INJECTABLE; INTRAVENOUS

IMAGENT

VESSELON SPV LLC	0.92MG/VIAL; 0.092MG/VIAL	N021191 001	May 31, 2002
------------------	---------------------------	-------------	--------------

DINOPROST TROMETHAMINE

INJECTABLE; INJECTION

PROSTIN F2 ALPHA

PHARMACIA AND UPJOHN	EQ 5MG BASE/ML	N017434 001	
----------------------	----------------	-------------	--

**DISCONTINUED DRUG PRODUCT LIST**

6-124(of 375)

\*\* See List Footnote

DIPHEMANIL METHYLSULFATETABLET;ORAL  
PRANTAL

SCHERING 100MG

N008114 004

DIPHENHYDRAMINE HYDROCHLORIDECAPSULE;ORAL  
BENADRYL

MCNEIL CONS	25MG	N005845 007
	50MG	N005845 001
DIPHENHYDRAMINE HYDROCHLORIDE		
ALRA	25MG	A080519 004
	50MG	A080519 003
ANABOLIC	50MG	A083275 001
ELKINS SINK	25MG	A085701 001
	50MG	A085701 002
HALSEY	50MG	A087914 001 Jun 04, 1984
HEATHER	25MG	A084524 001
	50MG	A083953 001
HIKMA INTL PHARMS	50MG	A083567 001
IMPAX LABS	25MG	A080807 001
	50MG	A080807 002
IVAX SUB TEVA PHARMS	25MG	A080762 001
	50MG	A080762 002
LANNETT	25MG	A080868 001
	50MG	A080868 002
LEDERLE	25MG	A086874 001
	50MG	A086875 001
LNK	25MG	A087977 001 Jan 27, 1983
	50MG	A087978 001 Jan 27, 1983
MK LABS	25MG	A083087 001
	50MG	A083087 002
MUTUAL PHARM	25MG	A084506 001
NEWTRON PHARMS	25MG	A086543 001
	50MG	A086544 001
NEXGEN PHARMA INC	25MG	A083634 001
PERRIGO	25MG	A083061 001
	50MG	A083061 002
PIONEER PHARMS	25MG	A089101 001 Dec 20, 1985
	50MG	A088880 001 Dec 20, 1985
PUREPAC PHARM	25MG	A085156 001
	50MG	A085150 001
PVT FORM	25MG	A083027 001
	50MG	A083027 002
ROXANE	50MG	A080635 001
SANDOZ	25MG	A080832 001
	25MG	A080845 002
	50MG	A080832 002
	50MG	A080845 001
SUN PHARM INDUSTRIES	25MG	A089488 001 Jan 02, 1987
	50MG	A089489 001 Jan 02, 1987
SUPERPHARM	25MG	A089040 001 May 15, 1985
	50MG	A089041 001 May 15, 1985
TEVA	25MG	A085874 001
	50MG	A085874 002
VALEANT PHARM INTL	25MG	A080596 001
	50MG	A080592 001
VANGARD	25MG	A088034 001 Oct 27, 1982
	50MG	A087630 001
WATSON LABS	25MG	A080728 001
	25MG	A083797 001
	25MG	A085138 001
	50MG	A080727 001
	50MG	A083797 002
	50MG	A085083 001
WHITEWORTH TOWN PLSN	25MG	A083441 001
	50MG	A080800 001

**DISCONTINUED DRUG PRODUCT LIST**

6-125(of 375)

\*\* See List Footnote

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR;ORAL			
BELIX			
HALSEY	12.5MG/5ML	A086586	001 Oct 03, 1983
BENADRYL		N005845	004
MCNEIL CONS	12.5MG/5ML		
DIBENIL		A088304	001 Dec 16, 1983
CENCI	12.5MG/5ML		
DIPHEN		A084640	001
USL PHARMA	12.5MG/5ML		
DIPHENHYDRAMINE HYDROCHLORIDE			
BUNDY	12.5MG/5ML	A083674	001
CENCI	12.5MG/5ML	A087941	001 Dec 17, 1982
KV PHARM	12.5MG/5ML	A085621	001
LANNETT	12.5MG/5ML	A080939	002
LEDERLE	12.5MG/5ML	A086937	001
MK LABS	12.5MG/5ML	A083088	002
NASKA	12.5MG/5ML	A088680	001 May 31, 1985
PERRIGO	12.5MG/5ML	A083063	001
PUREPAC PHARM	12.5MG/5ML	A083237	001 Jan 25, 1982
PVT FORM	12.5MG/5ML	A085287	001
ROXANE	12.5MG/5ML	A080643	001
HYDRAMINE			
ALPHARMA US PHARMS	12.5MG/5ML	A080763	002
INJECTABLE;INJECTION			
BENADRYL			
MCNEIL CONS	10MG/ML	N006146	001
+	50MG/ML **	N006146	002
BENADRYL PRESERVATIVE FREE			
+	MCNEIL CONS	50MG/ML **	N009486 001
DIPHENHYDRAMINE HYDROCHLORIDE			
BEL MAR	10MG/ML	A080822	001
EUROHLTH INTL SARL	50MG/ML	A083183	001
LYPHOMED	10MG/ML	A087066	001
WATSON LABS	10MG/ML	A083533	001
WATSON LABS TEVA	10MG/ML	A080873	001
	50MG/ML	A080873	002
WYETH AYERST	50MG/ML	A080577	001
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE			
ABRAXIS PHARM	50MG/ML	A080586	002
INTL MEDICATION	50MG/ML	A084094	001
WATSON LABS TEVA	50MG/ML	A080873	003
SYRUP;ORAL			
ANTITUSSIVE			
PERRIGO	12.5MG/5ML	A071292	001 Apr 10, 1987
BELDIN			
HALSEY	12.5MG/5ML	A089179	001 Jun 05, 1986
BENYLIN			
PARKE DAVIS	12.5MG/5ML	N006514	004
DIPHEN			
MORTON GROVE	12.5MG/5ML	A070118	001 Oct 01, 1985
DIPHENHYDRAMINE HYDROCHLORIDE			
ALPHARMA US PHARMS	12.5MG/5ML	A070497	001 Apr 25, 1989
CUMBERLAND SWAN	12.5MG/5ML	A073611	001 Aug 20, 1992
HI TECH PHARMA	12.5MG/5ML	A072416	001 Sep 28, 1990
HYDRAMINE			
ALPHARMA US PHARMS	12.5MG/5ML	A070205	001 Jan 28, 1986
SILPHEN			
SILARX	12.5MG/5ML	A072646	001 Feb 27, 1992
VICKS FORMULA 44			
WARNER CHILCOTT	12.5MG/5ML	A070524	001 Jan 14, 1987
<u>DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE</u>			
SOLUTION;ORAL			
BENYLIN			
PARKE DAVIS	12.5MG/5ML;30MG/5ML	N019014	001 Jun 11, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-126(of 375)

\*\* See List Footnote

DIPHENIDOL HYDROCHLORIDETABLET;ORAL  
VONTROL

GLAXOSMITHKLINE EQ 25MG BASE

N016033 001

DIPHENYL PYRALINE HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL  
HISPRIL

GLAXOSMITHKLINE 5MG

N011945 001

DIPIVEFRIN HYDROCHLORIDESOLUTION/DROPS;OPHTHALMIC  
AKPRO

AKORN 0.1%

A074382 001 Sep 29, 1995

DIPIVEFRIN HYDROCHLORIDE

BAUSCH AND LOMB 0.1%

A074188 001 May 19, 1995

FALCON PHARMS 0.1%

A073636 001 Jun 30, 1994

PROPINE

ALLERGAN 0.1%

N018239 001

DIPYRIDAMOLE

INJECTABLE;INJECTION

DIPYRIDAMOLE

HOSPIRA 5MG/ML

A074601 001 Dec 19, 1997

MYLAN LABS LTD 5MG/ML

A075769 001 Nov 27, 2002

TEVA PHARMS USA 5MG/ML

A074952 001 Nov 26, 1997

IV PERSANTINE

+ BOEHRINGER INGELHEIM 5MG/ML \*\*

N019817 001 Dec 13, 1990

TABLET;ORAL

DIPYRIDAMOLE

GLENMARK GENERICS 25MG

A088999 001 Feb 05, 1991

50MG

A089000 001 Feb 05, 1991

75MG

A089001 001 Feb 05, 1991

IDT AUSTRALIA LTD 25MG

A086944 002 Apr 16, 1991

50MG

A086944 001 Feb 25, 1992

75MG

A086944 003 Feb 25, 1992

LANNETT 25MG

A040898 001 Apr 23, 2008

50MG

A040898 002 Apr 23, 2008

75MG

A040898 003 Apr 23, 2008

OXFORD PHARMS 25MG

A040542 001 Apr 21, 2006

50MG

A040542 002 Apr 21, 2006

75MG

A040542 003 Apr 21, 2006

PUREPAC PHARM 25MG

A089425 001 Jul 12, 1990

50MG

A089426 001 Jul 12, 1990

75MG

A089427 001 Jul 12, 1990

WATSON LABS 50MG

A087160 001 Jun 07, 1996

DIRITHROMYCIN

TABLET, DELAYED RELEASE;ORAL

DYNABAC

LILLY RES LABS 250MG

N050678 001 Jun 19, 1995

DISOPYRAMIDE PHOSPHATE

CAPSULE;ORAL

DISOPYRAMIDE PHOSPHATE

AUROLIFE PHARMA LLC EQ 100MG BASE

A070470 001 Dec 10, 1985

EQ 150MG BASE

A070471 001 Dec 10, 1985

INTERPHARM EQ 100MG BASE

A071190 001 Jan 15, 1987

EQ 150MG BASE

A071191 001 Jan 15, 1987

IVAX SUB TEVA PHARMS EQ 100MG BASE

A070186 001 Nov 18, 1985

EQ 150MG BASE

A070187 001 Nov 18, 1985

MYLAN EQ 100MG BASE

A070138 001 Jun 14, 1985

EQ 150MG BASE

A070139 001 Jun 14, 1985

SUN PHARM INDUSTRIES EQ 100MG BASE

A070351 001 Dec 17, 1985

EQ 150MG BASE

A070352 001 Dec 17, 1985

SUPERPHARM EQ 100MG BASE

A070940 001 Feb 09, 1987

EQ 150MG BASE

A070941 001 Feb 09, 1987

WATSON LABS EQ 100MG BASE

A070240 001 Feb 02, 1986

EQ 150MG BASE

A070241 001 Feb 02, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-127(of 375)

\*\* See List Footnote

DISOPYRAMIDE PHOSPHATE

CAPSULE, EXTENDED RELEASE;ORAL

DISOPYRAMIDE PHOSPHATE

NESHER PHARMS EQ 150MG BASE

A071200 001 Dec 15, 1987

DISULFIRAM

TABLET;ORAL

ANTABUSE

+ TEVA WOMENS 250MG \*\*  
+ 500MG \*\*N007883 003  
N007883 002

DISULFIRAM

PAR PHARM 250MG A088792 001 Aug 14, 1984  
500MG A088793 001 Aug 14, 1984  
WATSON LABS 250MG A086889 001  
250MG A087973 001 Aug 05, 1983  
500MG A087974 001 Aug 05, 1983  
WATSON LABS TEVA 500MG A086890 001DIVALPROEX SODIUM

TABLET, DELAYED RELEASE;ORAL

DEPAKOTE CP

ABBOTT EQ 250MG BASE N019794 001 Jul 11, 1990  
EQ 500MG BASE N019794 002 Jul 11, 1990

DIVALPROEX SODIUM

MYLAN EQ 125MG VALPROIC ACID A077254 001 Jul 29, 2008  
EQ 250MG VALPROIC ACID A077254 002 Jul 29, 2008  
EQ 500MG VALPROIC ACID A077254 003 Jul 29, 2008

TABLET, EXTENDED RELEASE;ORAL

DIVALPROEX SODIUM

G AND W LABS INC EQ 500MG VALPROIC ACID A078700 001 Aug 03, 2009

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

BAXTER HLTHCARE EQ 12.5MG BASE/ML A074381 001 Sep 26, 1996  
HOSPIRA EQ 1.25GM BASE/100ML A074634 001 Sep 27, 1996  
LUITPOLD EQ 12.5MG BASE/ML A074545 001 Jun 25, 1998  
TELIGENT PHARMA INC EQ 12.5MG BASE/ML A074098 001 Feb 21, 1995  
TEVA PARENTERAL EQ 12.5MG BASE/ML A074206 001 Oct 19, 1993  
WATSON LABS EQ 12.5MG BASE/ML A074114 001 Nov 30, 1993  
WATSON LABS INC EQ 12.5MG BASE/ML A074279 001 Feb 18, 1998  
EQ 12.5MG BASE/ML A074995 001 Mar 31, 1998

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%

HOSPIRA EQ 50MG BASE/100ML N020269 001 Oct 19, 1993  
EQ 100MG BASE/100ML N020269 002 Oct 19, 1993  
EQ 200MG BASE/100ML N020269 003 Oct 19, 1993

DOBUTREX

+ LILLY EQ 12.5MG BASE/ML

N017820 002

DOCETAXEL

INJECTABLE; INJECTION

DOCEFREZ

+ SUN PHARMA GLOBAL 20MG/VIAL N022534 001 May 03, 2011  
+ 80MG/VIAL N022534 002 May 03, 2011

DOCETAXEL

APOTEX INC 20MG/0.5ML (40MG/ML) N022312 001 Jan 11, 2012  
80MG/2ML (40MG/ML) N022312 002 Jan 11, 2012  
+ HOSPIRA INC 120MG/6ML (20MG/ML) N022234 006 Jun 23, 2016  
PFIZER LABS 20MG/2ML (10MG/ML) N202356 001 Mar 13, 2014  
80MG/8ML (10MG/ML) N202356 002 Mar 13, 2014  
130MG/13ML (10MG/ML) N202356 003 Mar 13, 2014  
200MG/20ML (10MG/ML) N202356 004 Mar 13, 2014

TAXOTERE

+ SANOFI AVENTIS US 40MG/ML \*\*

N020449 001 May 14, 1996

DOLASETRON MESYLATE

INJECTABLE; INJECTION

ANZEMET

+ US PHARM HOLDINGS 12.5MG/0.625ML (20MG/ML) N020624 002 Sep 11, 1997  
+ 100MG/5ML (20MG/ML) N020624 001 Sep 11, 1997  
500MG/25ML (20MG/ML) N020624 003 Dec 11, 2001

**DISCONTINUED DRUG PRODUCT LIST**

6-128(of 375)

\*\* See List Footnote

DOLASETRON MESYLATE

TABLET;ORAL

ANZEMET

+ US PHARM HOLDINGS	50MG	N020623 001	Sep 11, 1997
+	100MG	N020623 002	Sep 11, 1997

DONEPEZIL HYDROCHLORIDE

SOLUTION;ORAL

ARICEPT

EISAI INC	5MG/5ML	N021719 001	Oct 18, 2004
-----------	---------	-------------	--------------

TABLET;ORAL

DONEPEZIL HYDROCHLORIDE

ACCORD HLTHCARE	5MG	A201335 001	Aug 29, 2011
	10MG	A201335 002	Aug 29, 2011
HIKMA PHARMS	5MG	A090247 001	May 31, 2011
	10MG	A090247 002	May 31, 2011

TABLET, ORALLY DISINTEGRATING;ORAL

DONEPEZIL HYDROCHLORIDE

SUN PHARM INDUSTRIES	5MG	A077975 002	Dec 11, 2009
	10MG	A077975 001	Dec 11, 2009

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

ABBOTT	40MG/ML	A070656 001	Jan 24, 1989
	80MG/ML	A070657 001	Jan 24, 1989
ABRAXIS PHARM	40MG/ML	A070012 001	Jun 12, 1985
	40MG/ML	A070058 001	Mar 20, 1985
	80MG/ML	A070013 001	Jun 12, 1985
	80MG/ML	A070059 001	Mar 20, 1985
	160MG/ML	A070364 001	Dec 04, 1985
BAXTER HLTHCARE	40MG/ML	N018398 001	
	80MG/ML	N018398 002	Mar 22, 1982
HOSPIRA	40MG/ML	A074403 001	May 23, 1996
IGI LABS INC	40MG/ML	A070087 001	Oct 23, 1985
	80MG/ML	A070089 001	Oct 23, 1985
	80MG/ML	A070090 001	Oct 23, 1985
	80MG/ML	A070091 001	Oct 23, 1985
	160MG/ML	A070092 001	Oct 23, 1985
	160MG/ML	A070093 001	Oct 23, 1985
	160MG/ML	A070094 001	Oct 23, 1985
INTL MEDICATION	40MG/ML	N018014 001	
LYPHOMED	40MG/ML	N018549 001	Mar 11, 1983
SMITH AND NEPHEW	40MG/ML	A070011 001	Aug 29, 1985
	40MG/ML	A070046 001	Aug 29, 1985
	80MG/ML	A070047 001	Aug 29, 1985
TELIGENT	40MG/ML	N018656 001	Jun 28, 1983
TEVA PARENTERAL	40MG/ML	A072999 001	Oct 23, 1991
	80MG/ML	A073000 001	Oct 23, 1991
WARNER CHILCOTT	40MG/ML	A070558 001	Sep 20, 1985
	40MG/ML	N018138 001	
	80MG/ML	A070559 001	Sep 20, 1985
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5%			
HOSPIRA	1.6MG/ML	N020542 001	Aug 30, 1995
INTROPIN			
HOSPIRA	40MG/ML	N017395 001	
	80MG/ML	N017395 002	
	160MG/ML	N017395 003	

DORIPENEM

INJECTABLE;IV (INFUSION)

DORIBAX

+ SHIONOGI INC	250MG/VIAL	N022106 002	Oct 05, 2010
+	500MG/VIAL	N022106 001	Oct 12, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-129(of 375)

\*\* See List Footnote

DORZOLAMIDE HYDROCHLORIDESOLUTION/DROPS;OPHTHALMIC  
DORZOLAMIDE HYDROCHLORIDE

APOTEX INC	EQ 2% BASE	A078395 001 Oct 28, 2008
ZAMBON SPA	EQ 2% BASE	A091034 001 Dec 04, 2013

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS;OPHTHALMIC

DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE		
APOTEX INC	EQ 2% BASE;EQ 0.5% BASE	A078201 001 Oct 28, 2008
LANNETT HOLDINGS INC	EQ 2% BASE;EQ 0.5% BASE	A201998 001 Dec 17, 2014

DOXACURIUM CHLORIDE

INJECTABLE; INJECTION

NUROMAX		
ABBVIE	EQ 1MG BASE/ML	N019946 001 Mar 07, 1991

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOXAPRAM HYDROCHLORIDE		
WATSON LABS	20MG/ML	A073529 001 Jan 30, 1992

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE		
ACTAVIS ELIZABETH	EQ 1MG BASE	A075574 001 Oct 18, 2000
	EQ 2MG BASE	A075574 002 Oct 18, 2000
	EQ 4MG BASE	A075574 003 Oct 18, 2000
	EQ 8MG BASE	A075574 004 Oct 18, 2000
FOSUN PHARMA	EQ 1MG BASE	A075646 001 Oct 18, 2000
	EQ 2MG BASE	A075646 002 Oct 18, 2000
	EQ 4MG BASE	A075646 003 Oct 18, 2000
	EQ 8MG BASE	A075646 004 Oct 18, 2000
GENPHARM	EQ 1MG BASE	A075466 001 Oct 18, 2000
	EQ 2MG BASE	A075466 002 Oct 18, 2000
	EQ 4MG BASE	A075466 003 Oct 18, 2000
	EQ 8MG BASE	A075466 004 Oct 18, 2000
IDT AUSTRALIA LTD	EQ 1MG BASE	A075432 001 Oct 18, 2000
	EQ 2MG BASE	A075432 002 Oct 18, 2000
	EQ 4MG BASE	A075432 003 Oct 18, 2000
	EQ 8MG BASE	A075432 004 Oct 18, 2000
IVAX SUB TEVA PHARMS	EQ 1MG BASE	A075453 001 Oct 18, 2000
	EQ 2MG BASE	A075453 002 Oct 18, 2000
	EQ 4MG BASE	A075453 003 Oct 18, 2000
	EQ 8MG BASE	A075453 004 Oct 18, 2000
NESHER PHARMS	EQ 1MG BASE	A075609 001 Oct 18, 2000
	EQ 2MG BASE	A075609 002 Oct 18, 2000
	EQ 4MG BASE	A075609 003 Oct 18, 2000
	EQ 8MG BASE	A075609 004 Oct 18, 2000
TEVA	EQ 1MG BASE	A075353 001 Jan 12, 2001
	EQ 2MG BASE	A075353 002 Jan 12, 2001
	EQ 4MG BASE	A075353 003 Jan 12, 2001
	EQ 8MG BASE	A075353 004 Jan 12, 2001
WATSON LABS INC	EQ 1MG BASE	A075426 001 Oct 18, 2000
	EQ 2MG BASE	A075426 002 Oct 18, 2000
	EQ 4MG BASE	A075426 003 Oct 18, 2000
	EQ 8MG BASE	A075426 004 Oct 18, 2000

DOXEPIPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIPIN HYDROCHLORIDE		
DAVA PHARMS INC	EQ 10MG BASE	A071685 001 Jan 05, 1988
	EQ 25MG BASE	A071686 001 Jan 05, 1988
	EQ 50MG BASE	A071673 001 Jan 05, 1988
	EQ 75MG BASE	A071674 001 Jan 05, 1988
	EQ 100MG BASE	A071675 001 Jan 05, 1988
	EQ 150MG BASE	A071676 001 Jan 05, 1988
NEW RIVER	EQ 10MG BASE	N016987 001
	EQ 25MG BASE	N016987 002
	EQ 50MG BASE	N016987 003
	EQ 75MG BASE	N016987 006
	EQ 100MG BASE	N016987 004

**DISCONTINUED DRUG PRODUCT LIST**

6-130(of 375)

\*\* See List Footnote

DOXE PIN HYDROCHLORIDE

CAPSULE; ORAL

DOXE PIN HYDROCHLORIDE

	EQ 150MG BASE	N016987 007	Apr 13, 1987
PAR PHARM	EQ 10MG BASE	A071697 001	Nov 09, 1987
	EQ 25MG BASE	A071437 001	Nov 09, 1987
	EQ 50MG BASE	A071595 001	Nov 09, 1987
	EQ 75MG BASE	A071608 001	Nov 09, 1987
	EQ 100MG BASE	A071422 001	Nov 09, 1987
PUREPAC PHARM	EQ 10MG BASE	A073054 001	Dec 28, 1990
	EQ 25MG BASE	A072109 001	Dec 28, 1990
	EQ 50MG BASE	A073055 001	Dec 28, 1990
	EQ 75MG BASE	A072386 001	Sep 08, 1988
	EQ 100MG BASE	A072110 001	Sep 08, 1988
	EQ 150MG BASE	A072387 001	Sep 08, 1988
QUANTUM PHARMICS	EQ 10MG BASE	A070972 001	Sep 29, 1987
	EQ 25MG BASE	A070973 001	Sep 29, 1987
	EQ 50MG BASE	A070931 001	Sep 29, 1987
	EQ 75MG BASE	A070932 001	Sep 29, 1987
	EQ 100MG BASE	A072375 001	Mar 15, 1989
	EQ 150MG BASE	A072376 001	Mar 15, 1989
SANDOZ	EQ 10MG BASE	A071487 001	Mar 02, 1987
	EQ 25MG BASE	A070827 001	May 15, 1986
	EQ 50MG BASE	A070828 001	May 15, 1986
	EQ 75MG BASE	A070825 001	May 15, 1986
	EQ 100MG BASE	A071562 001	Mar 02, 1987
SUN PHARM INDUSTRIES	EQ 25MG BASE	A071502 001	Feb 18, 1988
	EQ 50MG BASE	A071653 001	Feb 18, 1988
	EQ 75MG BASE	A071654 001	Feb 18, 1988
	EQ 100MG BASE	A071521 001	Feb 18, 1988
WATSON LABS	EQ 10MG BASE	A070952 001	Mar 04, 1987
	EQ 10MG BASE	A071485 001	Apr 30, 1987
	EQ 10MG BASE	A072985 001	Mar 29, 1991
	EQ 25MG BASE	A070953 001	May 15, 1986
	EQ 25MG BASE	A071486 001	Apr 30, 1987
	EQ 25MG BASE	A072986 001	Mar 29, 1991
	EQ 50MG BASE	A070954 001	May 15, 1986
	EQ 50MG BASE	A071238 001	Apr 30, 1987
	EQ 75MG BASE	A071326 001	Apr 30, 1987
	EQ 75MG BASE	A071763 001	Feb 09, 1988
	EQ 100MG BASE	A070955 001	May 15, 1986
	EQ 100MG BASE	A071239 001	Apr 30, 1987
	EQ 150MG BASE	A071764 001	Feb 09, 1988
WATSON LABS TEVA	EQ 50MG BASE	A072987 001	Mar 29, 1991
SINEQUAN			
+ PFIZER	EQ 10MG BASE **	N016798 003	
+	EQ 25MG BASE **	N016798 001	
+	EQ 50MG BASE **	N016798 002	
+	EQ 75MG BASE **	N016798 006	
+	EQ 100MG BASE **	N016798 005	
+	EQ 150MG BASE **	N016798 007	
CONCENTRATE; ORAL			
DOXE PIN HYDROCHLORIDE			
PHARM ASSOC	EQ 10MG BASE/ML	A075924 001	Jan 15, 2004
SINEQUAN			
+ PFIZER	EQ 10MG BASE/ML **	N017516 001	
TABLET; ORAL			
DOXE PIN HYDROCHLORIDE			
MYLAN PHARMS INC	EQ 3MG BASE	A202337 001	Jan 20, 2016
	EQ 6MG BASE	A202337 002	Jan 20, 2016

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

PHARMACIA AND UPJOHN	2MG/ML	A063165 001	Jan 30, 1991
	200MG/100ML	A063165 002	Jan 30, 1991

DOXORUBICIN HYDROCHLORIDE			
PHARMACIA AND UPJOHN	10MG/VIAL	N050467 001	
	20MG/VIAL	N050467 003	May 20, 1985
	50MG/VIAL	N050467 002	

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-131(of 375)

\*\* See List Footnote

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

SANDOZ INC	150MG/VIAL 2MG/ML	N050467 004 Jul 22, 1987 A200146 001 Jul 18, 2012
RUBEX		
BRISTOL MYERS SQUIBB	10MG/VIAL 50MG/VIAL 100MG/VIAL	A062926 001 Apr 13, 1989 A062926 002 Apr 13, 1989 A062926 003 Apr 13, 1989

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

PAR PHARM	EQ 75MG BASE	A065055 004 Apr 18, 2005
SANDOZ INC	EQ 50MG BASE	A065032 001 Jun 30, 2000
	EQ 100MG BASE	A065032 002 Jun 30, 2000
WATSON LABS	EQ 50MG BASE	A065041 001 Apr 28, 2000
	EQ 100MG BASE	A065041 002 Apr 28, 2000

FOR SUSPENSION; ORAL

DOXYCHEL

RACHELLE	EQ 25MG BASE/5ML	A061720 001
----------	------------------	-------------

TABLET; ORAL

DOXYCYCLINE

SANDOZ INC	EQ 50MG BASE	A065353 001 Nov 27, 2006
	EQ 75MG BASE	A065353 002 Nov 27, 2006
	EQ 100MG BASE	A065353 003 Nov 27, 2006
SUN PHARM INDUSTRIES	EQ 50MG BASE	A065471 001 Apr 17, 2009
	EQ 75MG BASE	A065471 002 Apr 17, 2009
	EQ 100MG BASE	A065471 003 Apr 17, 2009

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXY-LEMMON

TEVA	EQ 50MG BASE	A062497 001 Aug 23, 1984
	EQ 100MG BASE	A062497 002 Jun 15, 1984

DOXYCYCLINE HYCLATE

HALSEY	EQ 50MG BASE	A062119 002 May 24, 1985
	EQ 100MG BASE	A062119 001 May 24, 1985
HEATHER	EQ 50MG BASE	A062463 001 Dec 07, 1983
	EQ 100MG BASE	A062463 002 Dec 07, 1983
INTERPHARM	EQ 50MG BASE	A062763 001 Sep 02, 1988
	EQ 100MG BASE	A062763 002 Sep 02, 1988
MUTUAL PHARM	EQ 50MG BASE	A062418 001 Jan 28, 1983
	EQ 100MG BASE	A062418 002 Jan 28, 1983
PAR PHARM	EQ 50MG BASE	A062434 001 Oct 19, 1984
	EQ 100MG BASE	A062442 001 Dec 22, 1983
PVT FORM	EQ 50MG BASE	A062631 001 Jul 24, 1986
	EQ 100MG BASE	A062631 002 Jul 24, 1986
RANBAXY	EQ 50MG BASE	A062479 001 Dec 23, 1983
	EQ 100MG BASE	A062479 002 Dec 23, 1983
SUPERPHARM	EQ 50MG BASE	A062469 001 Oct 31, 1984
	EQ 100MG BASE	A062469 002 Oct 31, 1984
WARNER CHILCOTT	EQ 50MG BASE	A062594 001 Dec 05, 1985
	EQ 100MG BASE	A062594 002 Dec 05, 1985
WATSON LABS	EQ 50MG BASE	A061717 001
	EQ 50MG BASE	A062142 001
	EQ 100MG BASE	A061717 002
	EQ 100MG BASE	A062142 002

PERIOSTAT

+ COLLAGENEX	EQ 20MG BASE **	N050744 001 Sep 30, 1998
--------------	-----------------	--------------------------

VIBRAMYCIN

+ PFIZER	EQ 50MG BASE **	N050007 001
----------	-----------------	-------------

CAPSULE, COATED PELLETS; ORAL

DOXYCYCLINE HYCLATE

PLIVA	EQ 100MG BASE	A063187 001 Jun 30, 1992
-------	---------------	--------------------------

CAPSULE, DELAYED RELEASE; ORAL

DORYX

+ MAYNE PHARMA INTL	EQ 75MG BASE	N050582 002 Aug 13, 2001
+ MAYNE PHARMA INTL	EQ 100MG BASE	N050582 001 Jul 22, 1985

WARNER CHILCOTT	EQ 100MG BASE	A062653 001 Oct 30, 1985
-----------------	---------------	--------------------------



**DISCONTINUED DRUG PRODUCT LIST**

6-133(of 375)

\*\* See List Footnote

**DRONABINOL**CAPSULE;ORAL  
DRONABINOL

INSYS THERAP	2.5MG	A078501 001 Aug 19, 2011
	5MG	A078501 002 Aug 19, 2011
	10MG	A078501 003 Aug 19, 2011

**DROPERIDOL**

INJECTABLE;INJECTION

DROPERIDOL

ABRAXIS PHARM	2.5MG/ML	A070992 001 Nov 17, 1986
	2.5MG/ML	A070993 001 Nov 17, 1986
ASTRAZENECA	2.5MG/ML	A072018 001 Oct 20, 1988
HOSPIRA	2.5MG/ML	A071645 001 Apr 07, 1988
	2.5MG/ML	A072272 001 Aug 31, 1995
IGI LABS INC	2.5MG/ML	A072019 001 Oct 19, 1988
	2.5MG/ML	A072020 001 Oct 19, 1988
	2.5MG/ML	A072021 001 Oct 19, 1988
LUITPOLD	2.5MG/ML	A072335 001 Oct 24, 1988
SMITH AND NEPHEW	2.5MG/ML	A071750 001 Sep 06, 1988
SOLOPAK	2.5MG/ML	A071754 001 Sep 06, 1988
	2.5MG/ML	A071755 001 Sep 06, 1988
WATSON LABS	2.5MG/ML	A073520 001 Nov 27, 1991
	2.5MG/ML	A073521 001 Nov 27, 1991
	2.5MG/ML	A073523 001 Nov 27, 1991

**DROPERIDOL; FENTANYL CITRATE**

INJECTABLE;INJECTION

FENTANYL CITRATE AND DROPERIDOL

ASTRAZENECA	2.5MG/ML;EQ 0.05MG BASE/ML	A072026 001 Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	A072027 001 Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	A072028 001 Apr 13, 1989
HOSPIRA	2.5MG/ML;EQ 0.05MG BASE/ML	A071982 001 May 04, 1988
INNOVAR		

AKORN MFG 2.5MG/ML;EQ 0.05MG BASE/ML N016049 001

**DYCLONINE HYDROCHLORIDE**

SOLUTION;TOPICAL

DYCLONE

+	ASTRAZENECA	0.5% **	N009925 002
+		1% **	N009925 001

**DYDROGESTERONE**

TABLET;ORAL

GYNOREST

SOLVAY	5MG **	N017388 001
	10MG **	N017388 002

**DYPHYLLINE**

ELIXIR;ORAL

NEOTHYLLINE

TEVA 160MG/15ML N007794 003

INJECTABLE;INJECTION

NEOTHYLLINE

TEVA 250MG/ML N009088 001

TABLET;ORAL

DILOR

SAVAGE LABS 200MG A084514 001

DILOR-400

SAVAGE LABS 400MG A084751 001

LUFYLLIN

MYLAN SPECIALITY LP 200MG A084566 001  
400MG A084566 002

NEOTHYLLINE

TEVA 200MG N007794 001  
400MG N007794 002

**DISCONTINUED DRUG PRODUCT LIST**

6-134(of 375)

\*\* See List Footnote

ECHOTHIOPHATE IODIDE

FOR SOLUTION;OPHTHALMIC		
PHOSPHOLINE IODIDE		
WYETH PHARMS INC	0.03%	N011963 002
	0.06%	N011963 004
	0.25%	N011963 003

EDETA TE CALCIUM DISODIUM

TABLET;ORAL		
CALCIUM DISODIUM VERSENATE		
MEDICIS	500MG	N008922 002

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION		
EDROPHONIUM CHLORIDE		
HOSPIRA	10MG/ML	A040131 001 Feb 24, 1998
WATSON LABS	10MG/ML	A040044 001 Mar 20, 1996
EDROPHONIUM CHLORIDE PRESERVATIVE FREE		
WATSON LABS	10MG/ML	A040043 001 Mar 20, 1996
REVERSOL		
ORGANON USA INC	10MG/ML	A089624 001 May 13, 1988
TENSILON		
+ TELIGENT	10MG/ML **	N007959 001
TENSILON PRESERVATIVE FREE		
+ TELIGENT	10MG/ML **	N007959 002

EFAVIRENZ

CAPSULE;ORAL		
SUSTIVA		
+ BRISTOL MYERS SQUIBB	100MG **	N020972 002 Sep 17, 1998
TABLET;ORAL		
SUSTIVA		
+ BRISTOL MYERS SQUIBB	300MG **	N021360 001 Feb 01, 2002

EFLORNITHINE HYDROCHLORIDE

INJECTABLE; INJECTION		
ORNIDYL		
SANOFI AVENTIS US	200MG/ML	N019879 002 Nov 28, 1990

ELVITEGRAVIR

TABLET;ORAL		
VITEKTA		
+ GILEAD SCIENCES INC	85MG	N203093 001 Sep 24, 2014
+	150MG	N203093 002 Sep 24, 2014

ENALAPRIL MALEATE

TABLET;ORAL		
ENALAPRIL MALEATE		
APOTHECON	2.5MG	A075583 001 Aug 22, 2000
	5MG	A075583 002 Aug 22, 2000
	10MG	A075583 003 Aug 22, 2000
	20MG	A075583 004 Aug 22, 2000
IVAX SUB TEVA PHARMS	2.5MG	A075482 001 Aug 22, 2000
	5MG	A075482 002 Aug 22, 2000
	10MG	A075482 003 Aug 22, 2000
	20MG	A075482 004 Aug 22, 2000
KRKA DD NOVO MESTO	2.5MG	A075370 001 Aug 22, 2000
	5MG	A075370 002 Aug 22, 2000
	10MG	A075369 001 Aug 22, 2000
	20MG	A075369 002 Aug 22, 2000
MYLAN	2.5MG	A075472 001 Aug 22, 2000
	5MG	A075472 002 Aug 22, 2000
	10MG	A075472 003 Aug 22, 2000
	20MG	A075472 004 Aug 22, 2000
SANDOZ	2.5MG	A075048 001 Aug 22, 2000
	5MG	A075048 002 Aug 22, 2000
	10MG	A075048 003 Aug 22, 2000
	20MG	A075048 004 Aug 22, 2000
SANDOZ INC	2.5MG	A075621 001 Aug 22, 2000
	5MG	A075621 002 Aug 22, 2000
	10MG	A075621 003 Aug 22, 2000
	20MG	A075621 004 Aug 22, 2000

**DISCONTINUED DRUG PRODUCT LIST**

6-135(of 375)

\*\* See List Footnote

ENALAPRIL MALEATE

TABLET;ORAL

## ENALAPRIL MALEATE

SUN PHARM INDS LTD	2.5MG 5MG 10MG 20MG	A075556 001 Aug 22, 2000 A075556 002 Aug 22, 2000 A075556 003 Aug 22, 2000 A075556 004 Aug 22, 2000
WATSON LABS	2.5MG 5MG 10MG 20MG	A075501 001 Aug 22, 2000 A075501 002 Aug 22, 2000 A075501 003 Aug 22, 2000 A075501 004 Aug 22, 2000

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE;ORAL

## LEXXEL

ASTRAZENECA	5MG;2.5MG 5MG;5MG	N020668 002 Oct 28, 1998 N020668 001 Dec 27, 1996
-------------	----------------------	--

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

## ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

IVAX SUB TEVA PHARMS	5MG;12.5MG 10MG;25MG	A075736 001 Mar 25, 2003 A075736 002 Mar 25, 2003
UPSHER-SMITH LABS	5MG;12.5MG 10MG;25MG	A076116 001 Sep 19, 2001 A076116 002 Sep 19, 2001

ENALAPRILAT

INJECTABLE;INJECTION

## ENALAPRILAT

HOSPIRA	1.25MG/ML 1.25MG/ML	A075456 001 Aug 22, 2000 A075571 001 Aug 22, 2000
VASOTEC + BIOVAIL LABS INTL	1.25MG/ML	N019309 001 Feb 09, 1988

ENFLURANE

LIQUID;INHALATION

## ENFLURANE

ABBOTT	99.9%	A070803 001 Sep 08, 1987
PIRAMAL CRITICAL	99.9%	A074396 001 Jul 29, 1994
ETHRANE BAXTER HLTHCARE	99.9%	N017087 001

ENOXACIN

TABLET;ORAL

## PENETREX

SANOFI AVENTIS US	200MG 400MG	N019616 004 Dec 31, 1991 N019616 005 Dec 31, 1991
-------------------	----------------	--

ENOXAPARIN SODIUM

INJECTABLE;SUBCUTANEOUS

## LOVENOX (PRESERVATIVE FREE)

+ SANOFI AVENTIS US	90MG/0.6ML (150MG/ML) **	N020164 006 Jun 02, 2000
---------------------	--------------------------	--------------------------

ENTACAPONE

TABLET;ORAL

## ENTACAPONE

MYLAN PHARMS INC	200MG	A202394 001 May 13, 2013
------------------	-------	--------------------------

EPINEPHRINE

AEROSOL, METERED;INHALATION

## BRONKAID MIST

STERLING	0.25MG/INH	N016803 001
----------	------------	-------------

## EPINEPHRINE

ARMSTRONG PHARMS	0.2MG/INH	A087907 001 May 23, 1984
------------------	-----------	--------------------------

## PRIMATENE MIST

WYETH CONS	0.2MG/INH	N016126 001
------------	-----------	-------------

INJECTABLE;INJECTION

## SUS-PHRINE SULFITE FREE

FOREST LABS	1.5MG/AMP 5MG/ML	N007942 003 Feb 05, 1999 N007942 001
-------------	---------------------	---

INJECTABLE;INTRAMUSCULAR

## EPI E Z PEN JR

MYLAN SPECIALITY LP	0.15MG/DELIVERY	N019430 004 Aug 03, 1995
---------------------	-----------------	--------------------------

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-136(of 375)

\*\* See List Footnote

EPINEPHRINE

INJECTABLE; INTRAMUSCULAR

EPIPEN E Z PEN

MYLAN SPECIALITY LP 0.3MG/DELIVERY

N019430 003 Aug 03, 1995

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

TWINJECT 0.15

IMPAK LABS INC EQ 0.15MG/DELIVERY

N020800 002 May 28, 2004

TWINJECT 0.3

IMPAK LABS INC EQ 0.3MG/DELIVERY

N020800 001 May 30, 2003

EPINEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

BRONITIN MIST

WYETH CONS 0.3MG/INH

N016126 002

MEDIHALER-EPI

3M 0.3MG/INH

N010374 003

EPINEPHRINE BITARTRATE; ETIDOCAINe HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA 0.005MG/ML;1% \*\*  
+ 0.005MG/ML;1.5% \*\*  
+ DENTSPLY PHARM 0.005MG/ML;1.5% \*\*N017751 006  
N017751 007  
N021384 001EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANESE FORTE

ASTRAZENECA 0.005MG/ML;4%

N014763 008

EPINEPHRINE; ETIDOCAINe HYDROCHLORIDE

INJECTABLE; INJECTION

DURANEST

+ ASTRAZENECA 0.005MG/ML;0.5% \*\*

N017751 004

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINe HYDROCHLORIDE W/ EPINEPHRINE  
CARLISLE 0.01MG/ML;2%  
0.02MG/ML;2%A084720 001  
A084732 001LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE  
BELMORA LLC 0.01MG/ML;2%  
0.02MG/ML;2%A080504 004 Oct 19, 1983  
A080504 005 Oct 19, 1983HOSPIRA 0.005MG/ML;1%  
0.005MG/ML;1.5%A089649 001 Jun 21, 1988  
A089650 001 Jun 21, 1988WEST-WARD PHARMS INT 0.01MG/ML;1%  
0.01MG/ML;2%

A080406 001

A080406 002

LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE  
ABBOTT 0.01MG/ML;1%  
BEL MAR 0.01MG/ML;1%

A083154 001

DELL LABS 0.01MG/ML;1%  
0.01MG/ML;2%

A080820 001

INTL MEDICATION 0.01MG/ML;1%  
WATSON LABS 0.01MG/ML;1%

A080757 001

0.01MG/ML;1%  
0.01MG/ML;2%

A083389 001

0.01MG/ML;1%  
WATSON LABS 0.01MG/ML;1%

A083390 001

0.01MG/ML;1%  
0.01MG/ML;2%

A086402 001

LIDOCATON  
PHARMATON 0.01MG/ML;2%  
0.02MG/ML;2%

A080377 003

A085463 001

A080377 004

XYLOCAINE DENTAL WITH EPINEPHRINE  
DENTSPLY PHARM 0.01MG/ML;2%  
0.02MG/ML;2%

A084729 001 Aug 17, 1983

A084728 001 Aug 17, 1983

XYLOCAINE W/ EPINEPHRINE  
ASTRAZENECA 0.005MG/ML;1%  
0.005MG/ML;1.5%

N010418 006

N010418 010

0.005MG/ML;2%  
FRESENIUS KABI USA 0.01MG/ML;2%

N010418 008

N006488 003

PATCH; IONTOPHORESIS, TOPICAL  
LIDOSITE TOPICAL SYSTEM KIT  
VYTERIS 1.05MG/PATCH;100MG/PATCH

N021504 001 May 06, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-137(of 375)

\*\* See List Footnote

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

SOLUTION;IONTOPHORESIS		
IONTOCAINE		
IOMED	0.01MG/ML;2%	N020530 001 Dec 21, 1995
SOLUTION;IONTOPHORESIS, TOPICAL		
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE		
EMPI	0.01MG/ML;2%	N021486 001 Oct 26, 2004

EPINEPHRINE; PROCAINE HYDROCHLORIDE

INJECTABLE;INJECTION		
PROCAINE HYDROCHLORIDE W/ EPINEPHRINE		
BEL MAR	0.02MG/ML;1%	A080758 001
	0.02MG/ML;2%	A080759 001

EPIRUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION		
EPIRUBICIN HYDROCHLORIDE		
EBEWE PHARMA	50MG/25ML (2MG/ML)	A065339 001 Dec 22, 2009
	200MG/100ML (2MG/ML)	A065339 002 Dec 22, 2009
HOSPIRA	50MG/25ML (2MG/ML)	A065343 002 Apr 19, 2007
MUSTAFA NEVSAT	50MG/25ML (2MG/ML)	A090266 001 Apr 15, 2011
	200MG/100ML (2MG/ML)	A090266 002 Apr 15, 2011
MYLAN INSTITUTIONAL	50MG/25ML (2MG/ML)	A065371 001 Nov 28, 2007
	200MG/100ML (2MG/ML)	A065371 002 Nov 28, 2007
INJECTABLE;IV (INFUSION)		
EPIRUBICIN HYDROCHLORIDE		
HOSPIRA	50MG/VIAL	N050807 001 Sep 15, 2006
	200MG/VIAL	N050807 002 Sep 15, 2006

EPLERENONE

TABLET;ORAL		
INSPRA		
GD SEARLE LLC	100MG	N021437 003 Sep 27, 2002

ERPOSARTAN MESYLATE

TABLET;ORAL		
TEVETEN		
ABBVIE	EQ 300MG BASE	N020738 004 Dec 22, 1997

ERPOSARTAN MESYLATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL		
TEVETEN HCT		
ABBVIE	600MG;12.5MG	N021268 001 Nov 01, 2001
	600MG;25MG	N021268 002 Nov 01, 2001

EPTIFIBATIDE

INJECTABLE;INJECTION		
EPTIFIBATIDE		
TEVA PHARMS USA	75MG/100ML	A091555 001 Jun 05, 2015

ERGOCALCIFEROL

CAPSULE;ORAL		
DELTALIN		
LILLY	50,000 IU	A080884 001
VITAMIN D		
CHASE CHEM	50,000 IU	A080747 001
EVERYLIFE	50,000 IU	A080956 001
IMPAK LABS	50,000 IU	A080951 001
LANNETT	50,000 IU	A080825 001
VITARINE	50,000 IU	A084053 001
WEST WARD	50,000 IU	A083102 001

ERGOLOID MESYLATES

CAPSULE;ORAL		
HYDERGINE LC		
NOVARTIS	1MG	N018706 001 Jan 18, 1983
SOLUTION;ORAL		
HYDERGINE		
NOVARTIS	1MG/ML	N018418 001
TABLET;ORAL		
ERGOLOID MESYLATES		
MUTUAL PHARM	1MG	A088891 001 Nov 01, 1985
	1MG	A086433 001 May 27, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-138(of 375)

\*\* See List Footnote

ERGOLOID MESYLATES

TABLET;ORAL

ERGOLOID MESYLATES

1MG A087244 001 Aug 16, 1982

GERIMAL

WATSON LABS

1MG A088207 001 Mar 22, 1984

HYDERGINE

NOVARTIS

0.5MG N017993 003

+

1MG N017993 001

TABLET;SUBLINGUAL

ALKERGOT

SANDOZ

0.5MG A085153 001

1MG A087417 001

CIRCANOL

3M

0.5MG A084868 001

1MG A085809 001

DEAPRIL-ST

BRISTOL MYERS SQUIBB

1MG A085020 002

ERGOLOID MESYLATES

KV PHARM

0.5MG A085899 001

0.5MG A086265 001

1MG A085900 001

1MG A086264 001

LEDERLE

0.5MG A086984 001

1MG A086985 001

SUN PHARM INDUSTRIES 0.5MG A087407 001

1MG A087552 001

SUPERPHARM 0.5MG A089233 001 Sep 23, 1986

1MG A089234 001 Sep 23, 1986

VANGARD 0.5MG A088013 001 Sep 20, 1982

1MG A088014 001 Sep 20, 1982

WATSON LABS 0.5MG A084930 001

0.5MG A087233 001

1MG A085177 001

1MG A087183 001

GERIMAL

WATSON LABS

0.5MG A086189 001

1MG A086188 001

HYDERGINE

NOVARTIS

0.5MG N009087 002

1MG N009087 001

HYDROGENATED ERGOT ALKALOIDS

IVAX PHARMS

0.5MG A087186 001

1MG A087185 001

ERGOTAMINE TARTRATE

AEROSOL, METERED;INHALATION

MEDIHALER ERGOTAMINE

3M 0.36MG/INH N012102 001

TABLET;SUBLINGUAL

ERGOSTAT

WATSON LABS INC 2MG A088337 001 Jun 08, 1984

WIGRETTES

ORGANON USA INC 2MG A086750 001 Jul 29, 1982

ERLOTINIB HYDROCHLORIDE

TABLET;ORAL

ERLOTINIB HYDROCHLORIDE

MYLAN PHARMS INC EQ 25MG BASE A091002 001 Jun 11, 2014

EQ 100MG BASE A091002 002 Jun 11, 2014

EQ 150MG BASE A091002 003 Jun 11, 2014

TEVA PHARMS USA EQ 100MG BASE A091059 002 Aug 28, 2015

EQ 150MG BASE A091059 003 Aug 28, 2015

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS;ORAL

ERYC

PARKE DAVIS 250MG A062546 001 Jul 25, 1985

250MG A062618 001 Sep 25, 1985

WARNER CHILCOTT LLC 250MG A062338 001

**DISCONTINUED DRUG PRODUCT LIST**

6-139(of 375)

\*\* See List Footnote

**ERYTHROMYCIN**

CAPSULE, DELAYED REL PELLETS;ORAL			
ERYC 125			
PARKE DAVIS	125MG	A062648	001 Oct 24, 1985
ERYC SPRINKLES			
HOSPIRA	125MG	N050593	001 Jul 22, 1985
ERYTHROMYCIN			
BARR	250MG	A063098	001 May 04, 1989
GEL;TOPICAL			
E-GLADES			
MYLAN PHARMS INC	2%	A065009	001 Mar 18, 2002
EMGEL			
ALTANA	2%	A063107	001 Aug 23, 1991
LOTION;TOPICAL			
E-SOLVE 2			
SYOSSET	2%	A062467	001 Jul 03, 1985
OINTMENT;OPHTHALMIC			
ERYTHROMYCIN			
PHARMADERM	5MG/GM	A062446	001 Sep 26, 1983
PHARMAFAIR	5MG/GM	A062481	001 Apr 05, 1984
ILOTYCIN			
DISTA	0.5%	N050368	001
OINTMENT;TOPICAL			
AKNE-MYCIN			
+ DOW PHARM	2%	N050584	001 Jan 10, 1985
POWDER;FOR RX COMPOUNDING			
ERYTHROMYCIN			
PADDOCK LLC	100%	N050610	001 Nov 07, 1986
SOLUTION;TOPICAL			
A/T/S			
TARO	2%	A062405	001 Nov 18, 1982
C-SOLVE-2			
FOUGERA PHARMS	2%	A062468	001 Jul 03, 1985
ERYDERM			
ARBOR PHARMS INC	2%	A062290	001
ERYMAX			
MERZ PHARMS	2%	A062508	002 Jul 11, 1985
ERYTHRA-DERM			
SAPTALIS PHARMS	2%	A062687	001 Feb 05, 1988
ERYTHRO-STATIN			
HI TECH PHARMA	2%	A064101	001 Oct 22, 1996
ERYTHROMYCIN			
ALPHARMA US PHARMS	1.5%	A062328	001 Apr 19, 1982
	2%	A062326	001 Apr 19, 1982
	2%	A062327	001 Apr 19, 1982
	2%	A062342	001 Feb 25, 1982
	2%	A062957	001 Jul 21, 1988
BAUSCH AND LOMB	2%	A064039	001 Jan 27, 1994
FOUGERA PHARMS	2%	A064187	001 Sep 30, 1997
LILLY	2%	N050532	001
PHARMAFAIR	1.5%	A062485	001 Jul 11, 1984
	2%	A062616	001 Jul 25, 1985
RENAISSANCE PHARMA	2%	A064127	001 Feb 14, 1997
SANSAC			
DOW PHARM	2%	A062522	001 Jan 24, 1985
STATICIN			
+ WESTWOOD SQUIBB	1.5% **	N050526	001
T-STAT			
WESTWOOD SQUIBB	2% **	A062436	001 Mar 09, 1983
SWAB;TOPICAL			
C-SOLVE-2			
IVAX SUB TEVA PHARMS	2%	A062751	001 Jul 30, 1993
ERYCETTE			
+ JOHNSON AND JOHNSON	2% **	N050594	001 Feb 15, 1985
ERYTHROMYCIN			
FOUGERA PHARMS	2%	A065320	001 Jul 25, 2006
MYLAN PHARMS INC	2%	A064128	001 Jul 03, 1996
T-STAT			
WESTWOOD SQUIBB	2%	A062748	001 Jul 23, 1987

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-140(of 375)

\*\* See List Footnote

**ERYTHROMYCIN**

TABLET, DELAYED RELEASE;ORAL

E-BASE

BARR	333MG	A063028 001 May 15, 1990
	333MG	A063086 001 May 15, 1990
	500MG	A062999 001 Nov 25, 1988

E-MYCIN

ARBOR PHARMS INC	250MG	A060272 001
	333MG	A060272 002

ILOTYCIN

DISTA	250MG	A061910 001
-------	-------	-------------

R-P MYCIN

SOLVAY	250MG	A061659 001
--------	-------	-------------

ROBIMYCIN

ROBINS AH	250MG	A061633 001
-----------	-------	-------------

**ERYTHROMYCIN ESTOLATE**

CAPSULE;ORAL

ERYTHROMYCIN ESTOLATE

BARR	EQ 125MG BASE	A062162 001
	EQ 250MG BASE	A062162 002
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A062237 001
WATSON LABS	EQ 250MG BASE	A062087 001
ILOSONE		

LILLY	EQ 125MG BASE	A061897 001
	EQ 250MG BASE	A061897 002

FOR SUSPENSION;ORAL

ILOSONE

DISTA	EQ 125MG BASE/5ML	A061893 001
-------	-------------------	-------------

SUSPENSION;ORAL

ERYTHROMYCIN ESTOLATE

ALPHARMA US PHARMS	EQ 125MG BASE/5ML	A062353 001 Nov 18, 1982
	EQ 250MG BASE/5ML	A062409 001 Dec 16, 1982
G AND W LABS INC	EQ 125MG BASE/5ML	A062169 001 Oct 17, 1990
	EQ 250MG BASE/5ML	A062169 002 Oct 17, 1990
LIFE LABS	EQ 250MG BASE/5ML	A062362 001 Dec 17, 1982
ILOSONE		

LILLY	EQ 125MG BASE/5ML	A061894 001
	EQ 125MG BASE/5ML	N050010 001
	EQ 250MG BASE/5ML	A061894 002
	EQ 250MG BASE/5ML	N050010 002

SUSPENSION/DROPS;ORAL

ILOSONE

LILLY	EQ 100MG BASE/ML	A061894 003
-------	------------------	-------------

TABLET;ORAL

ILOSONE

LILLY	EQ 500MG BASE	A061896 001
-------	---------------	-------------

TABLET, CHEWABLE;ORAL

ILOSONE

DISTA	EQ 125MG BASE	A061895 001
	EQ 250MG BASE	A061895 002

**ERYTHROMYCIN ESTOLATE; SULFISOXAZOLE ACETYL**

SUSPENSION;ORAL

ILOSONE SULFA

LILLY	EQ 125MG BASE/5ML;EQ 600MG BASE/5ML	N050599 001 Sep 29, 1989
-------	-------------------------------------	--------------------------

**ERYTHROMYCIN ETHYLSUCCINATE**

GRANULE;ORAL

ERYTHROMYCIN ETHYLSUCCINATE

ANI PHARMS INC	EQ 200MG BASE/5ML	A062055 001
----------------	-------------------	-------------

PEDIAMYCIN

ROSS LABS	EQ 200MG BASE/5ML	A062305 001
-----------	-------------------	-------------

SUSPENSION;ORAL

E-MYCIN E

PHARMACIA AND UPJOHN	EQ 200MG BASE/5ML	A062198 001
	EQ 400MG BASE/5ML	A062198 002

E.E.S. 200

ARBOR PHARMS LLC	EQ 200MG BASE/5ML **	A061639 001
------------------	----------------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-141(of 375)

\*\* See List Footnote

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION;ORAL

E.E.S. 400

ARBOR PHARMS LLC	EQ 400MG BASE/5ML **	A061639 002
ERYTHROMYCIN ETHYLSUCCINATE		
ALPHARMA US PHARMS	EQ 200MG BASE/5ML	A062200 001
	EQ 400MG BASE/5ML	A062200 002
DISTA	EQ 200MG BASE/5ML	A062177 001
	EQ 400MG BASE/5ML	A062177 002
NASKA	EQ 400MG BASE/5ML	A062674 001 Mar 10, 1987
PARKE DAVIS	EQ 200MG BASE/5ML	A062231 001
	EQ 400MG BASE/5ML	A062231 002
PHARMAFAIR	EQ 200MG BASE/5ML	A062559 001 Mar 15, 1985
	EQ 400MG BASE/5ML	A062558 001 Mar 15, 1985
PEDIAMYCIN		
ARBOR PHARMS LLC	EQ 200MG BASE/5ML	A062304 001
PEDIAMYCIN 400		
ARBOR PHARMS LLC	EQ 400MG BASE/5ML	A062304 002
WYAMYCIN E		
WYETH AYERST	EQ 200MG BASE/5ML	A062123 002
	EQ 400MG BASE/5ML	A062123 001

SUSPENSION/DROPS;ORAL

PEDIAMYCIN

ROSS LABS	EQ 100MG BASE/2.5ML	A062305 002
-----------	---------------------	-------------

TABLET;ORAL

E.E.S. 400

ARBOR PHARMS LLC	EQ 400MG BASE	A061905 001
ERYTHROMYCIN ETHYLSUCCINATE		
BARR	EQ 400MG BASE	A062256 001
MYLAN	EQ 400MG BASE	A062847 001 Sep 14, 1988
TABLET, CHEWABLE;ORAL		
E.E.S.		
ARBOR PHARMS INC	EQ 200MG BASE	N050297 002
ERYPED		
ARBOR PHARMS INC	EQ 200MG BASE	N050297 003 Jul 05, 1988
PEDIAMYCIN		
ROSS LABS	EQ 200MG BASE	A062306 001

ERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYL

GRANULE;ORAL

ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL		
BARR	EQ 200MG BASE/5ML;EQ 600MG BASE/5ML	A062759 001 May 20, 1988
ERYZOLE		
ALRA	EQ 200MG BASE/5ML;EQ 600MG BASE/5ML	A062758 001 Jun 15, 1988
PEDIAZOLE		
ROSS LABS	EQ 200MG BASE/5ML;EQ 600MG BASE/5ML	N050529 001

ERYTHROMYCIN GLUCEPTATE

INJECTABLE;INJECTION

ILOTYCIN GLUCEPTATE

DISTA	EQ 250MG BASE/VIAL	N050370 001
	EQ 500MG BASE/VIAL	N050370 002
	EQ 1GM BASE/VIAL	N050370 003

ERYTHROMYCIN LACTOBIONATE

INJECTABLE;INJECTION

ERYTHROCIN

ABBOTT	EQ 500MG BASE/VIAL	A062586 001 Jan 04, 1988
	EQ 1GM BASE/VIAL	A062586 002 Jan 04, 1988
HOSPIRA	EQ 500MG BASE/VIAL	N050182 002
	EQ 1GM BASE/VIAL	N050182 003
	EQ 1GM BASE/VIAL	N050609 002 Sep 24, 1986

ERYTHROMYCIN

ELKINS SINK	EQ 500MG BASE/VIAL	A062563 001 Mar 28, 1985
	EQ 1GM BASE/VIAL	A062563 002 Mar 28, 1985

ERYTHROMYCIN LACTOBIONATE

ABRAXIS PHARM	EQ 500MG BASE/VIAL	A062604 001 Nov 24, 1986
	EQ 1GM BASE/VIAL	A062604 002 Nov 24, 1986
BAXTER HLTHCARE	EQ 500MG BASE/VIAL	A062993 001 May 09, 1989
	EQ 1GM BASE/VIAL	A062993 002 May 09, 1989
TEVA PARENTERAL	EQ 500MG BASE/VIAL	A063253 001 Jul 30, 1993

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-142(of 375)

\*\* See List Footnote

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROMYCIN LACTOBIONATE

EQ 1GM BASE/VIAL

A063253 002 Jul 30, 1993

ERYTHROMYCIN STEARATE

TABLET; ORAL

BRISTAMYCIN

BRISTOL

EQ 250MG BASE

A061304 001

EQ 250MG BASE

A061887 001

ERYPAR

PARKE DAVIS

EQ 250MG BASE

A062032 001

EQ 500MG BASE

A062032 002

WARNER CHILCOTT

EQ 250MG BASE

A062322 001

ERYTHROCIN STEARATE

ARBOR PHARMS LLC

EQ 125MG BASE

A060359 002

EQ 500MG BASE

A060359 003

ERYTHROMYCIN STEARATE

ANI PHARMS INC

EQ 250MG BASE

A061461 001

EQ 250MG BASE

A061591 001

EQ 500MG BASE

A061461 002

EQ 500MG BASE

A063179 001 May 15, 1990

LEDERLE

EQ 250MG BASE

A062089 001

EQ 500MG BASE

A062089 002

MYLAN

EQ 250MG BASE

A061505 001

EQ 500MG BASE

A061505 002

PUREPAC PHARM

EQ 250MG BASE

A061743 001

WATSON LABS

EQ 250MG BASE

A062121 002

EQ 500MG BASE

A062121 001

ETHRIL 250

BRISTOL MYERS SQUIBB

EQ 250MG BASE

A061605 001

ETHRIL 500

BRISTOL MYERS SQUIBB

EQ 500MG BASE

A061605 002

PFIZER-E

PFIZER

EQ 250MG BASE

A061791 001

EQ 500MG BASE

A061791 002

WYAMYCIN S

WYETH AYERST

EQ 250MG BASE

A061675 001

EQ 500MG BASE

A061675 002

ESCITALOPRAM OXALATE

CAPSULE; ORAL

ESCITALOPRAM OXALATE

MYLAN PHARMS INC

EQ 5MG BASE

A077660 001 Jul 31, 2007

EQ 10MG BASE

A077660 002 Jul 31, 2007

EQ 20MG BASE

A077660 003 Jul 31, 2007

TABLET; ORAL

ESCITALOPRAM OXALATE

MYLAN PHARMS INC

EQ 5MG BASE

A077550 001 May 14, 2015

EQ 10MG BASE

A077550 002 May 14, 2015

EQ 20MG BASE

A077550 003 May 14, 2015

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

BAXTER HLTHCARE

10MG/ML

N019386 003 Aug 15, 1988

20MG/ML

N019386 007 May 28, 2003

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

ESOMEPRAZOLE SODIUM

AUROBINDO PHARMA LTD

EQ 20MG BASE/VIAL

A204657 001 Aug 10, 2016

MYLAN LABS LTD

EQ 20MG BASE/VIAL

A202686 001 May 17, 2017

SUN PHARMA GLOBAL

EQ 20MG BASE/VIAL

A200882 001 Mar 18, 2013

NEXIUM IV

+ ASTRAZENECA PHARMS

EQ 20MG BASE/VIAL

N021689 001 Mar 31, 2005

**DISCONTINUED DRUG PRODUCT LIST**

6-143(of 375)

\*\* See List Footnote

ESOMEPRAZOLE STRONTIUM

CAPSULE, DELAYED RELEASE;ORAL

ESOMEPRAZOLE STRONTIUM

+ R2 PHARMA LLC 24.65MG

N202342 001 Aug 06, 2013

ESTAZOLAM

TABLET;ORAL

PROSOM

+ ABBOTT 1MG \*\*  
+ 2MG \*\*N019080 001 Dec 26, 1990  
N019080 002 Dec 26, 1990ESTRADIOL

FILM, EXTENDED RELEASE;TRANSDERMAL

ESCLIM

WOMEN FIRST HLTHCARE 0.025MG/24HR  
0.0375MG/24HR  
0.05MG/24HR  
0.075MG/24HR  
0.1MG/24HRN020847 001 Aug 04, 1998  
N020847 002 Aug 04, 1998  
N020847 003 Aug 04, 1998  
N020847 004 Aug 04, 1998  
N020847 005 Aug 04, 1998

ESTRADERM

+ NOVARTIS 0.05MG/24HR  
+ 0.1MG/24HRN019081 002 Sep 10, 1986  
N019081 003 Sep 10, 1986

ESTRADIOL

ORTHO MCNEIL PHARM 0.05MG/24HR  
0.075MG/24HR  
0.1MG/24HRN021048 001 Sep 20, 1999  
N021048 002 Sep 20, 1999  
N021048 003 Sep 20, 1999

FEMPATCH

PARKE DAVIS 0.025MG/24HR

N020417 001 Dec 03, 1996

VIVELLE

NOVARTIS 0.025MG/24HR  
0.0375MG/24HR  
0.05MG/24HR  
0.075MG/24HR  
0.1MG/24HRN020323 005 Aug 16, 2000  
N020323 001 Oct 28, 1994  
N020323 002 Oct 28, 1994  
N020323 003 Oct 28, 1994  
N020323 004 Oct 28, 1994

GEL;TOPICAL

ESTROGEL

ASCEND THERAPS US 0.06%

N021166 001 Feb 09, 2004

TABLET;ORAL

ESTRACE

BRISTOL MYERS SQUIBB 0.5MG  
1MG  
2MGA081295 001 Jun 30, 1993  
A084499 001  
A084500 001

ESTRADIOL

LANNETT HOLDINGS INC 0.5MG  
1MG  
2MG  
USL PHARMA 0.5MG  
1MG  
2MGA040138 001 Jan 30, 1998  
A040138 002 Jan 30, 1998  
A040138 003 Jan 30, 1998  
A040297 001 Apr 17, 2002  
A040297 002 Apr 17, 2002  
A040297 003 Apr 17, 2002

GYNODIOL

DURAMED PHARMS BARR 0.5MG  
1MG  
1.5MG  
2MGA040212 001 Dec 29, 1997  
A040212 002 Dec 29, 1997  
A040212 003 Dec 29, 1997  
A040212 004 Dec 29, 1997

INNOFEM

NOVO NORDISK INC 0.5MG  
1MG  
2MGA040312 001 Nov 19, 1999  
A040312 002 Nov 19, 1999  
A040312 003 Nov 19, 1999

TABLET;VAGINAL

VAGIFEM

+ NOVO NORDISK INC 25MCG \*\*

N020908 001 Mar 26, 1999

ESTRADIOL ACETATE

TABLET;ORAL

FEMTRACE

+ APIL 0.45MG  
+ 0.9MG  
+ 1.8MGN021633 001 Aug 20, 2004  
N021633 002 Aug 20, 2004  
N021633 003 Aug 20, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-144(of 375)

\*\* See List Footnote

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

PHARMACIA AND UPJOHN 1MG/ML  
3MG/MLA085470 001  
A085470 002

ESTRADIOL CYPIONATE

WATSON LABS 5MG/ML

A085620 001

ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATE

INJECTABLE; INTRAMUSCULAR

LUNELLE

PHARMACIA AND UPJOHN 5MG/0.5ML;25MG/0.5ML

N020874 001 Oct 05, 2000

ESTRADIOL CYPIONATE; TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTADIOL

PHARMACIA AND UPJOHN 2MG/ML;50MG/ML  
TESTOSTERONE CYPIONATE-ESTRADIOL CYPIONATE  
WATSON LABS 2MG/ML;50MG/MLN017968 001  
A085603 001 Mar 13, 1986ESTRADIOL VALERATE

INJECTABLE; INJECTION

ESTRADIOL VALERATE

SANDOZ INC 10MG/ML  
20MG/ML  
40MG/ML  
WATSON LABS 10MG/ML  
40MG/ML  
WATSON LABS INC 20MG/MLA040628 001 Oct 04, 2007  
A040628 002 Oct 04, 2007  
A040628 003 Oct 04, 2007  
A083546 001  
A083714 001  
A083547 001ESTRADIOL VALERATE; TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DITATE-DS

SAVAGE LABS 8MG/ML;180MG/ML  
TESTOSTERONE ENANTHATE AND ESTRADIOL VALERATE  
WATSON LABS 4MG/ML;90MG/ML  
8MG/ML;180MG/MLA086423 001  
A085865 001  
A085860 001ESTRADIOL; NORGESTIMATE

TABLET;ORAL

PREFEST

+ TEVA WOMENS 1MG,1MG;N/A,0.09MG \*\*

N021040 001 Oct 22, 1999

ESTROGENS, CONJUGATED

TABLET;ORAL

PREMARIN

WYETH PHARMS INC 2.5MG

N004782 002

ESTROGENS, CONJUGATED SYNTHETIC A

CREAM;VAGINAL

SYNTHETIC CONJUGATED ESTROGENS A

TEVA WOMENS 0.625MG/GM

N021788 001 Nov 28, 2008

TABLET;ORAL

CENESTIN

+ TEVA BRANDED PHARM 0.3MG \*\*  
+ 0.45MG \*\*  
+ 0.625MG \*\*  
+ 0.9MG \*\*  
+ 1.25MG \*\*N020992 001 Jun 21, 2002  
N020992 005 Feb 05, 2004  
N020992 002 Mar 24, 1999  
N020992 003 Mar 24, 1999  
N020992 004 Mar 13, 2000ESTROGENS, CONJUGATED SYNTHETIC B

TABLET;ORAL

ENJUVIA

TEVA BRANDED PHARM 0.3MG  
0.45MG  
0.625MG \*\*  
0.9MG  
1.25MG \*\*N021443 001 Dec 20, 2004  
N021443 002 Dec 20, 2004  
N021443 003 May 10, 2004  
N021443 005 Apr 27, 2007  
N021443 004 May 10, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-145(of 375)

\*\* See List Footnote

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET;ORAL-28

PREMPHASE (PREMARIN;CYCRIN 14/14)

WYETH PHARMS INC 0.625MG,0.625MG;N/A,5MG

N020303 002 Dec 30, 1994

PREMPRO (PREMARIN;CYCRIN)

WYETH PHARMS INC 0.625MG,0.625MG;2.5MG,2.5MG

N020303 001 Dec 30, 1994

ESTROGENS, CONJUGATED; MEPROBAMATE

TABLET;ORAL

MILPREM-200

MEDPOINTE PHARM HLC 0.45MG;200MG

N011045 002

MILPREM-400

MEDPOINTE PHARM HLC 0.45MG;400MG

N011045 001

PMB 200

WYETH AYERST 0.45MG;200MG

N010971 005

PMB 400

WYETH AYERST 0.45MG;400MG

N010971 003

ESTROGENS, ESTERIFIED

TABLET;ORAL

AMNESTROGEN

BRISTOL MYERS SQUIBB 0.3MG  
0.625MG  
1.25MG  
2.5MGA083266 001  
A083266 002  
A083266 003  
A083266 004

ESTERIFIED ESTROGENS

PVT FORM 0.625MG  
1.25MG  
2.5MGA083414 001  
A083765 001

SANDOZ 1.25MG

A085907 001  
A085302 001

ESTRATAB

SOLVAY 0.3MG  
0.625MG  
1.25MG  
2.5MGA086715 001  
A083209 001  
A083856 001  
A083857 001

EVEX

ROCHE PALO 0.625MG  
1.25MGA084215 001  
A083376 002

FEMOGEN

PVT FORM 0.625MG  
1.25MG  
2.5MGA085076 001  
A085008 001  
A085007 001ESTRONE

INJECTABLE;INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST 2MG/ML

A083488 001

ESTRONE

WATSON LABS 2MG/ML  
WATSON LABS TEVA 5MG/MLA083397 001  
A085239 001

NATURAL ESTROGENIC SUBSTANCE-ESTRONE

WATSON LABS 2MG/ML

A085237 001 Nov 23, 1982

THEELIN

PARKEDALE 1MG/ML  
2MG/ML  
5MG/MLN003977 001  
N003977 002  
N003977 003ESTROPIPATE

CREAM;VAGINAL

OGEN

PHARMACIA AND UPJOHN 1.5MG/GM

A084710 001

TABLET;ORAL

ESTROPIPATE

BARR 0.75MG

A040135 001 Nov 27, 1996

1.5MG

A040135 002 Nov 27, 1996

3MG

A040135 003 Nov 27, 1996

DURAMED PHARMS BARR 0.75MG

A040296 001 Nov 01, 1999

1.5MG

A040296 002 Nov 01, 1999

3MG

A040296 003 Nov 01, 1999

MYLAN 3MG

A040359 003 Aug 26, 1999

WATSON LABS 0.75MG

A081213 001 Sep 23, 1993

**DISCONTINUED DRUG PRODUCT LIST**

6-146(of 375)

\*\* See List Footnote

ESTROPIPATETABLET;ORAL  
ESTROPIPATE

WATSON LABS TEVA	1.5MG 6MG 3MG	A081214 001 Sep 23, 1993 A081216 001 Sep 23, 1993 A081215 001 Sep 23, 1993
OGEN .625	PHARMACIA AND UPJOHN 0.75MG	A083220 001
OGEN 1.25	PHARMACIA AND UPJOHN 1.5MG	A083220 002
OGEN 2.5	PHARMACIA AND UPJOHN 3MG	A083220 003
ORTHO-EST	SUN PHARM INDS INC 0.75MG 1.5MG	A089567 001 Feb 27, 1991 A089582 001 Jul 17, 1991

ESZOPICLONE

TABLET;ORAL

ESZOPICLONE  
WOCKHARDT LTD

1MG 2MG 3MG	A091165 001 Jul 14, 2011 A091165 002 Jul 14, 2011 A091165 003 Jul 14, 2011
-------------------	--

ETHACRYNIC ACID

TABLET;ORAL

EDECрин  
ATON

50MG	N016092 002
------	-------------

ETHAMBUTOL HYDROCHLORIDE

TABLET;ORAL

MYAMBUTOL  
STI PHARMA LLC

200MG 500MG	N016320 002 N016320 004
----------------	----------------------------

ETHCHLORVYNOL

CAPSULE;ORAL

ETHCHLORVYNOL  
BANNER PHARMACAPS

100MG 200MG 500MG 750MG	A084463 001 A084463 002 A084463 003 A084463 004
----------------------------------	--

PLACIDYL

ABBVIE	100MG 200MG 500MG 750MG	N010021 004 N010021 007 N010021 002 N010021 010
--------	----------------------------------	--

ETHINAMATE

CAPSULE;ORAL

VALMID  
DISTA

500MG	N009750 001
-------	-------------

ETHINYL ESTRADIOL

TABLET;ORAL

ESTINYL  
SCHERING

0.02MG 0.05MG 0.5MG	N005292 001 N005292 002 N005292 003
---------------------------	---

FEMINONE

PHARMACIA AND UPJOHN 0.05MG	N016649 001
-----------------------------	-------------

LYNORAL

ORGANON USA INC 0.01MG 0.05MG	N005490 003 N005490 002
----------------------------------	----------------------------

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET;ORAL-21

DEMULEN 1/35-21

GD SEARLE LLC	0.035MG;1MG **	N018168 001
---------------	----------------	-------------

DEMULEN 1/50-21

GD SEARLE LLC	0.05MG;1MG	N016927 001
---------------	------------	-------------

ZOVIA 1/35E-21

WATSON PHARMS TEVA	0.035MG;1MG	A072720 001 Dec 30, 1991
--------------------	-------------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-147(of 375)

\*\* See List Footnote

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET;ORAL-21			
ZOVIA 1/50E-21			
WATSON LABS	0.05MG;1MG		A072722 001 Dec 30, 1991
TABLET;ORAL-28			
DEMULEN 1/35-28			
GD SEARLE LLC	0.035MG;1MG **		N018160 001
DEMULEN 1/50-28			
GD SEARLE LLC	0.05MG;1MG **		N016936 001

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET;ORAL-28			
NORQUEST FE			
GD SEARLE LLC	0.035MG;75MG;1MG		N018926 001 Jul 18, 1986

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET;ORAL-28			
NORLESTRIN FE 1/50			
PARKE DAVIS	0.05MG;75MG;1MG		N016766 001
NORLESTRIN FE 2.5/50			
PARKE DAVIS	0.05MG;75MG;2.5MG		N016854 001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET;ORAL			
LYBREL			
+ WYETH PHARMS INC	0.02MG;0.09MG **		N021864 001 May 22, 2007
PREVEN EMERGENCY CONTRACEPTIVE KIT			
TEVA BRANDED PHARM	0.05MG;0.25MG		N020946 001 Sep 01, 1998
TABLET;ORAL-21			
ALESSE			
+ WYETH PHARMS	0.02MG;0.1MG **		N020683 001 Mar 27, 1997
AVIANE-21			
DURAMED PHARMS BARR	0.02MG;0.1MG		A075796 002 Apr 30, 2001
ENPRESSE-21			
DURAMED PHARMS BARR	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG		A075809 001 Jul 16, 2001
LESSINA-21			
BARR	0.02MG;0.1MG		A075803 001 Mar 20, 2002
LEVLITE			
+ BAYER HLTHCARE	0.02MG;0.1MG **		N020860 001 Jul 13, 1998
LEVONORGESTREL AND ETHINYL ESTRADIOL			
BARR	0.02MG;0.1MG		A075862 001 Apr 29, 2003
LEVORA 0.15/30-21			
WATSON LABS	0.03MG;0.15MG		A073592 001 Dec 13, 1993
NORDETTE-21			
TEVA BRANDED PHARM	0.03MG;0.15MG		N018668 001 May 10, 1982
PORTIA-21			
BARR	0.03MG;0.15MG		A075866 001 May 23, 2002
TRIPHASICL-21			
+ WYETH PHARMS	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG **		N019192 001 Nov 01, 1984
TRIVORA-21			
MAYNE PHARMA	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG		A074538 001 Dec 18, 1997

TABLET;ORAL-28			
ALESSE			
+ WYETH PHARMS	0.02MG;0.1MG **		N020683 002 Mar 27, 1997
LEVLITE			
+ BAYER HLTHCARE	0.02MG;0.1MG **		N020860 002 Jul 13, 1998
LEVONORGESTREL AND ETHINYL ESTRADIOL			
BARR	0.02MG;0.1MG		A075862 002 Apr 29, 2003
NORDETTE-28			
+ TEVA BRANDED PHARM	0.03MG;0.15MG **		N018782 001 Jul 21, 1982
TRIPHASICL-28			
+ WYETH PHARMS INC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.1 25MG **		N019190 001 Nov 01, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-148(of 375)

\*\* See List Footnote

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE;TRANSDERMAL

ORTHO EVRA

+ JANSSEN PHARMS 0.035MG/24HR;0.15MG/24HR \*\* N021180 001 Nov 20, 2001

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET;ORAL-21

BALZIVA-21

BARR 0.035MG; 0.4MG A076198 001 Apr 22, 2004

REVICON 21-DAY

ALLERGAN SALES LLC 0.035MG; 0.5MG N017566 001

GENCEPT 10/11-21

BARR 0.035MG, 0.035MG; 0.5MG, 1MG A072694 001 Feb 28, 1992

MODICON 21

ORTHO MCNEIL PHARM 0.035MG; 0.5MG N017488 001

N.E.E. 1/35 21

LPI 0.035MG; 1MG A071541 001 Dec 14, 1987

NORCEPT-E 1/35 21

ORTHO MCNEIL PHARM 0.035MG; 1MG A071545 001 Feb 09, 1989

NORETHIN 1/35E-21

WATSON PHARMS TEVA 0.035MG; 1MG A071480 001 Apr 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

WATSON LABS 0.035MG; 0.4MG A078379 001 Feb 23, 2010

0.035MG; 0.5MG A070684 001 Jan 29, 1987

WATSON PHARMS TEVA 0.035MG; 1MG A070685 001 Jan 29, 1987

NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)

WATSON LABS 0.035MG, 0.035MG; 0.5MG, 1MG A071043 001 Apr 01, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)

WATSON LABS TEVA 0.035MG, 0.035MG; 0.5MG, 1MG A071041 001 Sep 24, 1991

NORTREL 0.5/35-21

BARR 0.035MG; 0.5MG A072692 001 Feb 28, 1992

ORTHO-NOVUM 1/35-21

ORTHO MCNEIL PHARM 0.035MG; 1MG N017489 002

ORTHO-NOVUM 10/11-21

+ ORTHO MCNEIL JANSSEN 0.035MG, 0.035MG; 0.5MG, 1MG \*\* N018354 001 Jan 11, 1982

ORTHO-NOVUM 7/14-21

ORTHO MCNEIL PHARM 0.035MG, 0.035MG; 0.5MG, 1MG N019004 001 Apr 04, 1984

ORTHO-NOVUM 7/7/7-21

JANSSEN PHARMS 0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG N018985 001 Apr 04, 1984

G

OVCON-35

+ WARNER CHILCOTT 0.035MG; 0.4MG \*\* N018127 001

OVCON-50

WARNER CHILCOTT 0.05MG; 1MG N018128 001

TRI-NORINYL 21-DAY

MAYNE PHARMA 0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MG N018977 001 Apr 13, 1984

TABLET;ORAL-28

GENCEPT 10/11-28

BARR 0.035MG, 0.035MG; 0.5MG, 1MG A072697 001 Feb 28, 1992

N.E.E. 1/35 28

LPI 0.035MG; 1MG A071542 001 Dec 14, 1987

NORCEPT-E 1/35 28

ORTHO MCNEIL PHARM 0.035MG; 1MG A071546 001 Feb 09, 1989

NORETHIN 1/35E-28

WATSON LABS 0.035MG; 1MG A071481 001 Apr 12, 1988

NORETHINDRONE AND ETHINYL ESTRADIOL

MYLAN LABS LTD 0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG A200486 001 Dec 28, 2015

G

0.035MG; 0.5MG A200488 001 Oct 21, 2015

0.035MG; 1MG A200489 001 Oct 21, 2015

WATSON LABS 0.035MG, 0.035MG, 0.035MG; 0.5MG, 0.75MG, 1MG A076393 001 Feb 04, 2010

G

NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)

WATSON LABS 0.035MG, 0.035MG; 0.5MG, 1MG A071042 001 Sep 24, 1991

ORTHO-NOVUM 10/11-28

+ ORTHO MCNEIL JANSSEN 0.035MG, 0.035MG; 0.5MG, 1MG N018354 002 Jan 11, 1982

ORTHO-NOVUM 7/14-28

ORTHO MCNEIL PHARM 0.035MG, 0.035MG; 0.5MG, 1MG N019004 002 Apr 04, 1984

OVCON-35

+ WARNER CHILCOTT LLC 0.035MG; 0.4MG \*\* N017716 001

**DISCONTINUED DRUG PRODUCT LIST**

6-149(of 375)

\*\* See List Footnote

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET;ORAL-28

OVCON-50

WARNER CHILCOTT LLC 0.05MG;1MG

N017576 001

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET;ORAL

FEMHRT

+ APIL 0.005MG;1MG \*\*

N021065 002 Oct 15, 1999

TABLET;ORAL-21

ESTROSTEP 21

+ APIL 0.02MG, 0.03MG, 0.035MG;1MG,1MG,1MG \*\*

N020130 001 Oct 09, 1996

NORLESTRIN 21 1/50

PARKE DAVIS 0.05MG;1MG

N016749 001

NORLESTRIN 21 2.5/50

PARKE DAVIS 0.05MG;2.5MG

N016852 001

TABLET;ORAL-28

NORLESTRIN 28 1/50

PARKE DAVIS 0.05MG;1MG

N016723 001

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET;ORAL-21

ORTHO CYCLEN-21

JANSSEN PHARMS 0.035MG;0.25MG

N019653 001 Dec 29, 1989

ORTHO TRI-CYCLEN

JANSSEN PHARMS 0.035MG, 0.035MG, 0.035MG;0.18MG, 0.215MG, 0.25MG

N019697 002 Jul 03, 1992

TABLET;ORAL-28

NORGESTIMATE AND ETHINYL ESTRADIOL

WATSON LABS 0.025MG, 0.025MG, 0.025MG;0.18MG, 0.215MG,

A090479 001 Mar 09, 2011

0.25MG

0.035MG, 0.035MG, 0.035MG;0.18MG, 0.215MG,

A076626 001 Aug 17, 2006

0.25MG

0.035MG;0.25MG

A076627 001 Aug 17, 2006

ETHINYL ESTRADIOL; NORGESTREL

TABLET;ORAL-21

LO/OVRAL

PELAGIUS 0.03MG; 0.3MG

N017612 001

LOW-OGESTREL-21

MAYNE PHARMA 0.03MG; 0.3MG

A075288 001 Jul 28, 1999

OGESTREL 0.5/50-21

WATSON LABS 0.05MG; 0.5MG

A075406 001 Dec 15, 1999

OVRAL

WYETH PHARMS 0.05MG; 0.5MG

N016672 001

TABLET;ORAL-28

LO/OVRAL-28

WYETH PHARMS 0.03MG; 0.3MG \*\*

N017802 001

NORGESTREL AND ETHINYL ESTRADIOL

MYLAN LABS LTD 0.03MG; 0.3MG

A201828 001 Jun 21, 2016

0.05MG; 0.5MG

A202875 001 May 08, 2017

OVRAL-28

WYETH PHARMS 0.05MG; 0.5MG

N016806 001

ETHOPROPAZINE HYDROCHLORIDE

TABLET;ORAL

PARSIDOL

PARKE DAVIS 10MG

N009078 003

50MG

N009078 006

100MG

N009078 008

ETHOTOIN

TABLET;ORAL

PEGANONE

RECORDATI RARE 500MG

N010841 003

ETHOXZOLAMIDE

TABLET;ORAL

CARDRASE

PHARMACIA AND UPJOHN 62.5MG

N011047 002

125MG

N011047 001

**DISCONTINUED DRUG PRODUCT LIST**

6-150(of 375)

\*\* See List Footnote

ETHOXZOLAMIDE

TABLET;ORAL  
ETHAMIDE  
ALLERGAN 125MG N016144 001

ETHYLESTRENOL

ELIXIR;ORAL  
MAXIBOLIN  
ORGANON USA INC 2MG/5ML N014006 002  
TABLET;ORAL  
MAXIBOLIN  
ORGANON USA INC 2MG N014005 002

ETHYNODIOL DIACETATE; MESTRANOL

TABLET;ORAL-20  
OVULEN  
GD SEARLE LLC 1MG;0.1MG N016029 002  
TABLET;ORAL-21  
OVULEN-21  
GD SEARLE LLC 1MG;0.1MG N016029 003  
TABLET;ORAL-28  
OVULEN-28  
GD SEARLE LLC 1MG;0.1MG N016705 001

ETIDOCAINE HYDROCHLORIDE

INJECTABLE;INJECTION  
DURANEST  
+ ASTRAZENECA 0.5% \*\* N017751 003  
+ 1% \*\* N017751 005

ETIDRONATE DISODIUM

INJECTABLE;INJECTION  
DIDRONEL  
MGI PHARMA INC 50MG/ML N019545 001 Apr 20, 1987  
TABLET;ORAL  
DIDRONEL  
+ APIL 200MG N017831 001  
+ 400MG N017831 002

ETODOLAC

CAPSULE;ORAL  
ETODOLAC  
ANI PHARMS INC 200MG A074899 001 Jul 08, 1997  
300MG A074899 002 Jul 08, 1997  
CHARTWELL MOLECULES 200MG A074842 001 Jul 17, 1997  
300MG A074842 002 Jul 17, 1997  
ECI PHARMS LLC 300MG A074929 001 Jan 30, 1998  
IDT AUSTRALIA LTD 200MG A074840 001 Aug 29, 1997  
300MG A074840 002 Aug 29, 1997  
MYLAN 200MG A074932 001 May 16, 1997  
200MG A075071 001 Sep 30, 1998  
300MG A074932 002 May 16, 1997  
A075071 002 Sep 30, 1998  
SANDOZ 200MG A074942 001 Sep 30, 1997  
300MG A074942 002 Sep 30, 1997  
WATSON LABS 200MG A074844 001 Dec 23, 1997  
300MG A074844 002 Dec 23, 1997  
LODINE  
+ WYETH PHARMS INC 200MG \*\* N018922 002 Jan 31, 1991  
+ 300MG N018922 003 Jan 31, 1991

TABLET;ORAL  
ETODOLAC  
CHARTWELL MOLECULES 400MG A074841 001 Jun 27, 1997  
ECI PHARMS LLC 400MG A074927 001 Oct 30, 1997  
IVAX SUB TEVA PHARMS 400MG A074883 001 Feb 28, 1997  
500MG A074883 002 Nov 20, 1998  
MYLAN 400MG A075012 001 Sep 30, 1998  
500MG A075012 002 Sep 30, 1998  
MYLAN PHARMS INC 400MG A075104 001 Feb 06, 1998  
500MG A075104 002 Nov 20, 1998  
OXFORD PHARMS 400MG A074819 001 Feb 28, 1997  
500MG A074819 002 Apr 28, 1998

**DISCONTINUED DRUG PRODUCT LIST**

6-151(of 375)

\*\* See List Footnote

**ETODOLAC**TABLET;ORAL  
ETODOLAC

RANBAXY LABS LTD	400MG	A075226 001	Nov 24, 1998
	500MG	A075226 002	Nov 24, 1998
SANDOZ	400MG	A074839 001	Jul 11, 1997
	400MG	A074846 001	Feb 28, 1997
TEVA	400MG	A074847 001	Apr 23, 1999
	500MG	A074847 002	Apr 23, 1999
WATSON LABS	400MG	A074892 001	Apr 16, 1997
	400MG	A075069 001	Apr 16, 1998
	500MG	A074892 002	Oct 29, 1998

## LODINE

+ WYETH PHARMS INC	400MG **	N018922 004	Jul 29, 1993
+	500MG **	N018922 005	Jun 28, 1996

TABLET, EXTENDED RELEASE;ORAL

## ETODOLAC

ACTAVIS ELIZABETH	400MG	A075696 001	Jul 31, 2000
IDT AUSTRALIA LTD	400MG	A075943 001	Jul 26, 2002
	500MG	A075943 002	Jul 26, 2002
	600MG	A075943 003	Jul 26, 2002
WATSON LABS FLORIDA	400MG	A075829 001	Nov 30, 2001
	500MG	A075829 002	Nov 30, 2001

## LODINE XL

+ WYETH PHARMS INC	400MG **	N020584 001	Oct 25, 1996
	500MG **	N020584 003	Jan 20, 1998
+	600MG **	N020584 002	Oct 25, 1996

**ETONOGESTREL**

IMPLANT; IMPLANTATION

## IMPLANON

ORGANON USA INC	68MG/IMPLANT	N021529 001	Jul 17, 2006
-----------------	--------------	-------------	--------------

**ETOPOSIDE**

CAPSULE;ORAL

## VEPESID

+ DAVA PHARMS INC	50MG	N019557 001	Dec 30, 1986
+	100MG	N019557 002	Dec 30, 1986

INJECTABLE;INJECTION

## ETOPOSIDE

HOSPIRA	20MG/ML	A074320 001	Aug 30, 1995
	20MG/ML	A074351 001	Aug 30, 1995
PHARMACHEMIE BV	20MG/ML	A074227 001	Feb 22, 1996
PIERRE FABRE	20MG/ML	A074813 001	Jul 09, 1997
TEVA PARENTERAL	20MG/ML	A074510 001	Jun 29, 1995
TEVA PHARMS USA	20MG/ML	A074284 001	Feb 10, 1994
WATSON LABS	20MG/ML	A074228 001	Oct 15, 1996
WATSON LABS INC	20MG/ML	A074968 001	Jan 09, 1998

## TOPOSAR

TEVA PARENTERAL	20MG/ML	A074166 001	Feb 27, 1995
-----------------	---------	-------------	--------------

## VEPESID

+ CORDEN PHARMA	20MG/ML **	N018768 001	Nov 10, 1983
-----------------	------------	-------------	--------------

**ETOPOSIDE PHOSPHATE**

INJECTABLE;INJECTION

## ETOPOPHOS PRESERVATIVE FREE

BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N020906 001	Feb 27, 1998
	EQ 1GM BASE/VIAL	N020906 002	Feb 27, 1998

**ETRETINATE**

CAPSULE;ORAL

## TEGISON

ROCHE	10MG	N019369 001	Sep 30, 1986
	25MG	N019369 002	Sep 30, 1986

**EVANS BLUE**

INJECTABLE;INJECTION

## EVANS BLUE

PARKE DAVIS	0.5% **	N008041 001	
-------------	---------	-------------	--

**DISCONTINUED DRUG PRODUCT LIST**

6-152(of 375)

\*\* See List Footnote

**EZOGABINE**

TABLET;ORAL

POTIGA

+ GLAXOSMITHKLINE	50MG	N022345 001	Jun 10, 2011
+	200MG	N022345 002	Jun 10, 2011
+	300MG	N022345 003	Jun 10, 2011
+	400MG	N022345 004	Jun 10, 2011

**FAMCICLOVIR**

TABLET;ORAL

FAMVIR

+ NOVARTIS	125MG	N020363 003	Dec 11, 1995
+	250MG	N020363 001	Apr 26, 1996
+	500MG	N020363 002	Jun 29, 1994

**FAMOTIDINE**

INJECTABLE;INJECTION

FAMOTIDINE

APOTEX INC	10MG/ML	A075942 001	Aug 02, 2002
APOTHECON	10MG/ML	A075707 001	Apr 16, 2001
HOSPIRA	10MG/ML	A075705 001	Apr 16, 2001
	10MG/ML	A075870 001	Nov 23, 2001
	10MG/ML	A075905 001	Nov 23, 2001
WEST-WARD PHARMS INT	10MG/ML	A075799 001	Apr 30, 2002

FAMOTIDINE PRESERVATIVE FREE

APOTEX INC	10MG/ML	A076324 001	Nov 27, 2002
APOTHECON	10MG/ML	A075708 001	Apr 16, 2001
HOSPIRA	10MG/ML	A075669 001	Apr 16, 2001
WEST-WARD PHARMS INT	10MG/ML	A075789 001	Apr 30, 2002

FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)

APOTEX INC	10MG/ML	A076322 001	Nov 27, 2002
------------	---------	-------------	--------------

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

ABBVIE	0.4MG/ML	A075729 001	Dec 17, 2001
--------	----------	-------------	--------------

PEPCID

+ MERCK	10MG/ML **	N019510 001	Nov 04, 1986
---------	------------	-------------	--------------

PEPCID PRESERVATIVE FREE

+ MERCK	10MG/ML **	N019510 004	Nov 04, 1986
---------	------------	-------------	--------------

PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER

+ MERCK SHARP DOHME	0.4MG/ML **	N020249 001	Feb 18, 1994
---------------------	-------------	-------------	--------------

TABLET;ORAL

FAMOTIDINE

ACTAVIS ELIZABETH	20MG	A075650 001	Sep 14, 2001
	40MG	A075650 002	Sep 14, 2001
APOTEX	10MG	A075610 001	Mar 12, 2002
MYLAN PHARMS INC	20MG	A075457 001	Apr 18, 2001
	40MG	A075457 002	Apr 18, 2001
SANDOZ	10MG	A076101 001	Oct 21, 2002
	20MG	A075302 001	Apr 16, 2001
	20MG	A075607 001	May 10, 2001
	20MG	A075793 001	Apr 16, 2001
	40MG	A075302 002	Apr 16, 2001
	40MG	A075607 002	May 10, 2001
	40MG	A075793 002	Apr 16, 2001
SUN PHARM INDUSTRIES	20MG	A075639 002	Dec 12, 2001
	40MG	A075639 001	Dec 12, 2001
WATSON LABS	10MG	A075404 001	Nov 28, 2001
	20MG	A075062 002	Apr 16, 2001
	40MG	A075062 001	Apr 16, 2001

TABLET, CHEWABLE;ORAL

PEPCID AC

+ J AND J CONSUMER INC	10MG **	N020801 001	Sep 24, 1998
------------------------	---------	-------------	--------------

TABLET, ORALLY DISINTEGRATING;ORAL

FLUXID

UCB INC	20MG	N021712 001	Sep 24, 2004
	40MG	N021712 002	Sep 24, 2004

PEPCID RPD

MERCK	20MG	N020752 001	May 28, 1998
	40MG	N020752 002	May 28, 1998

**DISCONTINUED DRUG PRODUCT LIST**

6-153(of 375)

\*\* See List Footnote

**FELODIPINE**TABLET, EXTENDED RELEASE;ORAL  
FELODIPINE

WOCKHARDT LTD	2.5MG	A091484	001	Aug 15, 2012
	5MG	A091484	002	Aug 15, 2012
	10MG	A091484	003	Aug 15, 2012
PLENDIL				
+ ASTRAZENECA	2.5MG **	N019834	004	Sep 22, 1994
+ +	5MG **	N019834	001	Jul 25, 1991
+ +	10MG **	N019834	002	Jul 25, 1991

**FENOFIBRATE**

CAPSULE;ORAL

ANTARA (MICRONIZED)				
LUPIN ATLANTIS	87MG	N021695	002	Nov 30, 2004
LIPIDIL				
ABBVIE	100MG	N019304	001	Dec 31, 1993
LIPOFEN				
CIPHER PHARMS INC	100MG	N021612	002	Jan 11, 2006
TRICOR (MICRONIZED)				
+ ABBVIE	67MG **	N019304	002	Feb 09, 1998
+ +	134MG **	N019304	003	Jun 30, 1999
+ +	200MG **	N019304	004	Jun 30, 1999

TABLET;ORAL

FENOFIBRATE				
MYLAN	107MG	A076520	002	Dec 29, 2005
TRICOR				
+ ABBVIE INC	54MG **	N021203	001	Sep 04, 2001
+ +	160MG **	N021203	003	Sep 04, 2001
TRIGLIDE				
SKYEPHARMA AG	50MG	N021350	001	May 07, 2005

**FENOLDOPAM MESYLATE**

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE				
LUITPOLD	EQ 10MG BASE/ML	A076656	001	Dec 01, 2003
TEVA PARENTERAL	EQ 10MG BASE/ML	A077826	001	Mar 07, 2007

**FENOPROFEN CALCIUM**

CAPSULE;ORAL

FENOPROFEN CALCIUM				
AM THERAP	EQ 200MG BASE	A072307	001	Aug 22, 1988
	EQ 300MG BASE	A072308	001	Aug 22, 1988
AUROLIFE PHARMA LLC	EQ 200MG BASE	A072394	001	Oct 17, 1988
	EQ 300MG BASE	A072395	001	Oct 17, 1988
HALSEY	EQ 200MG BASE	A072355	001	Aug 17, 1988
	EQ 300MG BASE	A072356	001	Aug 17, 1988
PAR PHARM	EQ 200MG BASE	A072437	001	Aug 22, 1988
	EQ 300MG BASE	A072438	001	Aug 22, 1988
QUANTUM PHARMICS	EQ 200MG BASE	A072214	001	Aug 17, 1988
	EQ 300MG BASE	A071738	001	Aug 17, 1988
WARNER CHILCOTT	EQ 200MG BASE	A072946	001	Apr 30, 1991
	EQ 300MG BASE	A072472	001	Apr 30, 1991
WATSON LABS	EQ 200MG BASE	A072294	001	Aug 17, 1988
	EQ 200MG BASE	A072981	001	Aug 19, 1991
	EQ 300MG BASE	A072293	001	Aug 17, 1988
	EQ 300MG BASE	A072982	001	Aug 19, 1991

NALFON

XSPIRE PHARMA	EQ 300MG BASE	N017604	002	
---------------	---------------	---------	-----	--

TABLET;ORAL

FENOPROFEN CALCIUM				
ACTAVIS ELIZABETH	EQ 600MG BASE	A072274	001	May 02, 1988
AM THERAP	EQ 600MG BASE	A072309	001	Aug 17, 1988
AUROLIFE PHARMA LLC	EQ 600MG BASE	A072396	001	Oct 17, 1988
DAVA PHARMS INC	EQ 600MG BASE	A072326	001	Aug 17, 1988
HALSEY	EQ 600MG BASE	A072357	001	Aug 17, 1988
IVAX SUB TEVA PHARMS	EQ 600MG BASE	A072557	001	Aug 29, 1988
PAR PHARM	EQ 600MG BASE	A072429	001	Aug 17, 1988
QUANTUM PHARMICS	EQ 600MG BASE	A072194	001	Aug 17, 1988
SUN PHARM INDUSTRIES	EQ 600MG BASE	A072902	001	Dec 21, 1990
USL PHARMA	EQ 600MG BASE	A072362	001	Aug 17, 1988

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-154(of 375)

\*\* See List Footnote

**FENOPROFEN CALCIUM**

TABLET;ORAL

FENOPROFEN CALCIUM

WATSON LABS	EQ 600MG BASE	A072165 001 Aug 17, 1988
	EQ 600MG BASE	A072602 001 Oct 11, 1988
WATSON LABS TEVA	EQ 600MG BASE	A072407 001 Aug 17, 1988
NALFON		
DISTA	EQ 600MG BASE	N017710 001

**FENTANYL**

FILM, EXTENDED RELEASE;TRANSDERMAL

FENTANYL-100		
NOVEN	100MCG/HR	A077775 004 Oct 16, 2009
FENTANYL-25		
NOVEN	25MCG/HR	A077775 001 Oct 16, 2009
FENTANYL-50		
NOVEN	50MCG/HR	A077775 002 Oct 16, 2009
FENTANYL-75		
NOVEN	75MCG/HR	A077775 003 Oct 16, 2009

**FENTANYL CITRATE**

FILM;BUCCAL

ONSOLIS		
BDSI	EQ 0.2MG BASE	N022266 001 Jul 16, 2009
	EQ 0.4MG BASE	N022266 002 Jul 16, 2009
	EQ 0.6MG BASE	N022266 003 Jul 16, 2009
	EQ 0.8MG BASE	N022266 004 Jul 16, 2009
	EQ 1.2MG BASE	N022266 005 Jul 16, 2009

INJECTABLE;INJECTION

FENTANYL CITRATE		
ABBOTT	EQ 0.05MG BASE/ML	A070636 001 Apr 30, 1990
	EQ 0.05MG BASE/ML	A070637 001 Apr 30, 1990
WATSON LABS	EQ 0.05MG BASE/ML	A073488 001 Jun 30, 1992
FENTANYL CITRATE PRESERVATIVE FREE		
WATSON LABS INC	EQ 0.05MG BASE/ML	A074917 001 Feb 03, 1998

TABLET;BUCCAL, SUBLINGUAL

FENTANYL CITRATE		
WATSON LABS	EQ 0.1MG BASE	A079075 001 Jan 07, 2011
	EQ 0.2MG BASE	A079075 002 Jan 07, 2011
	EQ 0.4MG BASE	A079075 003 Jan 07, 2011
	EQ 0.6MG BASE	A079075 004 Jan 07, 2011
	EQ 0.8MG BASE	A079075 005 Jan 07, 2011

FENTORA		
+ CEPHALON	EQ 0.3MG BASE **	N021947 006 Mar 02, 2007

TROCHE/LOZENGE;ORAL

FENTANYL		
CEPHALON	EQ 0.1MG BASE	N020195 007 Oct 30, 1995
	EQ 0.2MG BASE	N020195 001 Oct 04, 1993
	EQ 0.3MG BASE	N020195 002 Oct 04, 1993
	EQ 0.4MG BASE	N020195 003 Oct 04, 1993

**FERRIC AMMONIUM CITRATE**

FOR SOLUTION;ORAL

FERRISELTZ		
OTSUKA	600MG/PACKET	N020292 001 Oct 14, 1997

**FERROUS CITRATE, FE-59**

INJECTABLE;INJECTION

FERROUS CITRATE FE 59		
MALLINCKRODT	25uCi/ML	N016729 001

**FERROUS SULFATE; FOLIC ACID**

CAPSULE;ORAL

FOLVRON		
LEDERLE	182MG;0.33MG	N006012 003

**DISCONTINUED DRUG PRODUCT LIST**

6-155(of 375)

\*\* See List Footnote

FERUMOXIDES

INJECTABLE; INJECTION

FERIDEX I.V.

AMAG PHARMS INC

EQ 11.2MG IRON/ML

N020416 001 Aug 30, 1996

FERUMOXSTIL

SUSPENSION; ORAL

GASTROMARK

AMAG PHARMS INC

EQ 0.175MG IRON/ML

N020410 001 Dec 06, 1996

FESOTERODINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

FESOTERODINE FUMARATE

ALKEM LABS LTD

4MG

A204827 001 Dec 10, 2015

8MG

A204827 002 Dec 10, 2015

FEXOFENADINE HYDROCHLORIDE

CAPSULE; ORAL

ALLEGRA

SANOFI AVENTIS US

60MG \*\*

N020625 001 Jul 25, 1996

FEXOFENADINE HYDROCHLORIDE

BARR

60MG

A076169 001 Jul 13, 2005

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

BARR

60MG;120MG

A076236 001 Apr 14, 2005

IMPAX PHARMS

60MG;120MG

A076298 001 Nov 12, 2010

FIBRINOGEN, I-125

INJECTABLE; INJECTION

IBRIN

GE HEALTHCARE

154uCi/VIAL

N017879 001

RADIONUCLIDE-LABELED (125 I) FIBRINOGEN (HUMAN) SENSOR

ABBOTT

140uCi/ML

N017787 001

FINASTERIDE

TABLET; ORAL

FINASTERIDE

IVAX SUB TEVA PHARMS

5MG

A076340 001 Jun 19, 2006

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDE

IMPAX PHARMS

100MG

A076234 001 Aug 28, 2003

URISPAS

ORTHO MCNEIL JANSSEN

100MG

N016769 001

FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATE

APOTEX INC

50MG

A079164 001 Jul 09, 2009

100MG

A079164 002 Jul 09, 2009

150MG

A079164 003 Jul 09, 2009

IDT AUSTRALIA LTD

50MG

A076030 001 Oct 28, 2002

100MG

A076030 002 Oct 28, 2002

150MG

A076030 003 Oct 28, 2002

TAMBOCOR

CNTY LINE PHARMS

200MG

N018830 002 Oct 31, 1985

FLORBETAPIR F-18

SOLUTION; INTRAVENOUS

AMYVID

AVID RADIOPHARMS INC 10ML (13.5-51mCi/ML)

N202008 001 Apr 06, 2012

FLOXURIDINE

INJECTABLE; INJECTION

FUUDR

+ HOSPIRA

500MG/VIAL \*\*

N016929 001

**DISCONTINUED DRUG PRODUCT LIST**

6-156(of 375)

\*\* See List Footnote

**FLUCONAZOLE**

FOR SUSPENSION;ORAL

**FLUCONAZOLE**

SUN PHARM INDS LTD	50MG/5ML 200MG/5ML	A076332 001 Jul 29, 2004 A076332 002 Jul 29, 2004
TARO PHARM INDS	50MG/5ML 200MG/5ML	A076918 001 Dec 18, 2006 A076918 002 Dec 18, 2006

**INJECTABLE;INJECTION**

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER		
+ PFIZER	200MG/100ML (2MG/ML)	N019950 003 Sep 29, 1992
+	400MG/200ML (2MG/ML)	N019950 005 Jul 08, 1994

DIFLUCAN IN SODIUM CHLORIDE 0.9%		
+ PFIZER	200MG/100ML (2MG/ML)	N019950 001 Jan 29, 1990
+	400MG/200ML (2MG/ML)	N019950 006 Jan 29, 1990

DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
+ PFIZER	200MG/100ML (2MG/ML)	N019950 002 Jan 29, 1990
+	400MG/200ML (2MG/ML)	N019950 004 Jan 29, 1990

FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER		
MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076888 001 Mar 25, 2005
	400MG/200ML (2MG/ML)	A076888 002 Mar 25, 2005

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%		
TEVA PHARMS USA	200MG/100ML (2MG/ML)	A076653 001 Jul 29, 2004
	400MG/200ML (2MG/ML)	A076653 002 Jul 29, 2004

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
HOSPIRA	200MG/100ML (2MG/ML)	A076617 001 Jul 29, 2004
	400MG/200ML (2MG/ML)	A076617 002 Jul 29, 2004

MYLAN LABS LTD	200MG/100ML (2MG/ML)	A076889 001 Mar 25, 2005
	400MG/200ML (2MG/ML)	A076889 002 Mar 25, 2005
TEVA PHARMS	200MG/100ML (2MG/ML)	A076837 001 Jan 13, 2005

	400MG/200ML (2MG/ML)	A076837 002 Jan 13, 2005
--	----------------------	--------------------------

**TABLET;ORAL**

FLUCONAZOLE		
GEDEON RICHTER USA	50MG	A076432 001 Jul 29, 2004
	100MG	A076432 002 Jul 29, 2004
	150MG	A076432 003 Jul 29, 2004
	200MG	A076432 004 Jul 29, 2004

IDT AUSTRALIA LTD	50MG	A076086 001 Jul 29, 2004
	100MG	A076086 002 Jul 29, 2004
	150MG	A076086 003 Jul 29, 2004
	200MG	A076086 004 Jul 29, 2004

MYLAN PHARMS INC	50MG	A076042 001 Jul 29, 2004
	100MG	A076042 002 Jul 29, 2004
	150MG	A076042 003 Jul 29, 2004
	200MG	A076042 004 Jul 29, 2004

PLIVA	50MG	A076424 001 Jul 29, 2004
	100MG	A076424 002 Jul 29, 2004
	150MG	A076424 003 Jul 29, 2004
	200MG	A076424 004 Jul 29, 2004

RANBAXY LABS LTD	50MG	A076386 001 Jul 29, 2004
	100MG	A076386 002 Jul 29, 2004
	150MG	A076386 003 Jul 29, 2004
	200MG	A076386 004 Jul 29, 2004

ROXANE	50MG	A076213 001 Jul 29, 2004
	100MG	A076213 002 Jul 29, 2004
	150MG	A076213 003 Jul 29, 2004
	200MG	A076213 004 Jul 29, 2004

**FLUDARABINE PHOSPHATE**

INJECTABLE;INJECTION		
FLUDARA		

+ GENZYME CORP	50MG/VIAL **	N020038 001 Apr 18, 1991
----------------	--------------	--------------------------

**TABLET;ORAL**

OFORTA		
SANOFI AVENTIS US	10MG	N022273 001 Dec 18, 2008

**DISCONTINUED DRUG PRODUCT LIST**

6-157(of 375)

\*\* See List Footnote

FLUDEOXYGLUCOSE F-18

INJECTABLE; INJECTION			
FLUDEOXYGLUCOSE F18			
+ DOWNSTATE CLINCL	4-40mCi/ML **	N020306 001	Aug 19, 1994
+	4-90mCi/ML **	N020306 002	Sep 25, 2001
INJECTABLE; INTRAVENOUS			
FLUDEOXYGLUCOSE F18			
WEILL MEDCL COLL	10-100mCi/ML **	N021768 001	Aug 05, 2004

FLUDROCORTISONE ACETATE

TABLET; ORAL			
FLORINEF			
+ CASPER PHARMA LLC	0.1MG **	N010060 001	

FLUMAZENIL

INJECTABLE; INJECTION			
FLUMAZENIL			
BAXTER HLTHCARE CORP	0.5MG/5ML (0.1MG/ML)	A076755 002	Oct 12, 2004
	1MG/10ML (0.1MG/ML)	A076755 001	Oct 12, 2004
TEVA PHARMS USA	0.5MG/5ML (0.1MG/ML)	A076589 002	Oct 12, 2004
	1MG/10ML (0.1MG/ML)	A076589 001	Oct 12, 2004
ROMAZICON			
+ HOFFMANN LA ROCHE	1MG/10ML (0.1MG/ML) **	N020073 001	Dec 20, 1991
+	0.5MG/5ML (0.1MG/ML) **	N020073 002	Dec 20, 1991

FLUMETHASONE PIVALATE

CREAM; TOPICAL			
LOCORTEN			
NOVARTIS	0.03%	N016379 001	

FLUNISOLIDE

AEROSOL, METERED; INHALATION			
AEROBID			
ROCHE PALO	0.25MG/INH	N018340 001	Aug 17, 1984
SPRAY, METERED; NASAL			
NASALIDE			
IVAX RES	0.025MG/SPRAY	N018148 001	
NASAREL			
TEVA BRANDED PHARM	0.029MG/SPRAY	N020409 001	Mar 08, 1995

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL			
FLUOCET			
ALPHARMA US PHARMS	0.025%	A088360 001	Jan 16, 1984
FLUOCINOLONE ACETONIDE			
ALPHARMA US PHARMS	0.01%	A088361 001	Jan 16, 1984
PERRIGO NEW YORK	0.01%	A086810 001	Mar 04, 1982
	0.025%	A086811 001	Mar 04, 1982
PHARMADERM	0.01%	A088047 001	Dec 16, 1982
	0.025%	A088045 001	Dec 16, 1982
PHARMAFAIR	0.01%	A088499 001	Aug 02, 1984
	0.025%	A088506 001	Aug 02, 1984
TARO	0.01%	A040035 001	Oct 31, 1994
	0.01%	A087102 001	Apr 27, 1982
	0.025%	A040042 001	Oct 31, 1994
USL PHARMA	0.01%	A088757 001	Feb 11, 1985
	0.025%	A088756 001	Mar 28, 1985
FLUONID			
ALLERGAN HERBERT	0.025%	A087156 002	Sep 06, 1984
FLUOTREX			
SAVAGE LABS	0.01%	A088174 001	May 06, 1983
	0.025%	A088173 001	Mar 09, 1983
SYNALAR-HP			
MEDIMETRIKS PHARMS	0.2%	N016161 002	
GEL; TOPICAL			
FLUONID			
ALLERGAN HERBERT	0.025%	A087300 001	May 27, 1982
OINTMENT; TOPICAL			
FLUOCINOLONE ACETONIDE			
PHARMADERM	0.025%	A088046 001	Dec 16, 1982
PHARMAFAIR	0.025%	A088507 001	Feb 27, 1984
USL PHARMA	0.025%	A088742 001	Feb 08, 1985

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-158(of 375)

\*\* See List Footnote

FLUOCINOLONE ACETONIDE

OINTMENT;TOPICAL

FLUONID

ALLERGAN HERBERT 0.025%

A087157 001 Sep 06, 1984

FLUOTREX

SAVAGE LABS 0.025%

A088172 001 Mar 09, 1983

SOLUTION;TOPICAL

FLUOCINOLONE ACETONIDE

ALPHARMA US PHARMS 0.01%

A087159 001 Jun 16, 1982

BAUSCH AND LOMB 0.01%

A040059 001 Dec 20, 1993

MORTON GROVE 0.01%

A088312 001 Jan 27, 1984

PHARMADERM 0.01%

A088048 001 Dec 16, 1982

PHARMAFAIR 0.01%

A088449 001 Feb 08, 1984

FLUONID

ALLERGAN HERBERT 0.01%

A087158 001 Mar 17, 1983

FLUOTREX

SAVAGE LABS 0.01%

A088171 001 Mar 09, 1983

FLUOCINONIDE

CREAM;TOPICAL

FLUOCINONIDE

PERRIGO NEW YORK 0.05%

A071790 001 Jul 13, 1988

LIDEX

+ CNTY LINE PHARMS 0.05%

N016908 002

LIDEX-E

+ CNTY LINE PHARMS 0.05%

N016908 003

SOLUTION;TOPICAL

FLUOCINONIDE

TARO 0.05%

A072857 001 Aug 02, 1989

TEVA PHARMS 0.05%

A072522 001 Sep 28, 1990

FLUORESCIN SODIUM

INJECTABLE; INJECTION

FUNDUSCEIN-25

+ NOVARTIS 25% \*\*

N017869 001

FLUOROMETHOLONE

CREAM;TOPICAL

OXYLONE

PHARMACIA AND UPJOHN 0.025%

N011748 001

SUSPENSION/DROPS;OPHTHALMIC

FLUOR-OP

NOVARTIS 0.1%

A070185 001 Feb 27, 1986

FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS;OPHTHALMIC

TOBRASONE

ALCON 0.1%;0.3%

N050628 001 Jul 21, 1989

FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS;OPHTHALMIC

FML-S

ALLERGAN 0.1%;10%

N019525 001 Sep 29, 1989

FLUOROURACIL

INJECTABLE; INJECTION

ADRUCIL

PHARMACIA AND UPJOHN 50MG/ML

A081222 001 Jun 28, 1991

50MG/ML

N017959 001

TEVA PARENTERAL 50MG/ML

A040023 001 Oct 18, 1991

50MG/ML

A081225 001 Aug 28, 1991

FLUOROURACIL

ABIC 50MG/ML

A088929 001 Mar 04, 1986

ABRAXIS PHARM 50MG/ML

A089152 001 Mar 21, 1986

50MG/ML

A089428 001 Jan 12, 1987

BEDFORD 50MG/ML

A089519 001 Mar 12, 1987

EBWE PHARMA 500MG/10ML (50MG/ML)

A089508 001 Jan 26, 1988

FRESENIUS KABI USA 50MG/ML

A040772 001 Aug 11, 2008

50MG/ML

A040291 001 Mar 24, 1999

MARCHAR 50MG/ML

A040379 001 Nov 15, 2000

SANDOZ 2.5GM/50ML (50MG/ML)

A087791 001 Jan 18, 1983

A091299 001 May 02, 2011

**DISCONTINUED DRUG PRODUCT LIST**

6-159(of 375)

\*\* See List Footnote

**FLUOROURACIL**

INJECTABLE; INJECTION

FLUOROURACIL

	5GM/100ML (50MG/ML)		
SMITH AND NEPHEW	50MG/ML	A091299	002 May 02, 2011
	50MG/ML	A088766	001 Dec 28, 1984
	50MG/ML	A088767	001 Dec 28, 1984
SPECTRUM PHARMS	50MG/ML	A089434	001 Mar 26, 1987
+	500MG/10ML (50MG/ML) **	A087792	001 Oct 13, 1982
+	2.5GM/50ML (50MG/ML)	N012209	001
		N012209	002 Jul 29, 2016

SOLUTION;TOPICAL

FLUOROPLEX

ELORAC 1%

N016765 001

**FLUOXETINE HYDROCHLORIDE**

CAPSULE;ORAL

FLUOXETINE

SUN PHARM INDUSTRIES	EQ 10MG BASE	A075787	001 Jan 29, 2002
	EQ 20MG BASE	A075787	002 Jan 29, 2002
WATSON LABS	EQ 10MG BASE	A075662	001 Jan 29, 2002
	EQ 20MG BASE	A075662	002 Jan 29, 2002

FLUOXETINE HYDROCHLORIDE

ANI PHARMS INC	EQ 10MG BASE	A076287	001 May 20, 2008
	EQ 20MG BASE	A076287	002 May 20, 2008
BARR	EQ 40MG BASE	A076251	001 May 18, 2005
CARLSBAD	EQ 10MG BASE	A076022	001 Jan 30, 2002
	EQ 20MG BASE	A076022	002 Jan 30, 2002
CR DOUBLE CRANE	EQ 10MG BASE	A076165	001 Feb 01, 2002
	EQ 20MG BASE	A076165	002 Feb 01, 2002
MYLAN	EQ 10MG BASE	A075207	001 Jan 30, 2002
	EQ 20MG BASE	A075207	002 Jan 30, 2002
	EQ 40MG BASE	A075207	003 May 25, 2007
MYLAN PHARMS INC	EQ 10MG BASE	A075577	001 Jan 29, 2002
	EQ 20MG BASE	A075577	002 Jan 29, 2002
PAR PHARM	EQ 10MG BASE	A076922	001 Dec 16, 2004
	EQ 20MG BASE	A076922	002 Dec 16, 2004
SANDOZ	EQ 10MG BASE	A075807	001 Jan 29, 2002
	EQ 10MG BASE	A077469	001 Nov 17, 2008
	EQ 20MG BASE	A075807	002 Jan 29, 2002
	EQ 20MG BASE	A077469	002 Nov 17, 2008
WOCKHARDT LTD	EQ 10MG BASE	A078143	001 Jan 16, 2008
	EQ 20MG BASE	A078143	002 Jan 16, 2008
	EQ 40MG BASE	A078143	003 Jan 16, 2008

PROZAC

ELI LILLY AND CO	EQ 60MG BASE	N018936	004 Jun 15, 1999
SARAFEM			
+ ELI LILLY AND CO	EQ 10MG BASE **	N018936	007 Jul 06, 2000
+	EQ 20MG BASE **	N018936	008 Jul 06, 2000

SOLUTION;ORAL

FLUOXETINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 20MG BASE/5ML	A075690	001 Jan 31, 2002
APOTEX INC	EQ 20MG BASE/5ML	A075292	001 Feb 07, 2002
HI TECH PHARMA	EQ 20MG BASE/5ML	A075525	001 Jun 27, 2002
LANNETT	EQ 20MG BASE/5ML	A076458	001 May 14, 2004

PROZAC

+ LILLY	EQ 20MG BASE/5ML **	N020101	001 Apr 24, 1991
---------	---------------------	---------	------------------

TABLET;ORAL

FLUOXETINE HYDROCHLORIDE

BARR	EQ 10MG BASE	A075810	001 Feb 01, 2002
IVAX SUB TEVA PHARMS	EQ 10MG BASE	A075865	001 Feb 28, 2002
	EQ 40MG BASE	A075865	003 Aug 30, 2004

SANDOZ EQ 10MG BASE

PROZAC		A076024	001 Jan 29, 2002
+ LILLY	EQ 10MG BASE **	N020974	001 Mar 09, 1999
+	EQ 20MG BASE **	N020974	002 Mar 09, 1999

**DISCONTINUED DRUG PRODUCT LIST**

6-160(of 375)

\*\* See List Footnote

FLUOXYMESTERONE

TABLET;ORAL ANDROID-F			
VALEANT PHARM INTL	10MG	A087196	001
FLUOXYMESTERONE			
VALEANT PHARM INTL	10MG	A088221	001 May 05, 1983
WATSON LABS	2MG	A088260	001 Dec 06, 1983
	5MG	A088265	001 Dec 06, 1983
	10MG	A088309	001 Dec 06, 1983
HALOTESTIN			
PHARMACIA AND UPJOHN	2MG	N010611	002
	5MG	N010611	006
	10MG	N010611	010
ORA-TESTRYL			
BRISTOL MYERS SQUIBB	2MG	N011359	001
	5MG	N011359	002

FLUPHENAZINE DECANOATE

INJECTABLE;INJECTION FLUPHENAZINE DECANOATE			
HOSPIRA	25MG/ML	A074966	001 Apr 16, 1998
MYLAN LABS LTD	25MG/ML	A075918	001 Aug 17, 2001
TEVA PARENTERAL	25MG/ML	A074795	001 Sep 10, 1996
PROLIXIN DECANOATE			
+ BRISTOL MYERS SQUIBB	25MG/ML **	N016727	001

FLUPHENAZINE ENANTHATE

INJECTABLE;INJECTION PROLIXIN ENANTHATE			
APOTHECON	25MG/ML	N016110	001

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE;ORAL FLUPHENAZINE HYDROCHLORIDE			
ANI PHARMS INC	5MG/ML	A073058	001 Aug 30, 1991
PERMITIL SCHERING	5MG/ML	N016008	001
PROLIXIN APOTHECON	5MG/ML	A070533	001 Nov 07, 1985
ELIXIR;ORAL FLUPHENAZINE HYDROCHLORIDE			
ANI PHARMS INC	2.5MG/5ML	A081310	001 Apr 29, 1993
PROLIXIN + APOTHECON	2.5MG/5ML **	N012145	003
INJECTABLE;INJECTION PROLIXIN			
APOTHECON	2.5MG/ML	N011751	005
TABLET;ORAL FLUPHENAZINE HYDROCHLORIDE			
WATSON LABS	1MG	A088555	001 Dec 18, 1987
	2.5MG	A088544	001 Dec 18, 1987
	5MG	A088527	001 Dec 18, 1987
	10MG	A088550	001 Dec 18, 1987
PERMITIL SCHERING	0.25MG	N012034	001
	2.5MG	N012034	004
	5MG	N012034	005
	10MG	N012034	006
PROLIXIN + APOTHECON	1MG	N011751	004
+	2.5MG	N011751	001
+	5MG	N011751	003
+	10MG	N011751	002
TABLET, EXTENDED RELEASE;ORAL PERMITIL SCHERING	1MG	N012419	004

**DISCONTINUED DRUG PRODUCT LIST**

6-161(of 375)

\*\* See List Footnote

FLUPREDNISOLONETABLET;ORAL  
ALPHADROL

PHARMACIA AND UPJOHN 1.5MG

N012259 002

FLURANDRENOLIDELOTION;TOPICAL  
FLURANDRENOLIDE

ALPHARMA US PHARMS 0.05%

A087203 001 Apr 29, 1982

OINTMENT;TOPICAL  
CORDRAN

+ AQUA PHARMS 0.025% \*\*

N012806 004

FLURANDRENOLIDE; NEOMYCIN SULFATECREAM;TOPICAL  
CORDRAN N

LILLY 0.05%;EQ 3.5MG BASE/GM

N050346 001

OINTMENT;TOPICAL  
CORDRAN N

LILLY 0.05%;EQ 3.5MG BASE/GM

N050345 001

FLURAZEPAM HYDROCHLORIDECAPSULE;ORAL  
DALMANEVALEANT PHARM INTL 15MG \*\*  
+ 30MG \*\*N016721 001  
N016721 002

FLURAZEPAM HYDROCHLORIDE

AUROLIFE PHARMA LLC	15MG	A071717 002	Jul 31, 1991
	30MG	A071717 001	Jul 31, 1991
HALSEY	15MG	A071808 001	Jan 07, 1988
	30MG	A071809 001	Jan 07, 1988
HIKMA INTL PHARMS	15MG	A071107 001	Dec 08, 1986
HIKMA PHARMS	30MG	A071108 001	Dec 08, 1986
PAR PHARM	15MG	A070444 001	Mar 20, 1986
	30MG	A070445 001	Mar 20, 1986
PUREPAC PHARM	15MG	A071927 001	Sep 09, 1987
	30MG	A071551 001	Sep 09, 1987
SUN PHARM INDUSTRIES	15MG	A070454 001	Aug 04, 1986
	30MG	A070455 001	Aug 04, 1986
SUPERPHARM	15MG	A071659 001	Aug 04, 1988
	30MG	A071660 001	Aug 04, 1988
USL PHARMA	15MG	A070562 001	Jul 09, 1987
	30MG	A070563 001	Jul 09, 1987
WARNER CHILCOTT	15MG	A071767 001	Dec 04, 1987
	30MG	A071768 001	Dec 04, 1987
WATSON LABS	15MG	A071205 001	Nov 25, 1986
	15MG	A072368 001	Mar 30, 1989
	30MG	A071068 001	Nov 25, 1986
	30MG	A072369 001	Mar 30, 1989

FLURBIPROFENTABLET;ORAL  
ANSAID

PHARMACIA AND UPJOHN	50MG	N018766 002	Oct 31, 1988
	100MG	N018766 003	Oct 31, 1988

FLURBIPROFEN

AUROLIFE PHARMA LLC	50MG	A074448 001	Jul 28, 1995
	100MG	A074448 002	Jul 28, 1995
IVAX SUB TEVA PHARMS	50MG	A074411 001	May 31, 1995
	100MG	A074411 002	May 31, 1995
PLIVA	50MG	A074647 001	Apr 01, 1997
	100MG	A074647 002	Apr 01, 1997
TEVA	50MG	A074405 002	May 24, 1995
	100MG	A074405 001	May 24, 1995
THERAGEN	100MG	A074560 002	May 16, 1997

**DISCONTINUED DRUG PRODUCT LIST**

6-162(of 375)

\*\* See List Footnote

FLUTAMIDE

CAPSULE;ORAL			
EULEXIN			
+ SCHERING	125MG		N018554 001 Jan 27, 1989
FLUTAMIDE			
FOSUN PHARMA	125MG		A075818 001 Sep 18, 2001
MYLAN	125MG		A076224 001 May 09, 2003

FLUTICASONE PROPIONATE

AEROSOL, METERED;INHALATION			
FLOVENT			
GLAXOSMITHKLINE	0.044MG/INH		N020548 001 Mar 27, 1996
	0.11MG/INH		N020548 002 Mar 27, 1996
	0.22MG/INH		N020548 003 Mar 27, 1996
CREAM;TOPICAL			
CUTIVATE			
+ FOUGERA PHARMS	0.05% **		N019958 001 Dec 18, 1990
FLUTICASONE PROPIONATE			
NESHER PHARMS	0.05%		A076865 001 Sep 10, 2004
OINTMENT;TOPICAL			
FLUTICASONE PROPIONATE			
TARO PHARM INDNS	0.005%		A077145 001 Jun 14, 2005
POWDER;INHALATION			
FLOVENT			
GLAXOSMITHKLINE	0.044MG/INH		N020549 001 Nov 07, 1997
	0.088MG/INH		N020549 002 Nov 07, 1997
	0.22MG/INH		N020549 003 Nov 07, 1997
SPRAY, METERED;NASAL			
FLONASE			
+ GLAXOSMITHKLINE	0.05MG/SPRAY **		N020121 001 Oct 19, 1994

FLUVASTATIN SODIUM

CAPSULE;ORAL			
LESCOL			
+ NOVARTIS	EQ 20MG BASE		N020261 001 Dec 31, 1993
+	EQ 40MG BASE		N020261 002 Dec 31, 1993

FLUVOXAMINE MALEATE

CAPSULE, EXTENDED RELEASE;ORAL			
LUVOX CR			
+ JAZZ PHARMS	100MG		N022033 001 Feb 28, 2008
+	150MG		N022033 002 Feb 28, 2008
TABLET;ORAL			
FLUVOXAMINE MALEATE			
ACTAVIS ELIZABETH	25MG		A075901 001 Dec 28, 2000
	50MG		A075901 002 Dec 28, 2000
	100MG		A075901 003 Dec 28, 2000
ANI PHARMS INC	25MG		A075898 001 Mar 12, 2001
	50MG		A075898 002 Mar 12, 2001
	100MG		A075898 003 Mar 12, 2001
ECI PHARMS LLC	25MG		A075900 001 Feb 23, 2006
	50MG		A075900 002 Feb 23, 2006
	100MG		A075900 003 Feb 23, 2006
MYLAN	50MG		A075950 001 Oct 15, 2001
	100MG		A075950 002 Oct 15, 2001
SUN PHARM INDUSTRIES	25MG		A076125 001 Apr 29, 2002
	50MG		A076125 002 Apr 29, 2002
	100MG		A076125 003 Apr 29, 2002
SYNTTHON PHARMS	25MG		A075899 001 Jan 17, 2001
	50MG		A075899 002 Jan 17, 2001
	100MG		A075899 003 Jan 17, 2001
UPSHER-SMITH LABS	25MG		A075887 001 Jan 05, 2001
	50MG		A075887 002 Jan 05, 2001
	100MG		A075887 003 Jan 05, 2001
WATSON LABS	25MG		A075894 001 Apr 18, 2001
	50MG		A075894 002 Apr 18, 2001
	100MG		A075894 003 Apr 18, 2001
LUVOX			
+ SOLVAY	25MG **		N020243 001 Dec 05, 1994
+	50MG **		N020243 002 Dec 05, 1994
+	100MG **		N020243 003 Dec 05, 1994

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-163(of 375)

\*\* See List Footnote

FLUVOXAMINE MALEATE

TABLET;ORAL

LUVOX

+

150MG \*\*

N020243 004 Dec 05, 1994

FOLIC ACID

INJECTABLE;INJECTION

FOLIC ACID

BEN VENUE

5MG/ML

A081066 001 Dec 29, 1993

FOLVITE

WYETH PHARMS INC

5MG/ML

N005897 008

TABLET;ORAL

FOLIC ACID

BARR

1MG

A089177 001 Jan 08, 1986

CONTRACT PHARMACAL

1MG

A085061 001

EVERYLIFE

1MG

A080755 001

HALSEY

1MG

A083598 001

IMPAKX LABS

1MG

A080686 001

IVAX SUB TEVA PHARMS

1MG

A083000 001

JUBILANT CADISTA

1MG

A040514 001 Jun 14, 2005

LANNETT

1MG

A080816 001

LILLY

1MG

N006135 003

MK LABS

1MG

A083526 001

NEXGEN PHARMA INC

1MG

A084915 001

PHARMERAL

1MG

A084158 001

PIONEER PHARMS

1MG

A088949 001 Sep 13, 1985

PUREPAC PHARM

1MG

A080784 001

SANDOZ

1MG

A084472 001

SUN PHARM INDUSTRIES

1MG

A040582 001 Jul 18, 2005

TABLICAPS

1MG

A083133 002

UDL

1MG

A088199 001 Mar 29, 1983

USL PHARMA

1MG

A087828 001 May 13, 1982

VALEANT PHARM INTL

1MG

A080903 001

VANGARD

1MG

A088730 001 Mar 23, 1984

VINTAGE PHARMS

1MG

A086296 001

WATSON LABS

1MG

A083141 001

1MG

A085141 002

WHITEWORTH TOWN PLSN

1MG

A080691 002

FOLICET

MISSION PHARMA

1MG

A087438 001

FOLVITE

WYETH PHARMS INC

1MG

N005897 004

FOLLITROPIN ALFA/BETA

INJECTABLE;INTRAMUSCULAR, SUBCUTANEOUS

FOLLISTIM

ORGANON USA INC

75 IU/VIAL

N020582 001 Sep 29, 1997

150 IU/VIAL

N020582 002 Sep 29, 1997

INJECTABLE;SUBCUTANEOUS

FOLLISTIM AQ

ORGANON USA INC

75 IU/0.5ML

N021273 001 Aug 26, 2005

150 IU/0.18ML

N021211 003 Feb 11, 2004

150 IU/0.5ML

N021273 002 Aug 26, 2005

GONAL-F

EMD SERONO

37.5 IU/VIAL

N020378 003 May 25, 2000

37.5 IU/VIAL

N021765 001 Mar 25, 2004

75 IU/VIAL

N020378 001 Sep 29, 1997

150 IU/VIAL

N020378 002 Sep 29, 1997

150 IU/VIAL

N021765 003 Mar 25, 2004

FOMEPIZOLE

INJECTABLE;INJECTION

FOMEPIZOLE

MYLAN INSTITUTIONAL

1.5GM/1.5ML (1GM/ML)

A079033 001 Apr 07, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-164(of 375)

\*\* See List Footnote

FOMIVIRSEN SODIUM

INJECTABLE; INJECTION  
 VITRAVENE PRESERVATIVE FREE  
 NOVARTIS 6.6MG/ML N020961 001 Aug 26, 1998

FORMOTEROL FUMARATE

POWDER; INHALATION  
 FORADIL  
 + NOVARTIS 0.012MG/INH N020831 001 Feb 16, 2001  
 FORADIL CERTIHALER  
 NOVARTIS 0.0085MG/INH N021592 001 Dec 15, 2006

FOSAPREPITANT DIMEGLUMINE

POWDER; INTRAVENOUS  
 EMEND  
 + MERCK AND CO INC EQ 115MG BASE/VIAL \*\* N022023 001 Jan 25, 2008

FOSCARNET SODIUM

INJECTABLE; INJECTION  
 FOSCARNET SODIUM  
 HOSPIRA 2.4GM/100ML A077174 001 May 31, 2005

FOSINOPRIL SODIUM

TABLET; ORAL  
 FOSINOPRIL SODIUM  
 ACTAVIS LABS FL INC 10MG A076620 001 Oct 15, 2004  
 20MG A076620 002 Oct 15, 2004  
 40MG A076620 003 Oct 15, 2004  
 RANBAXY LABS LTD 10MG A076580 001 Apr 23, 2004  
 20MG A076580 002 Apr 23, 2004  
 40MG A076580 003 Apr 23, 2004  
 UPSHER-SMITH LABS 10MG A076188 001 Oct 08, 2004  
 20MG A076188 002 Oct 08, 2004  
 40MG A076188 003 Oct 08, 2004  
 WATSON LABS 10MG A076987 001 Dec 23, 2004  
 20MG A077531 001 Aug 31, 2006  
 20MG A076987 002 Dec 23, 2004  
 40MG A077531 002 Aug 31, 2006  
 40MG A076987 003 Dec 23, 2004  
 40MG A077531 003 Aug 31, 2006  
 MONOPRIL  
 + BRISTOL MYERS SQUIBB 10MG \*\* N019915 002 May 16, 1991  
 + 20MG \*\* N019915 003 May 16, 1991  
 + 40MG \*\* N019915 004 Mar 28, 1995

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL  
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE  
 ACTAVIS LABS FL INC 10MG;12.5MG A076608 001 Dec 03, 2004  
 20MG;12.5MG A076608 002 Dec 03, 2004  
 MYLAN 10MG;12.5MG A077705 001 Aug 14, 2006  
 20MG;12.5MG A077705 002 Aug 14, 2006  
 SUN PHARM INDs LTD 10MG;12.5MG A076739 001 Dec 17, 2004  
 20MG;12.5MG A076739 002 Dec 17, 2004  
 TEVA 10MG;12.5MG A076945 001 Jul 05, 2006  
 20MG;12.5MG A076945 002 Jul 05, 2006  
 WATSON LABS 10MG;12.5MG A077144 001 Aug 16, 2005  
 20MG;12.5MG A077144 002 Aug 16, 2005  
 MONOPRIL-HCT  
 + BRISTOL MYERS SQUIBB 10MG;12.5MG \*\* N020286 002 Nov 30, 1994  
 + 20MG;12.5MG \*\* N020286 001 Nov 30, 1994

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION  
 FOSPHENYTOIN SODIUM  
 APOTEX INC EQ 50MG PHENYTOIN NA/ML A078126 001 Aug 06, 2007  
 HOSPIRA EQ 50MG PHENYTOIN NA/ML A078158 001 Aug 06, 2007  
 TEVA PHARMS USA EQ 50MG PHENYTOIN NA/ML A076886 001 Aug 06, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-165(of 375)

\*\* See List Footnote

**FOSPROPOFOL DISODIUM**SOLUTION; INTRAVENOUS  
LUSEDRA

EISAI INC 1050MG/30ML (35MG/ML) N022244 001 Dec 12, 2008

**FURAZOLIDONE**SUSPENSION; ORAL  
FUROXONESHIRE 50MG/15ML N011323 002  
TABLET; ORAL  
FUROXONE  
SHIRE 100MG N011270 002**FUROSEMIDE**INJECTABLE; INJECTION  
FUROSEMIDE

ABRAXIS PHARM	10MG/ML	N018507 001 Jul 30, 1982
	10MG/ML	N019036 001 Aug 13, 1984
ACCORD HLTHCARE	10MG/ML	A070017 001 Dec 15, 1986
ASTRAZENECA	10MG/ML	A070014 001 Sep 09, 1985
HOSPIRA	10MG/ML	A070578 001 Jul 08, 1987
	10MG/ML	A072080 001 Aug 13, 1991
	10MG/ML	A074337 001 Oct 31, 1994
IGI LABS INC	10MG/ML	A070095 001 Sep 09, 1985
	10MG/ML	A070096 001 Sep 09, 1985
INTL MEDICATION	10MG/ML	N018025 001
+ LUITPOLD	10MG/ML **	N018579 001 Nov 30, 1983
MARSAM PHARMS LLC	10MG/ML	A074017 001 Jun 30, 1994
SMITH AND NEPHEW	10MG/ML	A070023 001 Feb 05, 1986
	10MG/ML	A070078 001 Feb 05, 1986
WARNER CHILCOTT	10MG/ML	N018420 001 Feb 26, 1982
WATSON LABS	10MG/ML	A070019 001 Sep 22, 1986
	10MG/ML	A070604 001 Jan 02, 1987
WEST-WARD PHARMS INT	10MG/ML	A071439 001 Sep 14, 1990
	10MG/ML	N018267 001
WYETH AYERST	10MG/ML	N018670 001 Jul 20, 1982
LASIX		
+ SANOFI AVENTIS US	10MG/ML **	N016363 001
SOLUTION; ORAL		
LASIX		
SANOFI AVENTIS US	10MG/ML	N017688 001
TABLET; ORAL		
FUROSEMIDE		
DAVA PHARMS INC	20MG	N018415 001 Jul 27, 1982
	40MG	N018415 002 Jul 27, 1982
	80MG	N018415 003 Nov 26, 1984
INTL MEDICATION	20MG	N018753 001 Feb 28, 1984
	40MG	N018753 002 Feb 28, 1984
KALAPHARM	20MG	N018868 001 Jun 28, 1983
	40MG	N018868 002 Jun 28, 1983
SANDOZ	40MG	N018750 002 Jul 30, 1984
SUN PHARM INDNS INC	20MG	A091258 001 Apr 01, 2014
	40MG	A091258 002 Apr 01, 2014
	80MG	N018790 001 Nov 29, 1983
SUN PHARM INDUSTRIES	20MG	A091258 003 Apr 01, 2014
	80MG	A070043 001 Sep 26, 1985
	80MG	A070100 001 Jan 26, 1988
SUPERPHARM	20MG	N018370 002 Jun 26, 1984
	40MG	N018370 001 Feb 10, 1983
WARNER CHILCOTT	20MG	N018419 001 Jan 31, 1983
	40MG	N018419 002 Jan 31, 1983
	80MG	N018419 003 Nov 13, 1984
WATSON LABS	20MG	A070412 001 Feb 26, 1986
	20MG	A071379 001 Jan 02, 1987
	20MG	N018369 001 May 14, 1982
	40MG	A070413 001 Feb 26, 1986
	40MG	A070450 001 Nov 22, 1985
	40MG	N018369 002 May 14, 1982
	80MG	A071594 001 Feb 09, 1988
WATSON LABS TEVA	20MG	A070449 001 Nov 22, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-166(of 375)

\*\* See List Footnote

FUROSEMIDETABLET;ORAL  
FUROSEMIDE

80MG

A070528 001 Jan 07, 1986

GABAPENTINCAPSULE;ORAL  
GABAPENTIN

CSPC OUYI PHARM CO	100MG	A075477 001 Mar 23, 2005
	300MG	A075477 002 Mar 23, 2005
	400MG	A075477 003 Mar 23, 2005
HIKMA	100MG	A078150 001 Sep 25, 2007
	300MG	A078150 002 Sep 25, 2007
	400MG	A078150 003 Sep 25, 2007
SANDOZ	100MG	A075428 001 Jan 24, 2006
	100MG	A075539 001 Apr 06, 2005
	300MG	A075428 002 Jan 24, 2006
	300MG	A075539 002 Apr 06, 2005
	400MG	A075428 003 Jan 24, 2006
	400MG	A075539 003 Apr 06, 2005
SUN PHARM INDS LTD	100MG	A076606 001 Oct 07, 2005
	300MG	A076606 002 Oct 07, 2005
	400MG	A076606 003 Oct 07, 2005
SUN PHARM INDUSTRIES	100MG	A076537 001 Jun 30, 2005
	300MG	A076537 002 Jun 30, 2005
	400MG	A076537 003 Jun 30, 2005
WATSON LABS	100MG	A075485 003 May 11, 2007
	300MG	A075485 002 May 11, 2007
	400MG	A075485 001 May 11, 2007

TABLET;ORAL

GABAPENTIN

HIKMA PHARMS	600MG	A078782 001 Jul 21, 2011
	800MG	A078782 002 Jul 21, 2011
RANBAXY	600MG	A076605 001 Sep 14, 2005
	800MG	A076605 002 Sep 14, 2005
SANDOZ	600MG	A076120 001 Jan 27, 2006
	600MG	A076877 001 Jul 06, 2006
	800MG	A076120 002 Jan 27, 2006
	800MG	A076877 002 Jul 06, 2006
TEVA	600MG	A075827 001 Dec 15, 2004
	800MG	A075827 002 Dec 15, 2004

GADODIAMIDE

INJECTABLE; INJECTION

OMNISCAN

GE HEALTHCARE	14.35GM/50ML (287MG/ML)	N022066 001 Sep 05, 2007
---------------	-------------------------	--------------------------

GADOFOSEVESET TRISODIUM

SOLUTION; INTRAVENOUS

ABLAVAR

LANTHEUS MEDCL	2440MG/10ML (244MG/ML)	N021711 001 Dec 22, 2008
	3660MG/15ML (244MG/ML)	N021711 002 Dec 22, 2008

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

GALANTAMINE HYDROBROMIDE

IMPAX LABS	EQ 8MG BASE	A078484 001 May 27, 2009
	EQ 16MG BASE	A078484 002 May 27, 2009
	EQ 24MG BASE	A078484 003 May 27, 2009

SOLUTION; ORAL

RAZADYNE

JANSSEN PHARMS	4MG/ML **	N021224 001 Jun 22, 2001
----------------	-----------	--------------------------

TABLET; ORAL

GALANTAMINE HYDROBROMIDE

ACTAVIS ELIZABETH	EQ 4MG BASE	A077585 001 Sep 15, 2009
	EQ 8MG BASE	A077585 002 Sep 15, 2009
	EQ 12MG BASE	A077585 003 Sep 15, 2009
MYLAN	EQ 4MG BASE	A077603 001 Aug 28, 2008
	EQ 8MG BASE	A077603 002 Aug 28, 2008
	EQ 12MG BASE	A077603 003 Aug 28, 2008

**DISCONTINUED DRUG PRODUCT LIST**

6-167(of 375)

\*\* See List Footnote

GALLAMINE TRIETHIODIDE

INJECTABLE; INJECTION

FLAXEDIL

DAVIS AND GECK

20MG/ML

N007842 001

100MG/ML

N007842 002

GALLIUM CITRATE GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

GE HEALTHCARE

1mCi/ML

N017700 001

NEOSCAN

GE HEALTHCARE

2mCi/ML

N017655 001

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

CHAPTER 7 TRUSTEE

25MG/ML \*\*

N019961 002 Jan 17, 1991

GANCICLOVIR

CAPSULE; ORAL

CYTOVENE

+ ROCHE PALO

250MG \*\*

N020460 001 Dec 22, 1994

+

500MG \*\*

N020460 002 Dec 12, 1997

GANCICLOVIR

RANBAXY LABS LTD

250MG

A076457 001 Jun 27, 2003

500MG

A076457 002 Jun 27, 2003

IMPLANT; IMPLANTATION

VITRASERT

BAUSCH AND LOMB

4.5MG

N020569 001 Mar 04, 1996

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

GANCICLOVIR SODIUM

WEST-WARD PHARMS INT EQ 500MG BASE/VIAL

A076222 001 Jul 16, 2003

GATIFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

GATIFLOXACIN

APOTEX INC

0.3%

A079084 001 Aug 19, 2011

GEFITINIB

TABLET; ORAL

IRESSA

ASTRAZENECA

250MG

N021399 001 May 05, 2003

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION

GEMCITABINE HYDROCHLORIDE

SAGENT PHARMS

EQ 200MG BASE/VIAL

A091597 001 May 07, 2013

EQ 1GM BASE/VIAL

A091597 002 May 07, 2013

GEMFIBROZIL

CAPSULE; ORAL

GEMFIBROZIL

MYLAN

300MG

A073466 001 Jan 25, 1993

PUREPAC PHARM

300MG

A072929 001 Jan 29, 1993

LOPID

PFIZER PHARMS

200MG

N018422 001

300MG

N018422 002

TABLET; ORAL

GEMFIBROZIL

FOSUN PHARMA

600MG

A074615 001 Sep 29, 1995

MYLAN

600MG

A074452 001 Feb 16, 1995

PUREPAC PHARM

600MG

A074360 001 Aug 31, 1994

WATSON LABS

600MG

A074156 001 Oct 24, 1994

600MG

A074442 001 Apr 28, 1995

GEMTUZUMAB OZOGAMICIN

INJECTABLE; INJECTION

MYLOTARG

WYETH PHARMS INC

5MG/VIAL

N021174 001 May 17, 2000

**DISCONTINUED DRUG PRODUCT LIST**

6-168(of 375)

\*\* See List Footnote

GENTAMICIN SULFATE

CREAM;TOPICAL

GARAMYCIN

SCHERING

EQ 0.1% BASE \*\*

A060462 001

GENTAFAIR

PHARMAFAIR

EQ 0.1% BASE

A062458 001 Sep 01, 1983

GENTAMICIN SULFATE

ALPHARMA US PHARMS

EQ 0.1% BASE

A062471 001 Sep 27, 1983

FOUGERA PHARMS INC

EQ 0.1% BASE

A062531 001 Jul 05, 1984

PHARMADERM

EQ 1MG BASE/GM

A062530 001 Jul 05, 1984

TARO

EQ 0.1% BASE

A062427 001 May 26, 1983

INJECTABLE; INJECTION

APOGEN

KING PHARMS

EQ 10MG BASE/ML

A062289 001

EQ 40MG BASE/ML

A062289 002

BRISTAGEN

BRISTOL

EQ 40MG BASE/ML

A062288 001

GARAMYCIN

SCHERING

EQ 1MG BASE/ML \*\*

A061716 002

EQ 10MG BASE/ML \*\*

A061739 001

EQ 40MG BASE/ML \*\*

A061716 001

GENTAFAIR

PHARMAFAIR

EQ 40MG BASE/ML

A062493 001 Aug 28, 1985

GENTAMICIN

INTL MEDICATION

EQ 1MG BASE/ML

A062325 003 Jun 23, 1982

EQ 40MG BASE/ML

A062325 001

EQ 100MG BASE/100ML

A062325 004 Jun 23, 1982

GENTAMICIN SULFATE

ABBOTT

EQ 1.2MG BASE/ML

A062413 001 Aug 11, 1983

EQ 1.4MG BASE/ML

A062413 002 Aug 11, 1983

EQ 1.6MG BASE/ML

A062413 003 Aug 11, 1983

EQ 1.8MG BASE/ML

A062413 004 Aug 11, 1983

EQ 2MG BASE/ML

A062413 005 Aug 11, 1983

EQ 60MG BASE/100ML

A062413 006 Aug 11, 1983

EQ 70MG BASE/100ML

A062413 007 Aug 11, 1983

EQ 80MG BASE/100ML

A062413 008 Aug 11, 1983

EQ 90MG BASE/100ML

A062413 009 Aug 11, 1983

EQ 100MG BASE/100ML

A062413 010 Aug 11, 1983

FRESENIUS KABI USA

EQ 10MG BASE/ML

A062356 001 Mar 04, 1982

EQ 40MG BASE/ML

A062356 002 Mar 04, 1982

KALAPHARM

EQ 40MG BASE/ML

A062354 001 Apr 05, 1982

PHARM SPEC

EQ 40MG BASE/ML

A062340 001 Mar 28, 1983

SOLOPAK

EQ 10MG BASE/ML

A062507 001 Jun 06, 1985

EQ 40MG BASE/ML

A062507 002 Jun 06, 1985

TEVA PARENTERAL

EQ 10MG BASE/ML

A063149 001 Nov 21, 1991

EQ 40MG BASE/ML

A063106 002 Nov 21, 1991

WATSON LABS

EQ 10MG BASE/ML

A062318 002

EQ 40MG BASE/ML

A062318 001

WEST-WARD PHARMS INT

EQ 10MG BASE/ML

A062251 002

EQ 40MG BASE/ML

A062251 001

WYETH AYERST

EQ 10MG BASE/ML

A062264 001

EQ 40MG BASE/ML

A062264 002

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN

EQ 0.8MG BASE/ML

A062814 001 Aug 28, 1987

EQ 1.2MG BASE/ML

A062814 002 Aug 28, 1987

EQ 1.4MG BASE/ML

A062814 003 Aug 28, 1987

EQ 1.6MG BASE/ML

A062814 004 Aug 28, 1987

EQ 1.8MG BASE/ML

A062814 005 Aug 28, 1987

EQ 2MG BASE/ML

A062814 006 Aug 28, 1987

EQ 2.4MG BASE/ML

A062814 007 Aug 28, 1987

EQ 40MG BASE/100ML

A062814 008 Aug 28, 1987

EQ 60MG BASE/100ML

A062814 009 Aug 28, 1987

EQ 70MG BASE/100ML

A062814 010 Aug 28, 1987

EQ 80MG BASE/100ML

A062814 011 Aug 28, 1987

EQ 90MG BASE/100ML

A062814 012 Aug 28, 1987

EQ 100MG BASE/100ML

A062814 013 Aug 28, 1987

EQ 120MG BASE/100ML

A062814 014 Aug 28, 1987

BAXTER HLTHCARE

EQ 0.8MG BASE/ML

A062373 001 Sep 07, 1982

EQ 2.4MG BASE/ML

A062373 010 Sep 07, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-169(of 375)

\*\* See List Footnote

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	EQ 40MG BASE/100ML	A062373 003 Sep 07, 1982
	EQ 60MG BASE/100ML	A062373 004 Sep 07, 1982
HOSPIRA	EQ 1.2MG BASE/ML	A062588 001 Jan 06, 1986
	EQ 1.4MG BASE/ML	A062414 002 Aug 15, 1983
	EQ 1.4MG BASE/ML	A062588 002 Jan 06, 1986
	EQ 1.6MG BASE/ML	A062588 003 Jan 06, 1986
	EQ 1.8MG BASE/ML	A062414 004 Aug 15, 1983
	EQ 1.8MG BASE/ML	A062588 004 Jan 06, 1986
	EQ 2MG BASE/ML	A062414 005 Aug 15, 1983
	EQ 2MG BASE/ML	A062588 005 Jan 06, 1986
	EQ 60MG BASE/100ML	A062414 006 Aug 15, 1983
	EQ 60MG BASE/100ML	A062588 006 Jan 06, 1986
	EQ 70MG BASE/100ML	A062414 007 Aug 15, 1983
	EQ 70MG BASE/100ML	A062588 007 Jan 06, 1986
	EQ 80MG BASE/100ML	A062588 008 Jan 06, 1986
	EQ 90MG BASE/100ML	A062414 009 Aug 15, 1983
	EQ 90MG BASE/100ML	A062588 009 Jan 06, 1986
	EQ 100MG BASE/100ML	A062588 010 Jan 06, 1986

U-GENCIN

PHARMACIA AND UPJOHN	EQ 10MG BASE/ML	A062248 001
	EQ 40MG BASE/ML	A062248 002

INJECTABLE; INTRATHECAL

GARAMYCIN

+ SCHERING	EQ 2MG BASE/ML **	N050505 001
------------	-------------------	-------------

OINTMENT; OPHTHALMIC

GARAMYCIN

SCHERING	EQ 0.3% BASE	N050425 001
----------	--------------	-------------

GENTACIDIN

NOVARTIS	EQ 0.3% BASE	A062501 001 Jul 26, 1984
----------	--------------	--------------------------

GENTAFAIR

PHARMAFAIR	EQ 3MG BASE/GM	A062443 001 May 26, 1983
------------	----------------	--------------------------

OINTMENT; TOPICAL

GARAMYCIN

SCHERING	EQ 0.1% BASE **	A060463 001
----------	-----------------	-------------

GENTAFAIR

PHARMAFAIR	EQ 0.1% BASE	A062444 001 May 26, 1983
------------	--------------	--------------------------

GENTAMICIN SULFATE

ALPHARMA US PHARMS	EQ 0.1% BASE	A062496 001 Mar 14, 1984
G AND W LABS INC	EQ 0.1% BASE	A064054 001 Apr 29, 1994
PHARMADERM	EQ 0.1% BASE	A062534 001 Oct 10, 1984

SOLUTION/DROPS; OPHTHALMIC

GARAMYCIN

+ SCHERING	EQ 0.3% BASE **	N050039 002
------------	-----------------	-------------

GENTACIDIN

NOVARTIS	EQ 0.3% BASE	A062480 001 Mar 30, 1984
----------	--------------	--------------------------

GENTAFAIR

PHARMAFAIR	EQ 0.3% BASE	A062440 001 May 03, 1983
------------	--------------	--------------------------

GENTAMICIN SULFATE

ALCON PHARMS LTD	EQ 0.3% BASE	A062523 001 Nov 25, 1985
PACO	EQ 3MG BASE/ML	A062932 001 Nov 07, 1988

GENTIAN VIOLET

SUPPOSITORY; VAGINAL

GVS

SAVAGE LABS	0.4%	A083513 001
-------------	------	-------------

TAMPON; VAGINAL

GENAPAX

KEY PHARMS	5MG	A085017 001
------------	-----	-------------

GLATIRAMER ACETATE

FOR SOLUTION; SUBCUTANEOUS

COPAXONE

TEVA PHARMS USA	20MG/VIAL	N020622 001 Dec 20, 1996
-----------------	-----------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-170(of 375)

\*\* See List Footnote

**GLIMEPIRIDE**

TABLET;ORAL

GLIMEPIRIDE

ACTAVIS LABS FL INC	1MG	A076995 001	Apr 27, 2010
	2MG	A076995 002	Apr 27, 2010
	4MG	A076995 003	Apr 27, 2010
EPIC PHARMA LLC	1MG	A077274 001	Oct 06, 2005
	2MG	A077274 002	Oct 06, 2005
	4MG	A077274 003	Oct 06, 2005
HIKMA PHARMS	1MG	A078952 001	Aug 01, 2013
	2MG	A078952 002	Aug 01, 2013
	4MG	A078952 003	Aug 01, 2013
MYLAN	1MG	A077486 001	Feb 10, 2006
	2MG	A077486 002	Feb 10, 2006
	4MG	A077486 003	Feb 10, 2006
RANBAXY	3MG	A077366 001	Oct 06, 2005
	6MG	A077366 002	Oct 06, 2005
RANBAXY LABS LTD	1MG	A076875 001	Oct 06, 2005
	2MG	A076875 002	Oct 06, 2005
	4MG	A076875 003	Oct 06, 2005
	8MG	A076875 004	Oct 06, 2005
WATSON LABS	1MG	A077280 001	Feb 03, 2006
	2MG	A077280 002	Feb 03, 2006
	4MG	A077280 003	Feb 03, 2006

**GLIMEPIRIDE; ROSIGLITAZONE MALEATE**

TABLET;ORAL

AVANDARYL

SB PHARMCO	1MG;4MG **	N021700 001	Nov 23, 2005
+	2MG;4MG **	N021700 002	Nov 23, 2005
+	2MG;8MG **	N021700 004	Mar 30, 2007
+	4MG;4MG **	N021700 003	Nov 23, 2005
+	4MG;8MG **	N021700 005	Mar 30, 2007

**GLIPIZIDE**

TABLET;ORAL

GLIPIZIDE

ANI PHARMS INC	5MG	A074387 001	Mar 04, 1996
	10MG	A074387 002	Mar 04, 1996
BARR LABS INC	5MG	A074619 001	Apr 04, 1997
	10MG	A074619 002	Apr 04, 1997
MYLAN	5MG	A074438 001	Jun 20, 1995
	10MG	A074438 002	Jun 20, 1995
SANDOZ	5MG	A074542 001	Jun 20, 1995
	10MG	A074542 002	Jun 20, 1995
VINTAGE PHARMS LLC	5MG	A074378 001	Nov 28, 1994
	10MG	A074378 002	Nov 28, 1994
WATSON LABS	5MG	A074370 001	Nov 22, 1994
	10MG	A074370 002	Nov 22, 1994

GLUCOTROL

PFIZER	2.5MG	N017783 003	May 11, 1993
--------	-------	-------------	--------------

**GLIPIZIDE; METFORMIN HYDROCHLORIDE**

TABLET;ORAL

METAGLIP

BRISTOL MYERS SQUIBB	2.5MG;250MG **	N021460 001	Oct 21, 2002
+	2.5MG;500MG **	N021460 002	Oct 21, 2002
+	5MG;500MG **	N021460 003	Oct 21, 2002

**GLUCAGON HYDROCHLORIDE**

INJECTABLE;INJECTION

GLUCAGON

LILLY	EQ 1MG BASE/VIAL **	N012122 001
+	EQ 10MG BASE/VIAL **	N012122 002

**GLUTETHIMIDE**

CAPSULE;ORAL

DORIDEN

SANOFI AVENTIS US	500MG	N009519 008
-------------------	-------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-171(of 375)

\*\* See List Footnote

**GLUTETHIMIDE**TABLET;ORAL  
DORIDEN

SANOFI AVENTIS US	250MG	N009519 002
	500MG	N009519 005
GLUTETHIMIDE		
HALSEY	250MG	A089458 001 Oct 10, 1986
	500MG	A089459 001 Oct 10, 1986
LANNETT	250MG	A083475 001
	500MG	A085571 001
UCB INC	500MG	A085171 001
UPSHER-SMITH LABS	500MG	A083234 002
VITARINE	500MG	A087297 001
WATSON LABS	500MG	A084362 001
	500MG	A085763 001

**GLYBURIDE**TABLET;ORAL  
GLYBURIDE

ACTAVIS ELIZABETH	1.5MG	A075947 001 Nov 14, 2002
	3MG	A075947 002 Nov 14, 2002
	6MG	A075947 003 Nov 14, 2002
GLYBURIDE (MICRONIZED)		
FOSUN PHARMA	1.5MG	A075174 001 Jun 22, 1998
	3MG	A075174 002 Jun 22, 1998
SANOFI AVENTIS US	1.5MG	N020055 001 Apr 17, 1992
	3MG	N020055 002 Apr 17, 1992
	6MG	N020055 003 Mar 08, 2000
GLYNASE		
+ PHARMACIA AND UPJOHN	4.5MG **	N020051 003 Sep 24, 1993
MICRONASE		
+ PHARMACIA AND UPJOHN	1.25MG **	N017498 001 May 01, 1984
	2.5MG	N017498 002 May 01, 1984
+	5MG **	N017498 003 May 01, 1984

**GLYBURIDE; METFORMIN HYDROCHLORIDE**TABLET;ORAL  
GLUCOVANCE

+ BRISTOL MYERS SQUIBB	1.25MG;250MG **	N021178 001 Jul 31, 2000
+	2.5MG;500MG	N021178 002 Jul 31, 2000
+	5MG;500MG	N021178 003 Jul 31, 2000
GLYBURIDE AND METFORMIN HYDROCHLORIDE		
IMPAX LABS INC	1.25MG;250MG	A076731 001 Nov 19, 2004
	2.5MG;500MG	A076731 002 Nov 19, 2004
	5MG;500MG	A076731 003 Nov 19, 2004
TEVA	1.25MG;250MG	A076821 001 Jan 27, 2005
	2.5MG;500MG	A076821 002 Jan 27, 2005
	5MG;500MG	A076821 003 Jan 27, 2005

**GLYCINE**

SOLUTION;IRRIGATION

GLYCINE 1.5% IN PLASTIC CONTAINER		
BAXTER HLTHCARE	1.5GM/100ML	N018522 001 Feb 19, 1982
HOSPIRA	1.5GM/100ML	N017633 001

**GLYCOPYRROLATE**

INJECTABLE;INJECTION

GLYCOPYRROLATE		
ABRAXIS PHARM	0.2MG/ML	A088475 001 Jun 12, 1984
HOSPIRA	0.2MG/ML	A089393 001 Jun 15, 1988
TEVA PARENTERAL	0.2MG/ML	A081169 001 Sep 10, 1991
WATSON LABS	0.2MG/ML	A086947 001 Jun 24, 1983
ROBINUL		
ROBINS AH	0.2MG/ML	N014764 001
+ WEST-WARD PHARMS INT	0.2MG/ML **	N017558 001

TABLET;ORAL

GLYCOPYRROLATE		
EPIC PHARMA LLC	1MG	A040568 001 Dec 22, 2004
	2MG	A040568 002 Dec 22, 2004
HIKMA INTL PHARMS	1MG	A040836 001 Mar 05, 2009
	2MG	A040836 002 Mar 05, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-172(of 375)

\*\* See List Footnote

GLYCOPYRROLATE

TABLET;ORAL

## GLYCOPYRROLATE

WATSON LABS

1MG  
1MG  
2MG  
2MG  
2MGA085562 001  
A086902 001  
A085563 001  
A086178 001  
A086900 001GONADORELIN ACETATE

INJECTABLE;INJECTION

## LUTREPULSE KIT

FERRING

0.8MG/VIAL  
3.2MG/VIALN019687 001 Oct 10, 1989  
N019687 002 Oct 10, 1989GONADORELIN HYDROCHLORIDE

INJECTABLE;INJECTION

## FACTREL

WEST-WARD PHARMS INT

EQ 0.1MG BASE/VIAL  
EQ 0.2MG BASE/VIAL  
EQ 0.5MG BASE/VIALN018123 001 Sep 30, 1982  
N018123 002 Sep 30, 1982  
N018123 003 Sep 30, 1982GONADOTROPIN, CHORIONIC

INJECTABLE;INJECTION

## A.P.L.

FERRING

5,000 UNITS/VIAL  
10,000 UNITS/VIAL  
20,000 UNITS/VIALN017055 001  
N017055 002  
N017055 003

## CHORIONIC GONADOTROPIN

BEL MAR

5,000 UNITS/VIAL  
10,000 UNITS/VIALN017054 001  
N017054 002

FERRING

2,000 UNITS/VIAL  
2,000 UNITS/VIAL  
15,000 UNITS/VIALN017016 009 Dec 27, 1984  
N017016 011 Feb 16, 1990  
N017016 010 Feb 15, 1985

FRESENIUS KABI USA

20,000 UNITS/VIAL  
5,000 UNITS/VIAL  
15,000 UNITS/VIALN017016 004  
N017067 001  
N017067 004

FRESENIUS KABI USA

20,000 UNITS/VIAL

N017067 003

## FOLLUTEIN

BRISTOL MYERS SQUIBB

10,000 UNITS/VIAL

N017056 001

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OPHTHALMIC

## NEO-POLYCYN

DOW PHARM

0.025MG/ML;EQ 1.75MG BASE/ML;10,000  
UNITS/ML

A060427 001

## NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

IPHARM

0.025MG/ML;EQ 1.75MG BASE/ML;10,000  
UNITS/ML

A062818 001 Oct 11, 1988

WATSON LABS

0.025MG/ML;EQ 1.75MG BASE/ML;10,000  
UNITS/ML

A062788 001 Jun 11, 1987

## NEOMYCIN SULFATE AND POLYMYXIN B SULFATE GRAMICIDIN

PHARMAFAIR

0.025MG/ML;EQ 1.75MG BASE/ML;10,000  
UNITS/ML

A062383 001 Aug 31, 1982

GRANISETRON HYDROCHLORIDE

INJECTABLE;INJECTION

## GRANISETRON HYDROCHLORIDE

BAXTER HLTHCARE CORP

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)  
EQ 1MG BASE/ML (EQ 1MG BASE/ML)  
EQ 4MG BASE/4ML (EQ 1MG BASE/ML)A078197 001 Dec 31, 2007  
A078198 001 Jun 30, 2008  
A078198 002 Jun 30, 2008  
A078808 001 Apr 29, 2008

SANDOZ INC

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)

## GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE

TEVA PHARMS USA

EQ 1MG BASE/ML (EQ 1MG BASE/ML)

A077165 001 Dec 31, 2007

## KYTRIL

+ ROCHE

EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML) \*\*  
EQ 1MG BASE/ML (EQ 1MG BASE/ML) \*\*  
EQ 3MG BASE/ML \*\*  
EQ 4MG BASE/4ML (EQ 1MG BASE/ML) \*\*N020239 003 Sep 17, 2004  
N020239 004 Mar 11, 1994  
N020239 001 Dec 29, 1993  
N020239 002 Mar 11, 1994

## SOLUTION;ORAL

## GRANISOL

PEDIATRIX

EQ 2MG BASE/10ML

A078334 001 Feb 28, 2008

**DISCONTINUED DRUG PRODUCT LIST**

6-173(of 375)

\*\* See List Footnote

GRANISETRON HYDROCHLORIDE

SOLUTION;ORAL	KYTRIL	+ ROCHE	EQ 2MG BASE/10ML **	N021238 001 Jun 27, 2001
TABLET;ORAL	GRANISETRON HYDROCHLORIDE	BARR	EQ 1MG BASE	A078221 001 Dec 31, 2007
		EPIC PHARMA LLC	EQ 1MG BASE	A078260 001 Dec 31, 2007
	KYTRIL	+ ROCHE	EQ 1MG BASE **	N020305 001 Mar 16, 1995
		+	EQ 2MG BASE **	N020305 002 Jun 15, 1998

GREPAFLOXACIN HYDROCHLORIDE

TABLET;ORAL	RAXAR	OTSUKA	EQ 200MG BASE	N020695 001 Nov 06, 1997
			EQ 400MG BASE	N020695 002 May 14, 1998
			EQ 600MG BASE	N020695 003 May 14, 1998

GRISEOFULVIN, MICROCRYSTALLINE

CAPSULE;ORAL	GRISACTIN	WYETH AYERST	125MG	N050051 002
			250MG	N050051 001

SUSPENSION;ORAL

GRIFULVIN V	+ JOHNSON AND JOHNSON	125MG/5ML **	N050448 001
-------------	-----------------------	--------------	-------------

TABLET;ORAL

FULVICIN-U/F	CHARTWELL RX	250MG	A060569 002
		500MG	A060569 001

GRIFULVIN V

J AND J	125MG	A060618 001
	250MG	A060618 002
	500MG	A060618 003
VALEANT LUXEMBOURG	125MG	A062279 001
	250MG **	A062279 002

GRISACTIN

WYETH AYERST	500MG	A060212 001
--------------	-------	-------------

GRISEOFULVIN, MICROSIZE

SUSPENSION;ORAL	GRIFULVIN V	VALEANT LUXEMBOURG	125MG/5ML **	A062483 001 Jan 26, 1984
TABLET;ORAL	GRIFULVIN V	VALEANT LUXEMBOURG	500MG	A062279 003

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET;ORAL	FULVICIN P/G	CHARTWELL RX	125MG	A061996 001
			250MG	A061996 002
FULVICIN P/G 165				
CHARTWELL RX			165MG	A061996 003 Apr 06, 1982
FULVICIN P/G 330				
CHARTWELL RX			330MG	A061996 004 Apr 06, 1982
GRISACTIN ULTRA				
WYETH AYERST			125MG	A062178 001
			165MG	A062438 001 Nov 17, 1983
			250MG	A062178 002
			330MG	A062438 002 Nov 17, 1983
ULTRAGRIS-165				
PLIVA			165MG	A062645 001 Jun 30, 1992
ULTRAGRIS-330				
PLIVA			330MG	A062646 001 Jun 30, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-174(of 375)

\*\* See List Footnote

GUAIFENESIN; HYDROCODONE BITARTRATE

SOLUTION;ORAL

OBREDON

+ SOVEREIGN PHARMS

200MG/5ML;2.5MG/5ML

N205474 001 Nov 14, 2014

GUANABENZ ACETATE

TABLET;ORAL

GUANABENZ ACETATE

ANI PHARMS INC

EQ 4MG BASE

A074267 001 Jun 01, 1994

EQ 8MG BASE

A074267 002 Jun 01, 1994

SANDOZ

EQ 4MG BASE

A074517 001 Sep 30, 1998

EQ 8MG BASE

A074517 002 Sep 30, 1998

WATSON LABS

EQ 4MG BASE

A074025 001 Feb 28, 1994

EQ 8MG BASE

A074025 002 Feb 28, 1994

WYTENSIN

WYETH AYERST

EQ 4MG BASE

N018587 001 Sep 07, 1982

EQ 8MG BASE

N018587 002 Sep 07, 1982

EQ 16MG BASE

N018587 003 Sep 07, 1982

GUANADREL SULFATE

TABLET;ORAL

HYLOREL

PHARMACIA AND UPJOHN 10MG

25MG

N018104 001 Dec 29, 1982

N018104 002 Dec 29, 1982

GUANETHIDINE MONOSULFATE

TABLET;ORAL

GUANETHIDINE MONOSULFATE

WATSON LABS

EQ 10MG SULFATE

A086113 001 Mar 26, 1985

EQ 25MG SULFATE

A086114 001 Mar 26, 1985

ISMELIN

NOVARTIS

EQ 10MG SULFATE

N012329 001

EQ 25MG SULFATE

N012329 002

GUANETHIDINE MONOSULFATE; HYDROCHLOROTHIAZIDE

TABLET;ORAL

ESIMIL

NOVARTIS

10MG;25MG

N013553 001

GUANFACINE HYDROCHLORIDE

TABLET;ORAL

GUANFACINE HYDROCHLORIDE

WATSON LABS

EQ 1MG BASE

A074762 001 Jun 25, 1997

EQ 2MG BASE

A074762 002 Jun 25, 1997

TENEX

PROMIUS PHARMA

EQ 3MG BASE

N019032 003 Nov 07, 1988

TABLET, EXTENDED RELEASE;ORAL

GUANFACINE HYDROCHLORIDE

IMPAX LABS INC

EQ 1MG BASE

A202238 001 Oct 20, 2015

EQ 2MG BASE

A202238 002 Oct 20, 2015

EQ 3MG BASE

A202238 003 Oct 20, 2015

EQ 4MG BASE

A202238 004 Oct 20, 2015

HALAZEPAM

TABLET;ORAL

PAXIPAM

SCHERING

20MG

N017736 003

40MG

N017736 004

HALCINONIDE

CREAM;TOPICAL

HALOG

WESTWOOD SQUIBB

0.025%

N017818 001

HALOG-E

SUN PHARM INDs INC

0.1%

N018234 001

OINTMENT;TOPICAL

HALOG

BRISTOL MYERS SQUIBB

0.025%

N018125 001

SOLUTION;TOPICAL

HALOG

SUN PHARM INDs INC

0.1%

N017823 001

**DISCONTINUED DRUG PRODUCT LIST**

6-175(of 375)

\*\* See List Footnote

HALOBETASOL PROPIONATE

OINTMENT;TOPICAL

HALOBETASOL PROPIONATE

FOUGERA PHARMS

0.05%

A076903 001 Dec 16, 2004

HALOFANTRINE HYDROCHLORIDE

TABLET;ORAL

HALFAN

GLAXOSMITHKLINE

250MG

N020250 001 Jul 24, 1992

HALOPERIDOL

TABLET;ORAL

HALDOL

+ ORTHO MCNEIL

0.5MG \*\*

N015921 001

+ 1MG \*\*

N015921 002

+ 2MG \*\*

N015921 003

+ 5MG \*\*

N015921 004

+ 10MG \*\*

N015921 005

+ 20MG \*\*

N015921 006 Feb 02, 1982

HALDOL SOLUTAB

ORTHO MCNEIL PHARM

1MG

N017079 001

HALOPERIDOL

ANDA REPOSITORY

0.5MG

A071156 001 Jan 02, 1987

1MG

A071157 001 Jan 02, 1987

2MG

A071172 001 Jan 02, 1987

5MG

A071212 001 Jan 07, 1988

10MG

A071173 001 Jan 07, 1988

20MG

A071177 001 Jan 07, 1988

CYCLE PHARMS LTD

0.5MG

A071128 001 Feb 17, 1987

1MG

A071129 001 Feb 17, 1987

2MG

A071130 001 Feb 17, 1987

5MG

A071131 001 Feb 17, 1987

10MG

A071132 001 May 12, 1987

20MG

A071133 001 May 12, 1987

DURAMED PHARMS BARR

0.5MG

A071216 001 Dec 04, 1986

1MG

A071217 001 Dec 04, 1986

2MG

A071218 001 Dec 04, 1986

5MG

A071219 001 Dec 04, 1986

10MG

A071220 001 Jul 07, 1987

20MG

A071221 001 Jul 07, 1987

LEDERLE

0.5MG

A072727 001 Sep 19, 1989

1MG

A072728 001 Sep 19, 1989

2MG

A072729 001 Sep 19, 1989

5MG

A072730 001 Sep 19, 1989

10MG

A072731 001 Sep 19, 1989

20MG

A072732 001 Sep 19, 1989

PAR PHARM

20MG

A071328 001 Jul 20, 1987

PUREPAC PHARM

0.5MG

A071071 001 Nov 03, 1986

1MG

A071072 001 Nov 03, 1986

2MG

A071073 001 Nov 03, 1986

5MG

A071074 001 Nov 03, 1986

10MG

A071075 001 Aug 04, 1987

20MG

A071076 001 Aug 04, 1987

QUANTUM PHARMS

0.5MG

A071255 001 Feb 17, 1987

1MG

A071269 001 Feb 17, 1987

2MG

A071256 001 Feb 17, 1987

5MG

A071257 001 Feb 17, 1987

ROYCE LABS

0.5MG

A071722 001 Dec 24, 1987

1MG

A071723 001 Dec 24, 1987

2MG

A071724 001 Dec 24, 1987

5MG

A071725 001 Dec 24, 1987

10MG

A072121 001 Dec 24, 1987

20MG

A072122 001 Dec 24, 1987

SCS

0.5MG

A070720 001 Jun 10, 1986

1MG

A070721 001 Jun 10, 1986

2MG

A070722 001 Jun 10, 1986

5MG

A070723 001 Jun 10, 1986

10MG

A070724 001 Jun 10, 1986

20MG

A070725 001 Sep 24, 1986

VINTAGE

0.5MG

A071235 002 Nov 03, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-176(of 375)

\*\* See List Footnote

HALOPERIDOLTABLET;ORAL  
HALOPERIDOL

	1MG	A071235 003	Nov 03, 1986
	2MG	A071235 001	Nov 03, 1986
	5MG	A071235 004	Nov 03, 1986
	10MG	A071235 005	Jul 20, 1987
WATSON LABS	0.5MG	A070981 001	Mar 06, 1987
	0.5MG	A071571 001	Jun 03, 1988
	1MG	A070982 001	Mar 06, 1987
	1MG	A071572 001	Jun 03, 1988
	2MG	A070983 001	Mar 06, 1987
	2MG	A071573 001	Jun 03, 1988
	5MG	A070984 001	Mar 06, 1987
	5MG	A071374 001	Jun 03, 1988
	10MG	A071375 001	Jun 03, 1988
	10MG	A072113 001	Aug 27, 1991
	20MG	A071376 001	Jun 03, 1988
	20MG	A072353 001	Aug 27, 1991

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

## HALOPERIDOL DECANOATE

HOSPIRA	EQ 50MG BASE/ML	A075176 001	Feb 09, 2000
	EQ 100MG BASE/ML	A075176 002	Feb 09, 2000
SANDOZ INC	EQ 50MG BASE/ML	A076463 001	Jun 24, 2005
	EQ 100MG BASE/ML	A076463 002	Jun 24, 2005

HALOPERIDOL LACTATE

CONCENTRATE;ORAL

## HALDOL

ORTHO MCNEIL	EQ 2MG BASE/ML **	N015922 001	
HALOPERIDOL			
ALPHARMA	EQ 2MG BASE/ML	A070318 001	Apr 11, 1986
MORTON GROVE	EQ 2MG BASE/ML	A070710 001	Mar 07, 1986
SCS	EQ 2MG BASE/ML	A070726 001	Jun 10, 1986
TEVA	EQ 2MG BASE/ML	A071015 001	Aug 25, 1987
HALOPERIDOL INTENSOL			
CYCLE PHARMS LTD	EQ 2MG BASE/ML	A072045 001	Apr 12, 1988

INJECTABLE; INJECTION

## HALOPERIDOL

ABRAXIS PHARM	EQ 5MG BASE/ML	A071187 001	Jan 20, 1987
BAXTER HLTHCARE CORP	EQ 5MG BASE/ML	A076791 001	Aug 25, 2004
	EQ 5MG BASE/ML	A076828 001	Aug 25, 2004
MARSAM PHARMS LLC	EQ 5MG BASE/ML	A072516 001	Feb 25, 1993
	EQ 5MG BASE/ML	A072517 001	Feb 25, 1993
SANDOZ INC	EQ 5MG BASE/ML	A076464 001	Sep 29, 2004
SMITH AND NEPHEW	EQ 5MG BASE/ML	A070802 001	Dec 14, 1987
SOLOPAK	EQ 5MG BASE/ML	A070800 001	Dec 14, 1987
	EQ 5MG BASE/ML	A070801 001	Dec 14, 1987
	EQ 5MG BASE/ML	A070864 001	Dec 14, 1987
WATSON LABS	EQ 5MG BASE/ML	A070713 001	May 17, 1988
	EQ 5MG BASE/ML	A070714 001	May 17, 1988
	EQ 5MG BASE/ML	A070744 001	May 17, 1988

SOLUTION;ORAL

## HALOPERIDOL LACTATE

ACTAVIS MID ATLANTIC	EQ 1MG BASE/ML	A074536 001	Nov 02, 1995
----------------------	----------------	-------------	--------------

HALOPROGIN

CREAM;TOPICAL

## HALOTEX

WESTWOOD SQUIBB	1%	N016942 001	
SOLUTION;TOPICAL			
HALOTEX			
WESTWOOD SQUIBB	1%	N016943 001	

**DISCONTINUED DRUG PRODUCT LIST**

6-177(of 375)

\*\* See List Footnote

HALOTHANE

LIQUID; INHALATION

FLUOTHANE

WYETH AYERST

99.99%

N011338 001

HALOTHANE

BH

99.99%

A084977 001

HALOCARBON

99.99%

A080810 001

HOSPIRA

99.99%

A083254 001

HEPARIN CALCIUM

INJECTABLE; INJECTION

CALCIPARINE

SANOFI AVENTIS US

25,000 UNITS/ML

N018237 001

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN LOCK FLUSH

HOSPIRA

100 UNITS/ML

N005264 010

INT'L MEDICATION

10 UNITS/ML

A086357 001

500 UNITS/ML

A086357 002

LUITPOLD

10 UNITS/ML

A089063 001 Oct 09, 1985

100 UNITS/ML

A089064 001 Oct 09, 1985

PARKE DAVIS

10 UNITS/ML

N017346 006

SMITH AND NEPHEW

10 UNITS/ML

A087904 001 Apr 20, 1983

10 UNITS/ML

A087958 001 Apr 20, 1983

10 UNITS/ML

A088458 001 Jul 26, 1984

10 UNITS/ML

A088580 001 Oct 25, 1984

100 UNITS/ML

A087906 001 Apr 20, 1983

100 UNITS/ML

A087959 001 Apr 20, 1983

100 UNITS/ML

A088460 001 Jul 26, 1984

100 UNITS/ML

A088581 001 Oct 25, 1984

SOLOPAK

10 UNITS/ML

A087903 001 Apr 20, 1983

10 UNITS/ML

A088457 001 Oct 25, 1984

100 UNITS/ML

A087905 001 Apr 20, 1983

100 UNITS/ML

A088459 001 Jul 26, 1984

HEPARIN SODIUM

ABRAXIS PHARM

1,000 UNITS/ML

N017033 001

1,000 UNITS/ML

N017979 001

5,000 UNITS/ML

N017979 003

10,000 UNITS/ML

N017979 002

AKORN

1,000 UNITS/ML

N017486 001

5,000 UNITS/ML

N017486 002

10,000 UNITS/ML

N017486 003

20,000 UNITS/ML

N017486 004

40,000 UNITS/ML

N017486 005

CHAMBERLIN PARENTERL

1,000 UNITS/ML

N017130 001

5,000 UNITS/ML

N017130 002

10,000 UNITS/ML

N017130 003

20,000 UNITS/ML

N017130 004

DELL LABS

1,000 UNITS/ML

N017540 001

5,000 UNITS/ML

N017540 002

10,000 UNITS/ML

N017540 003

20,000 UNITS/ML

N017540 004

40,000 UNITS/ML

N017540 005

FRESENIUS KABI USA

1,000 UNITS/ML

N017651 005

5,000 UNITS/ML

N017029 002

10,000 UNITS/ML

N017651 003

20,000 UNITS/ML

N017651 008

HOSPIRA

2,500 UNITS/ML

A088099 001 Apr 28, 1983

10,000 UNITS/ML

A040095 001 Jul 26, 1996

LILLY

1,000 UNITS/ML

N005521 001

10,000 UNITS/ML

N005521 002

20,000 UNITS/ML

N005521 004

LUITPOLD

1,000 UNITS/ML

A087452 001 Oct 31, 1983

ORGANON USA INC

1,000 UNITS/ML

N000552 008

5,000 UNITS/ML

N000552 009

10,000 UNITS/ML

N000552 010

PARKE DAVIS

1,000 UNITS/ML

N017346 001

5,000 UNITS/ML

N017346 002

7,500 UNITS/ML

N017346 003

**DISCONTINUED DRUG PRODUCT LIST**

6-178(of 375)

\*\* See List Footnote

HEPARIN SODIUMINJECTABLE; INJECTION  
HEPARIN SODIUM

	10,000 UNITS/ML	N017346 004
	20,000 UNITS/ML	N017346 005
PHARM SPEC	1,000 UNITS/ML	N017780 001
	5,000 UNITS/ML	N017780 002
	10,000 UNITS/ML	N017780 003
	20,000 UNITS/ML	N017780 004
	40,000 UNITS/ML	N017780 005
PHARMACIA AND UPJOHN	1,000 UNITS/ML	N004570 001
	5,000 UNITS/ML	N004570 002
	10,000 UNITS/ML	N004570 003
SMITH AND NEPHEW	1,000 UNITS/ML	A088239 001 Jul 26, 1984
SOLOPAK	1,000 UNITS/ML	A087043 001
	5,000 UNITS/ML	A087077 001
	5,000 UNITS/0.5ML	A087395 001
	10,000 UNITS/ML	A087107 001
	10,000 UNITS/0.5ML	A087363 001
WATSON LABS	1,000 UNITS/ML	N017064 002
	2,500 UNITS/ML	N017064 015
	3,000 UNITS/ML	N017064 016
	4,000 UNITS/ML	N017064 017
	5,000 UNITS/ML	N017064 003
	6,000 UNITS/ML	N017064 018
	7,500 UNITS/ML	N017064 019
	10,000 UNITS/ML	N017064 004
	20,000 UNITS/ML	N017064 005
	40,000 UNITS/ML	N017064 006
WATSON LABS INC	1,000 UNITS/ML	A040007 001 Jun 07, 1996
	1,000 UNITS/ML	A040008 001 Oct 10, 1995
WEST-WARD PHARMS INT	1,000 UNITS/ML	N017007 001
	2,500 UNITS/ML	N017007 007
	5,000 UNITS/ML	N017007 002
	5,000 UNITS/0.5ML	N017007 010
	5,000 UNITS/0.5ML	N017037 013 Apr 07, 1986
	7,500 UNITS/ML	N017007 003
	10,000 UNITS/ML	N017007 004
	15,000 UNITS/ML	N017007 005
	20,000 UNITS/ML	N017007 006
HEPARIN SODIUM 1,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER		
MCGAW	200 UNITS/100ML	N019130 001 Dec 31, 1984
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
B BRAUN	200 UNITS/100ML	N019042 001 Mar 29, 1985
HEPARIN SODIUM 10,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER		
BAXTER HLTHCARE	2,000 UNITS/100ML	N018814 002 Jul 09, 1985
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5%		
HOSPIRA	10,000 UNITS/100ML	N018911 006 Jan 30, 1985
HEPARIN SODIUM 10,000 UNITS IN SODIUM CHLORIDE 0.45%		
HOSPIRA	10,000 UNITS/100ML	N018911 001 Jan 30, 1985
	10,000 UNITS/100ML	N018916 005 Jan 31, 1984
HEPARIN SODIUM 10,000 UNITS IN SODIUM CHLORIDE 0.9%		
HOSPIRA	10,000 UNITS/100ML	N018911 003 Jan 30, 1985
	10,000 UNITS/100ML	N018916 002 Jan 31, 1984
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5%		
HOSPIRA	5,000 UNITS/100ML	N018911 007 Jan 30, 1985
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
B BRAUN	5,000 UNITS/100ML	N019802 001 Jul 20, 1992
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.9%		
HOSPIRA	5,000 UNITS/100ML	N018911 005 Jan 30, 1985
	5,000 UNITS/100ML	N018916 003 Jan 31, 1984
HEPARIN SODIUM 2,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER		
MCGAW	200 UNITS/100ML	N019130 003 Dec 31, 1984
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
B BRAUN	200 UNITS/100ML	N019042 002 Mar 29, 1985
HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER		
BAXTER HLTHCARE	4,000 UNITS/100ML	N018814 001 Oct 31, 1983
HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER		
BAXTER HLTHCARE	5,000 UNITS/100ML	N018814 003 Jul 09, 1985

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-179(of 375)

\*\* See List Footnote

HEPARIN SODIUM

## INJECTABLE; INJECTION

HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER	10,000 UNITS/100ML	N018814 004 Jul 02, 1987
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5%		
HOSPIRA	5,000 UNITS/100ML	N018911 009 Jan 30, 1985
	10,000 UNITS/100ML	N018911 008 Jan 30, 1985
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER		
B BRAUN	5,000 UNITS/100ML	N019134 001 Mar 29, 1985
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
B BRAUN	5,000 UNITS/100ML	N019802 005 Jul 20, 1992
	10,000 UNITS/100ML	N019802 002 Jul 20, 1992
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9%		
HOSPIRA	5,000 UNITS/100ML	N018911 004 Jan 30, 1985
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
B BRAUN	5,000 UNITS/100ML	N019135 001 Mar 29, 1985
	5,000 UNITS/100ML	N019802 003 Jul 20, 1992
HOSPIRA	5,000 UNITS/100ML	N018916 009 Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
BAXTER HLTHCARE	500 UNITS/100ML	N018609 003 Apr 28, 1982
HEPARIN SODIUM 5,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER		
MCGAW	1,000 UNITS/100ML	N019130 002 Dec 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.45%		
HOSPIRA	100 UNITS/ML	N018911 002 Jan 30, 1985
	100 UNITS/ML	N018916 004 Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9%		
HOSPIRA	1,000 UNITS/100ML	N018916 001 Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
B BRAUN	1,000 UNITS/100ML	N019042 004 Mar 29, 1985
HEPARIN SODIUM PRESERVATIVE FREE		
HOSPIRA	2,000 UNITS/ML	N005264 013 Apr 07, 1986
	2,500 UNITS/ML	N005264 014 Apr 07, 1986
PHARMA SERVE NY	1,000 UNITS/ML	A086129 001
WATSON LABS INC	1,000 UNITS/ML	A089464 001 Jun 03, 1986
LIPO-HEPIN		
3M	1,000 UNITS/0.5ML	N017027 001
	1,000 UNITS/ML	N017027 006
	5,000 UNITS/0.5ML	N017027 002
	5,000 UNITS/ML	N017027 008
	7,500 UNITS/0.5ML	N017027 010
	10,000 UNITS/0.5ML	N017027 003
	10,000 UNITS/ML	N017027 009
	15,000 UNITS/0.5ML	N017027 011
	20,000 UNITS/0.5ML	N017027 004
	20,000 UNITS/ML	N017027 007
	40,000 UNITS/ML	N017027 005
LIQUAEMIN LOCK FLUSH		
ORGANON USA INC	100 UNITS/ML	N000552 007
LIQUAEMIN SODIUM		
ORGANON USA INC	1,000 UNITS/ML	N000552 004
	5,000 UNITS/ML	N000552 003
	10,000 UNITS/ML	N000552 005
	20,000 UNITS/ML	N000552 001
	40,000 UNITS/ML	N000552 002
LIQUAEMIN SODIUM PRESERVATIVE FREE		
ORGANON USA INC	1,000 UNITS/ML	N000552 011 Apr 11, 1986
	5,000 UNITS/ML	N000552 012 Apr 11, 1986
	10,000 UNITS/ML	N000552 013 Apr 11, 1986
PANHEPRIN		
HOSPIRA	1,000 UNITS/ML	N005264 004
	5,000 UNITS/ML	N005264 006
	10,000 UNITS/ML	N005264 007
	20,000 UNITS/ML	N005264 008
	40,000 UNITS/ML	N005264 009
SODIUM HEPARIN		
ABRAXIS PHARM	5,000 UNITS/ML	N017033 002
	10,000 UNITS/ML	N017033 003
	20,000 UNITS/ML	N017033 004
BAXTER HLTHCARE	1,000 UNITS/ML	N017036 001 Mar 04, 1988

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-180(of 375)

\*\* See List Footnote

HETACILLINFOR SUSPENSION;ORAL  
VERSAPEN

BRISTOL	EQ 112.5MG AMPICIL/ML	A061398 001
	EQ 112.5MG AMPICIL/5ML	N050060 001
	EQ 112.5MG AMPICIL/ML	N050060 003
	EQ 225MG AMPICIL/5ML	A061398 002

HETACILLIN POTASSIUMCAPSULE;ORAL  
VERSAPEN-K

BRISTOL	EQ 225MG AMPICIL	A061396 001
	EQ 450MG AMPICIL	A061396 002

HEXACHLOROPHENEAEROSOL;TOPICAL  
SEPTISOL

VESTAL LABS	0.23%	N017424 001
-------------	-------	-------------

TURGEX	3%	N018375 001
--------	----	-------------

XTTRIUM	3%	N018375 001
---------	----	-------------

EMULSION;TOPICAL

HEXA-GERM	3%	N017411 001
-----------	----	-------------

HUNTINGTON LABS	3%	N006882 001
-----------------	----	-------------

PHISOHEX	3%	N008402 001
----------	----	-------------

SANOFI AVENTIS US	3%	N017405 001
-------------------	----	-------------

	3%	
--	----	--

SOY-DOME		
----------	--	--

BAYER PHARMS	3%	N017405 001
--------------	----	-------------

TURGEX	3%	
--------	----	--

XTTRIUM	3%	N019055 001 Nov 30, 1984
---------	----	--------------------------

SOAP;TOPICAL

GAMOPHEN		
----------	--	--

ARBROOK	2%	N006270 003
---------	----	-------------

SOLUTION;TOPICAL

DIAL		
------	--	--

DIAL	0.25%	N017421 002
------	-------	-------------

GERMA-MEDICA		
--------------	--	--

HUNTINGTON LABS	1%	N017412 001
-----------------	----	-------------

GERMA-MEDICA "MG"		
-------------------	--	--

HUNTINGTON LABS	0.25%	N017412 002
-----------------	-------	-------------

SEPTI-SOFT		
------------	--	--

CALGON	0.25%	N017460 001
--------	-------	-------------

SEPTISOL		
----------	--	--

VESTAL LABS	0.25%	N017423 001
-------------	-------	-------------

SPONGE;TOPICAL

E-Z SCRUB		
-----------	--	--

BECTON DICKINSON	450MG	N017452 001
------------------	-------	-------------

HEXASCRUB		
-----------	--	--

PROF DSPLS	3%	N018363 001
------------	----	-------------

PHISO-SCRUB		
-------------	--	--

SANOFI AVENTIS US	3%	N017446 001
-------------------	----	-------------

SCRUBTEAM SURGICAL SPONGEBRUSH		
--------------------------------	--	--

3M	330MG	N017413 001
----	-------	-------------

HEXAFLUORENIUM BROMIDE

INJECTABLE;INJECTION

MYLAXEN		
---------	--	--

MEDPOINTE PHARM HLC	20MG/ML	N009789 003
---------------------	---------	-------------

HEXOCYCLIUM METHYLSULFATETABLET;ORAL  
TRAL

ABBVIE	25MG	N010599 001
--------	------	-------------

HEXYLCAINE HYDROCHLORIDESOLUTION;TOPICAL  
CYCLAINE

MERCK	5%	N008472 001
-------	----	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-181(of 375)

\*\* See List Footnote

HISTAMINE PHOSPHATE

INJECTABLE; INJECTION

HISTAMINE PHOSPHATE

LILLY

EQ 0.1MG BASE/ML

N000734 003

EQ 0.2MG BASE/ML

N000734 002

EQ 1MG BASE/ML

N000734 001

HISTRELIN ACETATE

INJECTABLE; INJECTION

SUPPRELIN

SHIRE

EQ 0.2MG BASE/ML

N019836 001 Dec 24, 1991

EQ 0.5MG BASE/ML

N019836 002 Dec 24, 1991

EQ 1MG BASE/ML

N019836 003 Dec 24, 1991

HOMATROPINE METHYLBROMIDE

TABLET; ORAL

HOMAPIN-10

MISSION PHARMA

10MG

A086308 001

HOMAPIN-5

MISSION PHARMA

5MG

A086309 001

TABLET, CHEWABLE; ORAL

EQUIPIN

MISSION PHARMA

3MG

A086310 001

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN

+ ENDO PHARMS

1.5MG/5ML; 5MG/5ML \*\*

N005213 002 Jul 26, 1988

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

IVAX SUB TEVA PHARMS 1.5MG/5ML; 5MG/5ML

A040285 001 Jul 19, 1999

HYDROPANE

HALSEY

1.5MG/5ML; 5MG/5ML

A088066 001 Jun 28, 1985

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

ACTAVIS ELIZABETH 1.5MG; 5MG

A040295 001 Dec 01, 2000

HYCODAN

+ ENDO PHARMS

1.5MG; 5MG \*\*

N005213 001 Jul 26, 1988

HYALURONIDASE

INJECTABLE; INJECTION

HYDASE

AKORN INC

150 UNITS/ML

N021716 001 Oct 25, 2005

VITRASE

BAUSCH AND LOMB

6,200 UNITS/VIAL

N021640 001 May 05, 2004

WYDASE

BAXTER HLTHCARE

150 UNITS/ML \*\*

N006343 002

150 UNITS/VIAL \*\*

N006343 006

1,500 UNITS/VIAL \*\*

N006343 005

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

APRESOLINE

+ NOVARTIS

20MG/ML \*\*

N008303 003

HYDRALAZINE HYDROCHLORIDE

ABRAXIS PHARM

20MG/ML

A089532 001 Aug 11, 1987

SMITH AND NEPHEW

20MG/ML

A088518 001 Apr 20, 1984

SOLOPAK

20MG/ML

A088517 001 Aug 22, 1985

TEVA PARENTERAL

20MG/ML

A040373 001 Feb 23, 2000

TABLET; ORAL

APRESOLINE

+ NOVARTIS

10MG \*\*

N008303 004

+

25MG \*\*

N008303 001

+

50MG \*\*

N008303 002

+

100MG \*\*

N008303 005

DRALZINE

TEVA

25MG

A084301 001

HYDRALAZINE HYDROCHLORIDE

ACTAVIS ELIZABETH

25MG

A088560 001 Oct 04, 1984

50MG

A088649 001 Oct 18, 1984

ACTAVIS GRP PTC

10MG

A091679 001 Mar 04, 2013

25MG

A091679 002 Mar 04, 2013

50MG

A091679 003 Mar 04, 2013

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-182(of 375)

\*\* See List Footnote

HYDRALAZINE HYDROCHLORIDE

TABLET;ORAL

HYDRALAZINE HYDROCHLORIDE

	100MG	A091679 004	Mar 04, 2013
ANDA REPOSITORY	10MG	A089359 001	Jul 25, 1986
	25MG	A089258 001	May 05, 1986
	50MG	A089259 001	May 05, 1986
	100MG	A088729 001	Apr 11, 1985
ASCOT	25MG	A088310 001	Dec 19, 1984
	50MG	A088311 001	Dec 19, 1984
CHARTWELL RX	10MG	A088846 001	Feb 26, 1985
	25MG	A088847 001	Feb 26, 1985
	50MG	A088848 001	Feb 26, 1985
	100MG	A088849 001	Feb 26, 1985
HALSEY	10MG	A089218 001	Jan 22, 1986
	25MG	A089130 001	Jan 15, 1986
	50MG	A089222 001	Jan 22, 1986
	100MG	A089178 001	Jan 15, 1986
HERITAGE PHARMS INC	10MG	A040858 001	Feb 26, 2010
	25MG	A040858 002	Feb 26, 2010
	50MG	A040858 003	Feb 26, 2010
	100MG	A040858 004	Feb 26, 2010
IMPAX LABS	25MG	A084922 001	
	50MG	A084923 001	
IVAX SUB TEVA PHARMS	10MG	A084443 001	
	25MG	A084437 001	
	50MG	A084469 002	
	100MG	A084581 001	
MUTUAL PHARM	10MG	A088728 001	Apr 11, 1985
	25MG	A084106 002	
	50MG	A084107 002	
MYLAN	10MG	A090413 001	Dec 08, 2010
	25MG	A090413 002	Dec 08, 2010
	50MG	A090413 003	Dec 08, 2010
	100MG	A090413 004	Dec 08, 2010
PUREPAC PHARM	25MG	A088177 001	Jul 29, 1983
	50MG	A088178 001	Aug 15, 1983
QUANTUM PHARMICS	10MG	A088671 001	May 01, 1984
	25MG	A088657 001	Jun 15, 1984
	50MG	A088652 001	May 08, 1984
	100MG	A088686 001	May 01, 1984
SUPERPHARM	10MG	A088787 001	Aug 28, 1984
	25MG	A088788 001	Aug 28, 1984
	50MG	A088789 001	Aug 28, 1984
UPSHER-SMITH LABS	10MG	A083241 001	
	25MG	A083560 001	
	50MG	A083561 001	
	50MG	A085088 001	
USL PHARMA	25MG	A087780 001	Mar 29, 1982
	50MG	A087751 001	Mar 29, 1982
VANGARD	25MG	A087712 001	
	50MG	A087908 001	May 07, 1982
VITARINE	25MG	A086088 001	
WATSON LABS	25MG	A084504 001	
	25MG	A085532 002	May 24, 1982
	50MG	A084503 001	
	50MG	A085533 002	May 25, 1982
WEST WARD	25MG	A088240 001	May 27, 1983
	50MG	A088241 001	May 27, 1983

HYDRALAZINE HYDROCHLORIDE; HYDROCHLORTHIAZIDE

CAPSULE;ORAL

APRESAZIDE

NOVARTIS	25MG;25MG	A084735 001
	50MG;50MG	A084810 001
	100MG;50MG	A084811 001

HYDRA-ZIDE	100MG;50MG	A088961 001	Oct 21, 1985
------------	------------	-------------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-183(of 375)

\*\* See List Footnote

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

SOLVAY	25MG;25MG	A087608 001 Feb 08, 1982
	50MG;50MG	A087213 001 Feb 08, 1982
	100MG;50MG	A087609 001 Feb 08, 1982
SUPERPHARM	25MG;25MG	A089200 001 Feb 09, 1987
	50MG;50MG	A089201 001 Feb 09, 1987
WATSON LABS	25MG;25MG	A085457 001 Mar 04, 1982
	50MG;50MG	A085446 001 Mar 04, 1982
	100MG;50MG	A085440 001 Mar 04, 1982

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 100/50

IVAX PHARMS	100MG;50MG	A088358 001 Apr 10, 1984
-------------	------------	--------------------------

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 25/25

IVAX PHARMS	25MG;25MG	A088356 001 Apr 10, 1984
-------------	-----------	--------------------------

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 50/50

IVAX PHARMS	50MG;50MG	A088357 001 Apr 10, 1984
-------------	-----------	--------------------------

TABLET; ORAL

APRESOLINE-ESIDRIX

NOVARTIS	25MG;15MG	N012026 002
----------	-----------	-------------

HYDRALAZINE AND HYDROCHLOROTHIAZIDE

WATSON LABS	25MG;15MG	A085827 001
-------------	-----------	-------------

HYDROCHLOROTHIAZIDE W/ HYDRALAZINE

WATSON LABS	25MG;15MG	A085373 001
-------------	-----------	-------------

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

CHARTWELL RX	25MG;15MG;0.1MG	A084897 001
--------------	-----------------	-------------

HYDRALAZINE HYDROCHLORIDE, HYDROCHLOROTHIAZIDE AND RESERPINE

IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A084291 001
----------------------	-----------------	-------------

HYDRALAZINE HYDROCHLORIDE-HYDROCHLOROTHIAZIDE-RESERPINE

MYLAN	25MG;15MG;0.1MG	A087085 001
-------	-----------------	-------------

HYDRALAZINE, HYDROCHLOROTHIAZIDE W/ RESERPINE

WATSON LABS	25MG;15MG;0.1MG	A085771 001
-------------	-----------------	-------------

HYDRAP-ES

SANDOZ	25MG;15MG;0.1MG	A084876 001
--------	-----------------	-------------

HYDROCHLOROTHIAZIDE W/ RESERPINE AND HYDRALAZINE

WATSON LABS	25MG;15MG;0.1MG	A083770 001
-------------	-----------------	-------------

HYDROSERPINE PLUS (R-H-H)

IVAX SUB TEVA PHARMS	25MG;15MG;0.1MG	A083877 001
----------------------	-----------------	-------------

RESERPINE, HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

SOLVAY	25MG;15MG;0.1MG	A088376 001 Oct 28, 1983
--------	-----------------	--------------------------

SUN PHARM INDUSTRIES	25MG;15MG;0.1MG	A088570 001 Apr 10, 1984
----------------------	-----------------	--------------------------

WATSON LABS	25MG;15MG;0.1MG	A085549 001
-------------	-----------------	-------------

	25MG;15MG;0.1MG	A087556 001
--	-----------------	-------------

RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HYDROCHLORIDE

LEDERLE	25MG;15MG;0.1MG	A087709 001 May 13, 1982
---------	-----------------	--------------------------

SER-A-GEN

SOLVAY	25MG;15MG;0.1MG	A087210 001
--------	-----------------	-------------

SER-AP-ES

NOVARTIS	25MG;15MG;0.1MG	N012193 005
----------	-----------------	-------------

UNIPRES

SOLVAY	25MG;15MG;0.1MG	A085893 001
--------	-----------------	-------------

	25MG;15MG;0.1MG	A086298 001
--	-----------------	-------------

HYDRALAZINE HYDROCHLORIDE; RESERPINE

TABLET; ORAL

DRALSERP

SANDOZ	25MG;0.1MG	A084617 001
--------	------------	-------------

SERPASIL-APRESOLINE

NOVARTIS	25MG;0.1MG	N009296 004
----------	------------	-------------

	50MG;0.2MG	N009296 002
--	------------	-------------

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

HIKMA INTL PHARMS	12.5MG	A077885 001 Nov 26, 2007
-------------------	--------	--------------------------

LANNETT HOLDINGS INC	12.5MG	A091662 001 Jan 27, 2012
----------------------	--------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-184(of 375)

\*\* See List Footnote

HYDROCHLOROTHIAZIDE

SOLUTION;ORAL

## HYDROCHLOROTHIAZIDE

MORTON GROVE	50MG/5ML	A089661 001 Jun 20, 1988
ROXANE	50MG/5ML	A088587 001 Jul 02, 1984

## HYDROCHLOROTHIAZIDE INTENSOL

ROXANE	100MG/ML	A088588 001 Jul 02, 1984
--------	----------	--------------------------

TABLET;ORAL

## ESIDRIX

NOVARTIS	25MG	N011793 005
	50MG	N011793 008
	100MG	N011793 009

## HYDRO-D

HALSEY	25MG	A086504 001
	50MG	A083891 002

## HYDROCHLOROTHIAZIDE

ABC HOLDING	50MG	A085672 001
ACTAVIS ELIZABETH	25MG	A085054 002
	50MG	A085208 001

ALRA	25MG	A086369 001
	50MG	A083554 001

ASCOT	25MG	A087539 001 Feb 03, 1982
	50MG	A087540 001 Feb 03, 1982

AUROLIFE PHARMA LLC	25MG	A083899 001
	50MG	A085219 001

BARR	50MG	A084771 001
CHARTWELL RX	25MG	A085683 001
	50MG	A083965 001

DAVA PHARMS INC	100MG	A087060 001
ELKINS SINK	50MG	A085152 002

FOSUN PHARMA	25MG	A087565 001 Mar 09, 1982
	50MG	A084912 001

HEATHER	50MG	A084135 001
HIKMA INTL PHARMS	25MG	A084878 002 Jul 12, 2006

IMPAX LABS	25MG	A084029 001
	50MG	A083607 002
	100MG	A085098 001

INWOOD LABS	25MG	A084776 001
	50MG	A085067 001

IVAX SUB TEVA PHARMS	50MG	A084658 001
	100MG	A085022 001

JUBILANT CADISTA	25MG	A040809 001 Sep 04, 2007
	50MG	A040809 002 Sep 04, 2007

LANNETT	25MG	A084325 001
	50MG	A084324 001

MAST MM	25MG	A086192 001
	50MG	A086192 002

MYLAN	25MG	A084880 001
	50MG	A085112 001

MYLAN PHARMS INC	12.5MG	A040770 001 Jan 23, 2007
PVT FORM	50MG	A086597 001

ROXANE	25MG	A085004 001
	50MG	A084536 002
	50MG	A085005 001

SOLVAY	25MG	A085323 001
SUN PHARM INDUSTRIES	25MG	A083972 001
	50MG	A083972 002
	100MG	A083972 003

SUPERPHARM	25MG	A088827 001 Dec 28, 1984
	50MG	A088828 001 Dec 28, 1984
	100MG	A088829 001 Dec 28, 1984

TEVA	25MG	A088924 001 Feb 07, 1985
	50MG	A088923 001 Feb 07, 1985

USL PHARMA	25MG	A087827 001 Apr 19, 1982
	50MG	A087752 001 Apr 19, 1982

VANGARD	25MG	A087638 001
	50MG	A087610 001

WARNER CHILCOTT	25MG	A087586 001 May 03, 1982
-----------------	------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-185(of 375)

\*\* See List Footnote

HYDROCHLOROTHIAZIDE

TABLET;ORAL

HYDROCHLOROTHIAZIDE

	50MG	A087587 001 May 03, 1982
WATSON LABS	25MG	A081189 001 Jan 24, 1992
	25MG	A083458 001
	25MG	A085232 002
	50MG	A083456 001
	50MG	A085233 001
	50MG	A086087 001
	50MG	A086594 001
	100MG	A081190 001 Jan 24, 1992
	100MG	A085099 001
	100MG	A087002 001
WATSON LABS TEVA	50MG	A083232 001
WEST WARD	25MG	A084899 001
WHITEWORTH TOWN PLSN	25MG	A083809 002
	50MG	A083809 001
	100MG	A085347 001
HYDRODIURIL		
+ MERCK	25MG **	N011835 003
+	50MG **	N011835 006
+	100MG **	N011835 007
ORETIC		
ABBVIE	25MG	N011971 001
	50MG	N011971 002
ZIDE		
SOLVAY	50MG	A083925 001

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET;ORAL

AVALIDE

+ SANOFI AVENTIS US	12.5MG;75MG **	N020758 001 Sep 30, 1997
+	25MG;300MG **	N020758 004 Mar 15, 2005
IRBESARTAN AND HYDROCHLOROTHIAZIDE		
TEVA	25MG;300MG	A077369 003 Mar 30, 2012
WATSON LABS INC	12.5MG;150MG	A091539 001 Oct 22, 2012
	12.5MG;300MG	A091539 002 Oct 22, 2012

HYDROCHLOROTHIAZIDE; LABETALOL HYDROCHLORIDE

TABLET;ORAL

NORMOZIDE

SCHERING	25MG;100MG	N019046 001 Apr 06, 1987
	25MG;200MG	N019046 002 Apr 06, 1987
	25MG;300MG	N019046 003 Apr 06, 1987
	25MG;400MG	N019046 004 Apr 06, 1987
TRANDATE HCT		
GLAXOSMITHKLINE	25MG;100MG	N019174 001 Apr 10, 1987
	25MG;200MG	N019174 002 Apr 10, 1987
	25MG;300MG	N019174 003 Apr 10, 1987
	25MG;400MG	N019174 004 Apr 10, 1987

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET;ORAL

LISINOPRIL AND HYDROCHLOROTHIAZIDE

SANDOZ	12.5MG;10MG	A075926 001 Jul 01, 2002
	12.5MG;20MG	A075926 002 Jul 01, 2002
	25MG;20MG	A075926 003 Jul 01, 2002
TEVA		
	12.5MG;10MG	A075869 001 Jul 01, 2002
	12.5MG;20MG	A075869 002 Jul 01, 2002
	25MG;20MG	A075869 003 Jul 01, 2002
PRINZIDE		
+ MERCK	12.5MG;10MG **	N019778 003 Nov 18, 1993
+	12.5MG;20MG **	N019778 001 Feb 16, 1989
+	25MG;20MG **	N019778 002 Feb 16, 1989

**DISCONTINUED DRUG PRODUCT LIST**

6-186(of 375)

\*\* See List Footnote

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

## TABLET;ORAL

LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE  
 WATSON LABS            12.5MG;50MG  
                         12.5MG;100MG  
                         25MG;100MG

A200180 001 Jan 12, 2011  
 A200180 002 Jan 12, 2011  
 A200180 003 Jan 12, 2011

HYDROCHLOROTHIAZIDE; METHYLDOPA

## TABLET;ORAL

ALDORIL 15 MERCK	15MG;250MG	N013402 001
ALDORIL 25 MERCK	25MG;250MG	N013402 002
ALDORIL D30 MERCK	30MG;500MG	N013402 003
ALDORIL D50 MERCK	50MG;500MG	N013402 004
METHYLDOPA AND HYDROCHLOROTHIAZIDE		
DAVA PHARMS INC	15MG;250MG	A072507 001 Jun 02, 1989
	25MG;250MG	A072508 001 Jun 02, 1989
	30MG;500MG	A072509 001 Jun 02, 1989
	50MG;500MG	A072510 001 Jun 02, 1989
FOSUN PHARMA	15MG;250MG	A070182 001 Jan 15, 1986
	25MG;250MG	A070183 001 Jan 15, 1986
	30MG;500MG	A070543 001 Jan 15, 1986
IVAX SUB TEVA PHARMS	15MG;250MG	A071458 001 Mar 08, 1988
	25MG;250MG	A071459 001 Mar 08, 1988
	30MG;500MG	A071460 001 Mar 08, 1988
	50MG;500MG	A071461 001 Mar 08, 1988
PAR PHARM	15MG;250MG	A070616 001 Feb 02, 1987
	25MG;250MG	A070612 001 Feb 02, 1987
	30MG;500MG	A070613 001 Feb 02, 1987
	50MG;500MG	A070614 001 Feb 02, 1987
PARKE DAVIS	15MG;250MG	A071897 001 Nov 23, 1987
	25MG;250MG	A071898 001 Nov 23, 1987
	30MG;500MG	A071899 001 Nov 23, 1987
	50MG;500MG	A071900 001 Nov 23, 1987
PUREPAC PHARM	15MG;250MG	A070853 001 Oct 08, 1986
	25MG;250MG	A070688 001 Apr 24, 1986
	30MG;500MG	A070854 001 Oct 08, 1986
	50MG;500MG	A070689 001 Apr 24, 1986
SANDOZ	15MG;250MG	A070829 001 Mar 09, 1987
	25MG;250MG	A070830 001 Mar 09, 1987
	50MG;500MG	A070544 001 Jan 15, 1986
TEVA	15MG;250MG	A071819 001 Apr 08, 1988
	25MG;250MG	A071820 001 Apr 08, 1988
	30MG;500MG	A071821 001 Apr 08, 1988
	50MG;500MG	A071822 001 Apr 08, 1988
WATSON LABS	15MG;250MG	A070365 001 Mar 19, 1986
	15MG;250MG	A070958 001 Feb 06, 1989
	15MG;250MG	A071920 001 Aug 29, 1988
	25MG;250MG	A070366 001 Apr 16, 1986
	25MG;250MG	A070959 001 Jan 19, 1989
	30MG;500MG	A071921 001 Aug 29, 1988
	30MG;500MG	A070367 001 Mar 19, 1986
	30MG;500MG	A071069 001 Jan 19, 1989
	50MG;500MG	A071922 001 Aug 29, 1988
	50MG;500MG	A070368 001 Apr 16, 1986
	50MG;500MG	A070960 001 Feb 06, 1989
	50MG;500MG	A071923 001 Aug 29, 1988

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

## TABLET;ORAL

LOPRESSOR HCT

+ US PHARMS HOLDINGS I 50MG;100MG \*\*

N018303 003 Dec 31, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-187(of 375)

\*\* See List Footnote

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET;ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

CHARTWELL RX	12.5MG; 7.5MG 12.5MG; 15MG 25MG; 15MG	A090096 001 Sep 25, 2008 A090096 002 Sep 25, 2008 A090096 003 Sep 25, 2008
UNIRETIC		
UCB INC	12.5MG; 7.5MG ** 12.5MG; 15MG ** 25MG; 15MG **	N020729 001 Jun 27, 1997 N020729 003 Feb 14, 2002 N020729 002 Jun 27, 1997

HYDROCHLOROTHIAZIDE; PINDOLOL

TABLET;ORAL

VISKAZIDE

NOVARTIS	25MG; 5MG 25MG; 10MG	N018872 001 Jul 22, 1987 N018872 002 Jul 22, 1987
----------	-------------------------	--

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

INDERIDE LA 120/50		
WYETH AYERST	50MG; 120MG	N019059 002 Jul 03, 1985
INDERIDE LA 160/50		
WYETH AYERST	50MG; 160MG	N019059 003 Jul 03, 1985
INDERIDE LA 80/50		
WYETH AYERST	50MG; 80MG	N019059 001 Jul 03, 1985

TABLET;ORAL

INDERIDE-40/25		
+ WYETH PHARMS INC	25MG; 40MG **	N018031 001
INDERIDE-80/25		
+ WYETH PHARMS INC	25MG; 80MG **	N018031 002
PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE		
DURAMED PHARMS BARR	25MG; 40MG 25MG; 80MG	A071126 001 Mar 02, 1987 A071127 001 Mar 02, 1987
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE		
ACTAVIS ELIZABETH	25MG; 40MG 25MG; 80MG	A070851 001 May 15, 1986 A070852 001 May 15, 1986
ANI PHARMS INC	25MG; 40MG 25MG; 80MG	A070704 001 Oct 01, 1986 A070705 001 Oct 01, 1986
FOSUN PHARMA	25MG; 40MG 25MG; 80MG	A071060 001 Aug 26, 1987 A071061 001 Aug 26, 1987
IVAX SUB TEVA PHARMS	25MG; 40MG 25MG; 80MG	A071552 001 Dec 01, 1988 A071553 001 Dec 01, 1988
WARNER CHILCOTT	25MG; 40MG 25MG; 80MG	A071771 001 Jan 26, 1988 A071772 001 Jan 26, 1988
WATSON LABS	25MG; 40MG 25MG; 80MG 25MG; 80MG 25MG; 80MG	A070301 001 Apr 18, 1986 A071498 001 Dec 18, 1991 A070305 001 Apr 18, 1986 A071501 001 Dec 18, 1991

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE		
SUN PHARM INDS LTD	12.5MG; EQ 10MG BASE 12.5MG; EQ 20MG BASE	A078211 001 Mar 04, 2009 A078211 002 Mar 04, 2009
	25MG; EQ 20MG BASE	A078211 003 Mar 04, 2009

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET;ORAL

H.R.-50		
WHITEWORTH TOWN PLSN	50MG; 0.125MG	A085338 001
HYDRO-RESERP		
ABC HOLDING	50MG; 0.125MG	A084714 002 Jun 29, 1982
HYDRO-SERP "25"		
SANDOZ	25MG; 0.125MG	A084827 001
HYDRO-SERP "50"		
SANDOZ	50MG; 0.125MG	A085213 001
HYDROCHLOROTHIAZIDE W/ RESERPINE		
IVAX SUB TEVA PHARMS	25MG; 0.1MG 25MG; 0.125MG 50MG; 0.1MG 50MG; 0.125MG	A083572 001 A083571 001 A083568 001 A083573 001

**DISCONTINUED DRUG PRODUCT LIST**

6-188(of 375)

\*\* See List Footnote

HYDROCHLOROTHIAZIDE; RESERPINE

## TABLET;ORAL

## HYDROCHLOROTHIAZIDE W/ RESERPINE

PHARMERAL	25MG; 0.125MG	A085421 001
	50MG; 0.125MG	A085420 001
ROXANE	50MG; 0.125MG	A084603 001
WATSON LABS	25MG; 0.125MG	A084466 001
	25MG; 0.125MG	A085317 001
	25MG; 0.125MG	A086330 002
	50MG; 0.125MG	A083666 001
	50MG; 0.125MG	A084467 001
	50MG; 0.125MG	A086331 001
HYDROPPRES 25		
MERCK	25MG; 0.125MG	N011958 002
HYDROPPRES 50		
MERCK	50MG; 0.125MG	N011958 003
RESERPINE AND HYDROCHLOROTHIAZIDE		
BARR	25MG; 0.125MG	A084580 001
	50MG; 0.125MG	A084579 001
SANDOZ	50MG; 0.125MG	A088200 001 Jan 31, 1984
RESERPINE AND HYDROCHLOROTHIAZIDE-50		
WEST WARD	50MG; 0.125MG	A088189 001 May 10, 1984
SERPASIL-ESISIDRIX #1		
NOVARTIS	25MG; 0.1MG	N011878 003
SERPASIL-ESISIDRIX #2		
NOVARTIS	50MG; 0.1MG	N011878 005

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

## TABLET;ORAL

## SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

ASCOT	25MG; 25MG	A088025 001 Nov 23, 1984
FOSUN PHARMA	25MG; 25MG	A086881 001
MUTUAL PHARM	25MG; 25MG	A087267 001
PUREPAC PHARM	25MG; 25MG	A087999 001 Nov 06, 1985
SUPERPHARM	25MG; 25MG	A089137 001 Aug 26, 1985
WATSON LABS	25MG; 25MG	A087398 001
SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE		
IVAX PHARMS	25MG; 25MG	A087004 002 May 24, 1982
LEDERLE	25MG; 25MG	A087511 001
PARKE DAVIS	25MG; 25MG	A087948 001 Feb 22, 1983
PUREPAC PHARM	25MG; 25MG	A088054 001 Aug 18, 1983
UPSHER SMITH	25MG; 25MG	A087553 001
USL PHARMA	25MG; 25MG	A087651 001
VANGARD	25MG; 25MG	A087655 001
WATSON LABS	25MG; 25MG	A085974 001
	25MG; 25MG	A086026 001

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

## TABLET;ORAL

## TIMOLIDE 10-25

MERCK

25MG; 10MG

N018061 001

HYDROCHLOROTHIAZIDE; TRIAMTERENE

## CAPSULE;ORAL

## DYAZIDE

GLAXOSMITHKLINE LLC 25MG; 50MG

N016042 002

## TRIAMTERENE AND HYDROCHLOROTHIAZIDE

ANI PHARMS INC 25MG; 37.5MG

A074970 001 Jan 06, 1998

NOVARTIS 25MG; 37.5MG

A074857 001 Sep 09, 1997

VITARINE 25MG; 50MG

A071737 001 Feb 12, 1988

## TABLET;ORAL

## TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AM THERAP 50MG; 75MG

A072022 001 Apr 17, 1988

QUANTUM PHARMICS 50MG; 75MG

A071980 001 Apr 17, 1988

WATSON LABS 50MG; 75MG

A071969 001 Apr 17, 1988

**DISCONTINUED DRUG PRODUCT LIST**

6-189(of 375)

\*\* See List Footnote

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET;ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN  
 ACTAVIS LABS FL INC 5MG;200MG  
 VICOPROFEN  
 + ABBVIE 7.5MG;200MG

A077454 001 Jun 23, 2010  
 N020716 001 Sep 23, 1997

HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDESYRUP;ORAL  
 CODAMINE

ALPHARMA US PHARMS 5MG/5ML;25MG/5ML

A075103 001 Sep 29, 2000

HYDROCODONE BITARTRATE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION;ORAL

HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE  
 TRIS PHARMA INC 5MG/5ML;60MG/5ML

A203839 001 Oct 28, 2014

HYDROCORTAMATE HYDROCHLORIDEOINTMENT;TOPICAL  
 MAGNACORT

PFIZER 0.5%

N010554 001

HYDROCORTISONEAEROSOL;TOPICAL  
 AEROSEB-HC

ALLERGAN HERBERT 0.5%

A085805 001

CREAM;TOPICAL

CORT-DOME

BAYER PHARMS 0.5%  
 1%N009585 003  
 N009585 001

DERMACORT

MONARCH PHARMS 1%

A083011 002

ELDECORT

VALEANT PHARM INTL 1%  
 2.5%A080459 001  
 A084055 001

FLEXICORT

WESTWOOD SQUIBB 0.5%  
 1%  
 2.5%A087136 003 Apr 08, 1982  
 A087136 002 Apr 08, 1982  
 A087136 001 Apr 08, 1982

H-CORT

PHARM ASSOC 0.5%

A086823 001

HC #1

BAYER PHARMS 0.5%

A080438 001

HC #4

BAYER PHARMS 1%

A080438 002

HC (HYDROCORTISONE)

C AND M PHARMA 0.5%  
 1%A080482 003  
 A080482 004

HI-COR

C AND M PHARMA 2.5%

A080483 001

HYDROCORTISONE

ALPHARMA US PHARMS 2.5%

A089754 001 Feb 01, 1989

ALTANA 0.5%

A080848 002

1%

A080848 003

AMBIX

1%

A086080 001

2.5%

A086271 001

EVERYLIFE

0.5%

A080452 001

1%

A080452 002

G AND W LABS

1%

A084059 001

INGRAM PHARM

0.5%

A080456 002

1%

A080456 003

IVAX PHARMS

1%

A085733 001

NASKA

1%

A089706 001 Mar 10, 1988

PERRIGO NEW YORK

0.5%

A084970 002

1%

A085026 001

PHARMADERM

1%

A088845 001 Feb 27, 1986

2.5%

A089413 001 Dec 16, 1986

PHARMAFAIR

1%

A087838 001 Jul 28, 1982

STIEFEL

1%

A086170 001

SYOSSET

0.5%

A085527 001

TARO

0.5%

A086154 001

**DISCONTINUED DRUG PRODUCT LIST**

6-190(of 375)

\*\* See List Footnote

HYDROCORTISONECREAM;TOPICAL  
HYDROCORTISONE

TEVA	1%	A086155 001
	0.5%	A080400 002
	1%	A080400 003
	1%	A085191 001
	2.5%	A080400 004
TOPIDERM	1%	A089273 001 Feb 17, 1989
USL PHARMA	1%	A088027 001 Sep 27, 1983
	2.5%	A088029 001 Sep 27, 1983
WHITEWORTH TOWN PLSN	1%	A080496 002
HYTONE		
VALEANT INTL	1%	A080472 003
	2.5%	A080472 004
NOGENIC HC		
IVAX PHARMS	1%	A087427 001 Apr 04, 1988
NUTRACORT		
DOW PHARM	0.5%	A080442 002
	1%	A080442 003
PENEDECORT		
ALLERGAN HERBERT	1%	A088216 001 Jun 06, 1984
PROCTOCORT		
MONARCH PHARMS	1%	A083011 001
SYNACORT		
MEDICIS	0.5%	A087459 001
	1%	A087458 001
	2.5%	A087457 001
GEL;TOPICAL		
NUTRACORT		
HEALTHPOINT	1%	A084698 001
PENEDECORT		
ALLERGAN HERBERT	1%	A088215 001 Jun 06, 1984
INJECTABLE; INJECTION		
CORTEF		
PHARMACIA AND UPJOHN	50MG/ML	N009864 001
LOTION;TOPICAL		
ACTICORT		
BAKER NORTON	1%	A086535 001
ALA-CORT		
CROWN LABS	1%	A083201 001
BALNEOL-HC		
SOLVAY	1%	A088041 001 Dec 03, 1982
BETA-HC		
BETA DERMAC	1%	A089495 001 Jan 25, 1988
CETACORT		
DOW PHARM	0.5%	A080426 002
	1%	A080426 001
CORT-DOME		
BAYER PHARMS	0.5%	N009895 003
	1%	N009895 001
DERMACORT		
SOLVAY	0.5%	A084573 002
	1%	A086462 001
EPICORT		
BLULINE	0.5%	A083219 002
GLYCORT		
HERAN	1%	A087489 001 Oct 03, 1983
H-CORT		
PHARM ASSOC	0.5%	A086824 001
HYDROCORTISONE		
ALPHARMA US PHARMS	0.5%	A087317 001 Jun 07, 1982
	1%	A087315 001 Jun 07, 1982
MERICON	0.5%	A085282 001
	1%	A085282 002 Feb 26, 1987
NASKA	1%	A089705 001 Apr 25, 1988
PERRIGO NEW YORK	0.5%	A085662 001
	1%	A085663 001
TARO	1%	A089024 001 Feb 12, 1986

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-191(of 375)

\*\* See List Footnote

HYDROCORTISONE

LOTION;TOPICAL

HYTONE

VALEANT INTL	1%	A080473 003
	2.5%	A080473 004 Nov 30, 1982

NUTRACORT

DOW PHARM	0.5%	A080443 002
	1%	A080443 003
	2.5%	A087644 001 Aug 24, 1982

STIE-CORT

PERRIGO CO	1%	A089066 001 Nov 25, 1985
------------	----	--------------------------

OINTMENT;TOPICAL

CORTRIL

PFIZER GLOBAL	1%	N009176 001
	2.5%	N009176 002

HC (HYDROCORTISONE)

C AND M PHARMA	0.5%	A080481 001
	1%	A080481 002

HYDROCORTISONE

ALTANA	0.5%	A080489 002
	1%	A080489 003

AMBIX	1%	A086079 001
	2.5%	A086272 001

NASKA	1%	A089704 001 Mar 10, 1988
PERRIGO NEW YORK	0.5%	A084969 003

	1%	A085028 001
--	----	-------------

PHARMADERM	1%	A088842 001 Feb 09, 1987
------------	----	--------------------------

TARO	0.5%	A086256 001
------	------	-------------

	2.5%	A040310 001 Dec 29, 2000
--	------	--------------------------

USL PHARMA	1%	A088061 001 Sep 27, 1983
------------	----	--------------------------

	2.5%	A088039 001 Sep 27, 1983
--	------	--------------------------

HYTONE

DERMIK LABS	1%	A080474 003
	2.5%	A080474 004

PENECLORT

ALLERGAN HERBERT	2.5%	A088217 001 Jun 06, 1984
------------------	------	--------------------------

POWDER;FOR RX COMPOUNDING

H-CORT

TORCH	100%	A087834 001 Mar 29, 1982
-------	------	--------------------------

HYDRO-RX

X GEN PHARMS	100%	A085982 001
--------------	------	-------------

HYDROCORTISONE

PADDOCK LLC	100%	A088082 001 Apr 08, 1983
-------------	------	--------------------------

SOLUTION;TOPICAL

PENECLORT

ALLERGAN HERBERT	1%	A088214 001 Jun 06, 1984
------------------	----	--------------------------

TEXACORT

MISSION PHARMA	1%	A080425 001
----------------	----	-------------

TABLET;ORAL

CORTRIL

PFIZER	10MG	N009127 005
	20MG	N009127 003

HYDROCORTISONE

BARR	20MG	A083999 001
------	------	-------------

ELKINS SINN	20MG	A080624 001
-------------	------	-------------

FERRANTE	10MG	A080568 001
----------	------	-------------

	20MG	A080568 002
--	------	-------------

IMPAX LABS	20MG	A080781 001
------------	------	-------------

INWOOD LABS	20MG	A080732 001
-------------	------	-------------

LANNETT	20MG	A085070 001
---------	------	-------------

NEXGEN PHARMA INC	20MG	A083140 001
-------------------	------	-------------

PANRAY	10MG	N009659 001
--------	------	-------------

	20MG	N009659 002
--	------	-------------

PARKE DAVIS	20MG	A084243 001
-------------	------	-------------

PUREPAC PHARM	10MG	A084247 003 Aug 31, 1982
---------------	------	--------------------------

	20MG	A080395 001
--	------	-------------

	20MG	A084247 002
--	------	-------------

ROXANE	10MG	A088539 001 Mar 21, 1984
--------	------	--------------------------

SANDOZ	20MG	A080642 002
--------	------	-------------

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-192(of 375)

\*\* See List Footnote

**HYDROCORTISONE**

TABLET;ORAL

## HYDROCORTISONE

WATSON LABS	20MG	A080355 001
WHITEWORTH TOWN PLSN	10MG	A080344 001
	20MG	A080344 002

## HYDROCORTONE

MERCK	10MG	N008506 007
	20MG	N008506 011

TABLET;VAGINAL

## CORTRIL

PFIPHARMECS	10MG	N009796 001
-------------	------	-------------

**HYDROCORTISONE ACETATE**

CREAM;TOPICAL

## HEMSOL-HC

ABLE	1%	A081274 001 Jun 19, 1992
HYDROCORTISONE ACETATE		
CENCI	1%	A080419 001 Jan 25, 1982
FERNDALE LABS	2.5%	A040259 001 Jul 29, 1999
PARKE DAVIS	1%	A089914 001 Jan 03, 1989
PUREPAC PHARM	0.5%	A086050 001
	1%	A086052 001

## MICORT-HC

SEBELA IRELAND LTD	2%	A040398 001 Mar 29, 2002
--------------------	----	--------------------------

INJECTABLE;INJECTION

## CORTEF ACETATE

PHARMACIA AND UPJOHN	50MG/ML	N009378 002
----------------------	---------	-------------

## CORTRIL

PFIZER	25MG/ML	N009164 001
--------	---------	-------------

## HYDROCORTISONE ACETATE

AKORN	25MG/ML	N009637 001
	50MG/ML	N009637 002
BEL MAR	25MG/ML	A083739 001
	50MG/ML	A083739 002
WATSON LABS	25MG/ML	A083128 001
	25MG/ML	A083759 001
	50MG/ML	A083759 002
	50MG/ML	A085214 001

## HYDROCORTONE

MERCK	25MG/ML	N008228 001
	50MG/ML	N008228 004

LOTION;TOPICAL

## DRICORT

INGRAM PHARM	0.5%	A086207 001
--------------	------	-------------

OINTMENT;OPHTHALMIC

## HYDROCORTISONE ACETATE

FERA PHARMS	0.5%	A080828 001
-------------	------	-------------

OINTMENT;OPHTHALMIC, OTIC

## HYDROCORTONE

MERCK	1.5%	N009018 003
-------	------	-------------

OINTMENT;TOPICAL

## CORTEF ACETATE

PHARMACIA AND UPJOHN	1%	N008917 002
+	2.5% **	N008917 001

PASTE;TOPICAL

## ORABASE HCA

COLGATE	0.5%	A083205 001
---------	------	-------------

POWDER;FOR RX COMPOUNDING

## HYDROCORTISONE ACETATE

X GEN PHARMS	100%	A085981 001
--------------	------	-------------

**HYDROCORTISONE ACETATE; NEOMYCIN SULFATE**

CREAM;TOPICAL

## NEO-CORTEF

PHARMACIA AND UPJOHN	1%;EQ 3.5MG BASE/GM	A061049 001
	2.5%;EQ 3.5MG BASE/GM	A061049 002

OINTMENT;OPHTHALMIC

## NEO-CORTEF

PHARMACIA AND UPJOHN	0.5%;EQ 3.5MG BASE/GM	A060610 001
----------------------	-----------------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-193(of 375)

\*\* See List Footnote

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

OINTMENT;OPHTHALMIC NEO-CORTEF	1.5%;EQ 3.5MG BASE/GM	A060610 002
OINTMENT;TOPICAL NEO-CORTEF		
PHARMACIA AND UPJOHN	0.5%;EQ 3.5MG BASE/GM	A060751 001
	1%;EQ 3.5MG BASE/GM	A060751 002
	2.5%;EQ 3.5MG BASE/GM	A060751 003
SUSPENSION/DROPS;OPHTHALMIC		
COR-OTICIN		
AKORN	1.5%;EQ 3.5MG BASE/ML	A060188 001
NEO-CORTEF		
PHARMACIA AND UPJOHN	0.5%;EQ 3.5MG BASE/ML	A060612 002
	1.5%;EQ 3.5MG BASE/ML	A060612 001

HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE

SUSPENSION;OPHTHALMIC TERRA-CORTRIL PFIZER	1.5%;EQ 5MG BASE/ML	A061016 001
--	---------------------	-------------

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED;TOPICAL HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%		
VINTAGE PHARMS	1%;1%	A089440 001 May 17, 1988
LOTION;TOPICAL PRAMOSONE		
FERNDALE LABS	0.5%;1%	A083213 002

HYDROCORTISONE ACETATE; UREA

CREAM;TOPICAL CARMOL HC FOUGERA PHARMS	1%;10%	A080505 001
--	--------	-------------

HYDROCORTISONE BUTYRATE

CREAM;TOPICAL LOCOID YAMANOUCHI	0.1%	N018795 001 Jan 07, 1983
OINTMENT;TOPICAL LOCOID YAMANOUCHI	0.1%	N019106 001 Jul 03, 1984
SOLUTION;TOPICAL LOCOID YAMANOUCHI	0.1%	N019819 001 Sep 15, 1988

HYDROCORTISONE CYPIONATE

SUSPENSION;ORAL CORTEF		
PHARMACIA AND UPJOHN	EQ 10MG BASE/5ML	N009900 001

HYDROCORTISONE SODIUM PHOSPHATE

INJECTABLE;INJECTION HYDROCORTONE MERCK	EQ 50MG BASE/ML	N012052 001
---	-----------------	-------------

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE;INJECTION A-HYDROCORT ABBOTT	EQ 100MG BASE/VIAL	A085928 001
	EQ 100MG BASE/VIAL	A089577 001 Apr 11, 1989
	EQ 250MG BASE/VIAL	A089578 001 Apr 11, 1989
	EQ 500MG BASE/VIAL	A089579 001 Apr 11, 1989
	EQ 1GM BASE/VIAL	A089580 001 Apr 11, 1989
HOSPIRA	EQ 100MG BASE/VIAL	A040666 001 Apr 06, 2006
	EQ 100MG BASE/VIAL	A085929 001
	EQ 250MG BASE/VIAL	A085930 001
	EQ 500MG BASE/VIAL	A085931 001
	EQ 1GM BASE/VIAL	A085932 001

## HYDROCORTISONE SODIUM SUCCINATE

ABRAXIS PHARM	EQ 100MG BASE/VIAL	A088667 001 Jun 08, 1984
	EQ 100MG BASE/VIAL	A088712 001 Jun 08, 1984
	EQ 250MG BASE/VIAL	A088668 001 Jun 08, 1984
	EQ 500MG BASE/VIAL	A088669 001 Jun 08, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-194(of 375)

\*\* See List Footnote

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

HYDROCORTISONE SODIUM SUCCINATE

BAXTER HLTHCARE	EQ 1GM BASE/VIAL EQ 100MG BASE/VIAL EQ 250MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 100MG BASE/VIAL EQ 100MG BASE/VIAL EQ 250MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A088670 001 Jun 08, 1984 A086619 001 A087567 001 A087568 001 A087569 001 A087532 001 Mar 19, 1982 A084737 002 A084738 001 A084737 001 A084747 001 A084748 001
-----------------	--	---

HYDROCORTISONE VALERATE

CREAM;TOPICAL

HYDROCORTISONE VALERATE

G AND W LABS INC	0.2%	A074489 001 Aug 12, 1998
WESTCORT		
+ RANBAXY LABS LTD	0.2% **	N017950 001
OINTMENT;TOPICAL		
HYDROCORTISONE VALERATE		
FOUGERA PHARMS	0.2%	A075085 001 Jul 31, 2001
WESTCORT		
+ RANBAXY	0.2% **	N018726 001 Aug 08, 1983

HYDROCORTISONE; NEOMYCIN SULFATE

CREAM;TOPICAL

NEO-CORT-DOME

BAYER PHARMS	0.5%;EQ 3.5MG BASE/GM 1%;EQ 3.5MG BASE/GM	N050237 006 Jun 05, 1984 N050237 005 Jun 05, 1984
--------------	--	--

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062394 001 Sep 29, 1982
OTOCORT		

WATSON LABS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A060730 002

SUSPENSION/DROPS;OPHTHALMIC

CORTISPORIN

MONARCH PHARMS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N050169 001
NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE		

PHARMAFAIR 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A062623 001 Sep 24, 1985

SUSPENSION/DROPS;OTIC

NEOMYCIN SULFATE, POLYMYXIN B SULFATE &amp; HYDROCORTISONE

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062617 001 Sep 18, 1985
OTICAIR		

PHARMAFAIR 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A062399 001 Nov 18, 1982

OTOBIONE

SCHERING	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A061816 001
OTOCORT		

ACTAVIS LABS FL INC 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

A062521 001 Jul 11, 1985

PEDIOTIC

MONARCH PHARMS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	A062822 001 Sep 29, 1987
----------------	-------------------------------------	--------------------------

MONARCH PHARMS 1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML

HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS;OTIC

OTOBIOPTIC

SCHERING	5MG/ML;EQ 10,000 UNITS BASE/ML	A062302 001
PYOCIDIN		

FOREST LABS 5MG/ML;EQ 10,000 UNITS BASE/ML

A061606 001

HYDROCORTISONE; TETRACYCLINE HYDROCHLORIDE

OINTMENT;OPHTHALMIC

ACHROMYCIN

LEDERLE	1.5%;1%	N050272 001
---------	---------	-------------

LEDERLE 1.5%;1%

**DISCONTINUED DRUG PRODUCT LIST**

6-195(of 375)

\*\* See List Footnote

HYDROCORTISONE; UREA

CREAM;TOPICAL		
ALPHADERM		
BIOGLAN	1%;10%	A086008 001
CALMURID HC		
PHARMACIA AND UPJOHN	1%;10%	A083947 001

HYDROFLUMETHIAZIDE

TABLET;ORAL		
DIUCARDIN		
WYETH AYERST	50MG	A083383 001
HYDROFLUMETHIAZIDE		
PAR PHARM	50MG	A088850 001 May 31, 1985
WATSON LABS	50MG	A088031 001 Apr 06, 1983
	50MG	A088528 001 Aug 15, 1984
SALURON		
+ SHIRE LLC	50MG	N011949 001

HYDROFLUMETHIAZIDE; RESERPINE

TABLET;ORAL		
HYDROFLUMETHIAZIDE AND RESERPINE		
USL PHARMA	50MG;0.125MG	A088195 001 Oct 26, 1983
WATSON LABS	25MG;0.125MG	A088127 001 Mar 22, 1983
	50MG;0.125MG	A088110 001 Mar 22, 1983
RESERPINE AND HYDROFLUMETHIAZIDE		
IVAX PHARMS	50MG;0.125MG	A088932 001 Jan 11, 1985
PAR PHARM	50MG;0.125MG	A088907 001 Sep 20, 1985
SALUTENSIN		
SHIRE	50MG;0.125MG	N012359 003
SALUTENSIN-DEMI		
SHIRE	25MG;0.125MG	N012359 004

HYDROMORPHONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL		
PALLADONE		
PURDUE PHARMA LP	12MG	N021044 001 Sep 24, 2004
	16MG	N021044 002 Sep 24, 2004
	24MG	N021044 003 Sep 24, 2004
	32MG	N021044 004 Sep 24, 2004

## INJECTABLE;INJECTION

DILAUDID-HP		
FRESENIUS KABI USA	250MG/VIAL	N019034 002 Aug 04, 1994
HYDROMORPHONE HYDROCHLORIDE		
HOSPIRA	10MG/ML	A074598 001 Jun 19, 1997
WATSON LABS	10MG/ML	A074317 001 Aug 23, 1995

## TABLET;ORAL

HYDROMORPHONE HYDROCHLORIDE		
NESHER PHARMS	2MG	A077311 001 Nov 09, 2005
	4MG	A077311 002 Nov 09, 2005
	8MG	A077311 003 Nov 09, 2005

HYDOXOCOBALAMIN

INJECTABLE;INJECTION		
ALPHAREDISOL		
MERCK	1MG/ML	A080778 001
CYANOKIT		
SERB SA	2.5GM/VIAL (5GM/KIT)	N022041 002 Dec 15, 2006
HYDOXOCOBALAMIN		
ABRAXIS PHARM	1MG/ML	A084921 001
WATSON LABS	1MG/ML	A085528 001
HYDROXOMIN		
BEL MAR	1MG/ML	A084629 001

HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS;OPHTHALMIC		
PAREDRINE		
PHARMICS	1%	N000004 004

**DISCONTINUED DRUG PRODUCT LIST**

6-196(of 375)

\*\* See List Footnote

HYDROXYCHLOROQUINE SULFATE

TABLET;ORAL

HYDROXYCHLOROQUINE SULFATE

SANDOZ

200MG

A040150 001 Jan 27, 1996

HYDROXYPROGESTERONE CAPROATE

INJECTABLE;INJECTION

DELALUTIN

+	BRISTOL MYERS SQUIBB	125MG/ML **
+		125MG/ML **
+		250MG/ML **
+		250MG/ML **

N010347 004  
N016911 001  
N010347 002  
N016911 002

HYDROXYPROGESTERONE CAPROATE

AKORN

125MG/ML

N018004 001

ALLERGAN SALES LLC

125MG/ML

N017439 001

250MG/ML

N017439 002

HYDROXYSTILBAMIDINE ISETHIONATE

INJECTABLE;INJECTION

HYDROXYSTILBAMIDINE ISETHIONATE

SANOFI AVENTIS US 225MG/AMP

N009166 001

HYDROXYUREA

CAPSULE;ORAL

HYDROXYUREA

BARR

250MG

A075143 002 Sep 21, 2000

BARR LABS INC

250MG

A075020 002 Jun 26, 2000

500MG

A075020 001 Jul 30, 1998

ROXANE

500MG

A074476 001 Aug 18, 1995

TABLET;ORAL

HYDROXYUREA

BARR

1GM

A075734 001 Aug 29, 2000

HYDROXYZINE HYDROCHLORIDE

INJECTABLE;INJECTION

HYDROXYZINE

BAXTER HLTHCARE

50MG/ML

A085551 002

HYDROXYZINE HYDROCHLORIDE

ALTANA

25MG/ML

A087273 001 Apr 20, 1982

50MG/ML

A087273 002 Apr 20, 1982

BAXTER HLTHCARE

25MG/ML

A085551 001

FRESENIUS KABI USA

25MG/ML

A088184 001 Mar 31, 1983

50MG/ML

A088185 001 Mar 31, 1983

HOSPIRA

25MG/ML

A087416 001

50MG/ML

A086821 001

PHARMAFAIR

50MG/ML

A087546 001

25MG/ML

A088862 001 Feb 14, 1986

25MG/ML

A089106 001 Feb 14, 1986

50MG/ML

A088881 001 Feb 14, 1986

50MG/ML

A089107 001 Feb 14, 1986

SMITH AND NEPHEW

25MG/ML

A087592 001

SOLOPAK

25MG/ML

A086822 001

25MG/ML

A087591 001

50MG/ML

A087310 001

50MG/ML

A087593 001

50MG/ML

A087595 001

50MG/ML

A087596 001

WATSON LABS

25MG/ML

A085778 001

25MG/ML

A087274 001

50MG/ML

A085779 001

50MG/ML

A087274 002

WYETH AYERST

25MG/ML

A086258 001

50MG/ML

A086258 002

ORGATRAX

ORGANON USA INC

25MG/ML

A087014 001

50MG/ML

A087014 002

VISTARIL

+ PFIZER

25MG/ML \*\*

N011111 001

+

50MG/ML \*\*

N011111 002

**DISCONTINUED DRUG PRODUCT LIST**

6-197(of 375)

\*\* See List Footnote

HYDROXYZINE HYDROCHLORIDE

SYRUP;ORAL

ATARAX

ROERIG	10MG/5ML **
HYDROXYZINE HYDROCHLORIDE	
ALPHARMA US PHARMS	10MG/5ML
KV PHARM	10MG/5ML
STI PHARMA LLC	10MG/5ML

N010485 001

A088785 001	Feb 03, 1988
A087730 001	Jul 01, 1982
A086880 001	

TABLET;ORAL

ATARAX

PFIZER	10MG **
	25MG **
	50MG **
	100MG **

N010392 001	
N010392 004	
N010392 006	
N010392 005	

HYDROXYZINE HYDROCHLORIDE

ABLE

10MG	A040559 001	Jul 22, 2004
25MG	A040562 001	Jul 22, 2004
50MG	A040563 001	Jul 22, 2004

ACTAVIS ELIZABETH

10MG	A089071 001	Jul 22, 1986
25MG	A089072 001	Jul 22, 1986
50MG	A089073 001	Jul 22, 1986

AUROLIFE PHARMA LLC

10MG	A087871 002	Dec 20, 1982
25MG	A087871 003	Dec 20, 1982
50MG	A087871 001	Dec 20, 1982

HALSEY

10MG	A089366 001	May 02, 1988
25MG	A089117 001	May 02, 1988
50MG	A089396 001	May 02, 1988

IVAX PHARMS

10MG	A087216 001	
25MG	A087410 001	
50MG	A087411 001	

KV PHARM

10MG	A087819 001	Jun 23, 1982
25MG	A087820 001	Jun 23, 1982
50MG	A087821 001	Jun 23, 1982
100MG	A087822 001	Jun 23, 1982

MUTUAL PHARM

10MG	A088409 001	Nov 15, 1983
25MG	A087857 001	Apr 18, 1983
50MG	A087860 001	Apr 18, 1983

PLIVA

100MG	A081054 001	Sep 25, 1995
10MG	A088120 001	Sep 25, 1984
25MG	A088121 001	Sep 25, 1984
50MG	A088122 001	Sep 25, 1984

PUREPAC PHARM

10MG	A088540 001	Oct 22, 1985
25MG	A088551 001	Oct 22, 1985
50MG	A088529 001	Oct 22, 1985

QUANTUM PHARMICS

10MG	A088246 002	
25MG	A085247 001	
50MG	A087245 001	

SANDOZ

10MG	A040899 001	Jun 10, 2008
25MG	A040899 002	Jun 10, 2008
50MG	A040899 003	Jun 10, 2008

SUN PHARM INDS INC

10MG	A089381 001	May 19, 1986
25MG	A089382 001	May 19, 1986
50MG	A089383 001	May 19, 1986
100MG	A087862 001	Apr 18, 1983

SUN PHARM INDUSTRIES

10MG	A088794 001	Dec 05, 1984
25MG	A088795 001	Dec 05, 1984
50MG	A088796 001	Dec 05, 1984
100MG	A089121 001	Mar 20, 1986

SUPERPHARM

10MG	A089122 001	Mar 20, 1986
25MG	A089123 001	Mar 20, 1986
50MG	A087602 001	Jan 22, 1982
100MG	A087603 001	Jan 22, 1982

USL PHARMA

10MG	A087604 001	Jan 22, 1982
25MG	A087605 001	Jan 22, 1982
50MG	A087606 001	Jan 22, 1982
100MG	A087607 001	Jan 22, 1982

VINTAGE

10MG	A087608 001	Jan 22, 1982
25MG	A087609 001	Jan 22, 1982
50MG	A087610 001	Jan 22, 1982
100MG	A087611 001	Jan 22, 1982

WATSON LABS

10MG	A081149 001	Mar 18, 1994
10MG	A086827 001	
10MG	A088348 001	Sep 15, 1983
25MG	A081150 001	Mar 18, 1994
25MG	A086829 001	
25MG	A088349 001	Sep 15, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-198(of 375)

\*\* See List Footnote

HYDROXYZINE HYDROCHLORIDE

TABLET;ORAL

HYDROXYZINE HYDROCHLORIDE

50MG	A081151 001 Mar 18, 1994
50MG	A086836 001
50MG	A088350 001 Sep 15, 1983

HYDROXYZINE PAMOATE

CAPSULE;ORAL

HY-PAM "25"

TEVA

EQ 25MG HCL

A088713 001 Mar 04, 1985

HYDROXYZINE PAMOATE

DURAMED PHARMS BARR

EQ 25MG HCL

A088593 001 Feb 29, 1984

EQ 50MG HCL

A088594 001 Feb 29, 1984

EQ 100MG HCL

A088595 001 Feb 29, 1984

IVAX SUB TEVA PHARMS

EQ 25MG HCL

A087761 001 Mar 05, 1982

EQ 50MG HCL

A087760 001 Mar 05, 1982

PAR PHARM

EQ 25MG HCL

A087656 001 Jun 11, 1982

EQ 25MG HCL

A089145 001 Mar 17, 1986

EQ 50MG HCL

A087657 001 Jun 11, 1982

EQ 50MG HCL

A089146 001 Mar 17, 1986

EQ 100MG HCL

A087658 001 Jun 11, 1982

SANDOZ

EQ 25MG HCL

A081127 001 Jun 28, 1991

EQ 50MG HCL

A081128 001 Jun 28, 1991

EQ 100MG HCL

A081129 001 Jun 28, 1991

SUPERPHARM

EQ 25MG HCL

A089031 001 Jan 02, 1987

EQ 50MG HCL

A089032 001 Jan 02, 1987

EQ 100MG HCL

A089033 001 Jan 02, 1987

VANGARD

EQ 25MG HCL

A088392 001 Sep 19, 1983

EQ 50MG HCL

A088393 001 Sep 19, 1983

WATSON LABS

EQ 25MG HYDROCHLORIDE

A081165 001 Jul 31, 1991

EQ 25MG HCL

A086698 001

EQ 25MG HCL

A086840 001 Jul 01, 1982

EQ 50MG HCL

A086695 001

EQ 50MG HCL

A086705 001 Jul 01, 1982

EQ 50MG HCL

A087767 001 Aug 16, 1982

EQ 100MG HCL

A086697 001

EQ 100MG HCL

A086728 001 Oct 05, 1982

EQ 100MG HCL

A087790 001 Aug 16, 1982

VISTARIL

PFIZER

EQ 100MG HCL \*\*

N011459 006

SUSPENSION;ORAL

VISTARIL

PFIZER

EQ 25MG HCL/5ML

N011795 001

IBANDRONATE SODIUM

TABLET;ORAL

BONIVA

+ HOFFMANN LA ROCHE

EQ 2.5MG BASE \*\*

N021455 001 May 16, 2003

IBANDRONATE SODIUM

MYLAN PHARMS INC

EQ 150MG BASE

A078995 001 Mar 19, 2012

IBUPROFEN

CAPSULE;ORAL

IBUPROFEN

CONTRACT PHARMACAL

200MG

A074782 001 Jul 06, 1998

MIDOL

BAYER

200MG \*\*

A070626 001 Sep 02, 1987

200MG \*\*

A071002 001 Sep 02, 1987

SOLUTION;INTRAVENOUS

CALDOLOR

CUMBERLAND PHARMS

400MG/4ML (100MG/ML)

N022348 001 Jun 11, 2009

SUSPENSION;ORAL

CHILDREN'S ADVIL

WYETH CONS

100MG/5ML

N019833 002 Sep 19, 1989

IBU

ABBOTT

100MG/5ML

N019784 001 Dec 18, 1989

MOTRIN

+ MCNEIL CONSUMER

100MG/5ML \*\*

N019842 001 Sep 19, 1989

**DISCONTINUED DRUG PRODUCT LIST**

6-199(of 375)

\*\* See List Footnote

**IBUPROFEN**

SUSPENSION/DROPS;ORAL

MOTRIN

MCNEIL

40MG/ML

N020476 001 May 25, 1995

TABLET;ORAL

ACHE-S-N-PAIN

LEDERLE

200MG

A071065 001 May 28, 1987

CAP-PROFEN

PERRIGO

200MG

A072097 001 Dec 08, 1987

IBU

BASF

400MG

A070083 001 Feb 22, 1985

400MG

N018197 001

600MG

A070088 001 Feb 08, 1985

600MG

A070099 001 Mar 29, 1985

800MG

A070745 001 Jul 23, 1986

IBU-TAB

ALRA

800MG

A071965 001 Aug 11, 1988

IBUPRIN

PLIVA

200MG

A071773 001 Jul 16, 1987

IBUPROFEN

ABBOTT

600MG

A070556 001 Jun 14, 1985

800MG

A071264 001 Jul 25, 1986

ANI PHARMS INC

200MG

A071144 001 Jan 20, 1987

200MG

A072901 001 Dec 19, 1991

200MG

A072903 001 Dec 19, 1991

AUROLIFE PHARMA LLC

300MG

A070736 002 Jun 12, 1986

400MG

A070736 003 Jun 12, 1986

600MG

A070736 001 Jun 12, 1986

800MG

A071938 001 Jan 14, 1988

CONTRACT PHARMACAL

200MG

A071735 001 Sep 10, 1987

200MG

A073691 001 Feb 25, 1994

200MG

A074931 001 Jul 20, 1998

HALSEY

200MG

A071027 001 Sep 29, 1987

300MG

A071028 001 Mar 23, 1987

400MG

A071029 001 Mar 23, 1987

600MG

A071030 001 Mar 23, 1987

800MG

A072137 001 Feb 05, 1988

IVAX SUB TEVA PHARMS

200MG

A071154 001 Oct 27, 1987

200MG

A072040 001 Apr 29, 1988

400MG

A071145 001 Sep 23, 1986

600MG

A071146 001 Sep 23, 1986

800MG

A071769 001 May 08, 1987

J AND J CONSUMER INC

400MG

A070081 001 Jun 16, 1986

LEDERLE

400MG

A070629 001 Sep 19, 1986

600MG

A070630 001 Sep 19, 1986

LEINER

300MG

A071266 001 Oct 15, 1986

MCNEIL

600MG

A070476 001 Jun 16, 1986

MYLAN

200MG

A071870 001 May 05, 1988

600MG

A070057 001 Sep 24, 1985

800MG

A071999 001 Dec 03, 1987

MYLAN PHARMS INC

400MG

A070045 001 Sep 24, 1985

NORTHSTAR HLTHCARE

400MG

A078132 001 Sep 10, 2007

600MG

A078132 002 Sep 10, 2007

800MG

A078132 003 Sep 10, 2007

OHM LABS

400MG

A070818 001 Dec 26, 1985

PAR PHARM

200MG

A071575 001 May 08, 1987

300MG

A070328 001 Aug 06, 1985

400MG

A070329 001 Aug 06, 1985

600MG

A070330 001 Aug 06, 1985

800MG

A070986 001 Jul 25, 1986

PERRIGO

200MG

A072098 001 Dec 08, 1987

PLIVA

400MG

A071666 001 Jun 18, 1987

600MG

A071667 001 Jun 18, 1987

800MG

A071668 001 Jun 18, 1987

PUREPAC PHARM

200MG

A071122 001 Oct 03, 1986

200MG

A071664 001 Feb 03, 1987

300MG

A071123 001 Sep 19, 1986

400MG

A071124 001 Sep 19, 1986

600MG

A071125 001 Sep 19, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-200(of 375)

\*\* See List Footnote

**IBUPROFEN**TABLET;ORAL  
IBUPROFEN

SANDOZ	800MG	A071964 001	Feb 01, 1988
	200MG	A070733 001	Sep 19, 1986
	200MG	A071807 001	Feb 25, 1988
	200MG	A074525 001	Dec 15, 1995
	200MG	A074533 001	Dec 15, 1995
	400MG	A072064 001	Jan 14, 1988
	600MG	A072065 001	Jan 14, 1988
	800MG	A072169 001	Dec 11, 1987
SUN PHARM INDUSTRIES	200MG	A070493 001	Dec 24, 1985
	200MG	A070908 001	Sep 26, 1986
	200MG	A071462 001	Oct 02, 1986
	400MG	A070079 001	Jul 24, 1985
	600MG	A070080 001	Jul 24, 1985
	800MG	A071448 001	Feb 18, 1987
SUPERPHARM	600MG	A070709 001	Apr 25, 1986
TEVA	200MG	A073141 001	May 29, 1992
	400MG	A073343 001	Jun 30, 1992
	600MG	A073344 001	Jun 30, 1992
	800MG	A073345 001	Jun 30, 1992
VINTAGE PHARMS	200MG	A072249 001	Jan 10, 1989
	300MG	A071230 001	Oct 22, 1986
	400MG	A071231 001	Oct 22, 1986
	600MG	A071232 001	Oct 22, 1986
	800MG	A072004 001	Nov 18, 1987
WATSON LABS	200MG	A070435 001	Mar 05, 1986
	200MG	A071765 001	Sep 04, 1987
	200MG	A071905 001	Mar 08, 1988
	300MG	A071338 001	Dec 01, 1986
	400MG	A070038 001	Sep 06, 1985
	400MG	A070436 001	Aug 21, 1985
	600MG	A070041 001	Sep 06, 1985
	600MG	A070437 001	Aug 21, 1985
	800MG	A071547 001	Jul 02, 1987
	800MG	A071911 001	Oct 13, 1987
<b>IBUPROHM</b>			
OHM LABS	400MG	A070469 001	Aug 29, 1985
<b>MEDIPREN</b>			
MCNEIL	200MG	A070475 001	Feb 06, 1986
	200MG	A071215 001	Jun 26, 1986
<b>MIDOL</b>			
BAYER	200MG	A070591 001	Sep 02, 1987
	200MG	A071001 001	Sep 02, 1987
<b>MOTRIN</b>			
+ MCNEIL CONSUMER	300MG **	N017463 003	
+	400MG **	N017463 002	
+	600MG **	N017463 004	
+	800MG **	N017463 005	May 22, 1985
MCNEIL PED	100MG	N020418 001	Nov 16, 1994
MOTRIN MIGRAINE PAIN			
J AND J CONSUMER INC	200MG	N019012 004	Feb 25, 2000
<b>NUPRIN</b>			
BRISTOL MYERS	200MG	A072035 001	Feb 16, 1988
	200MG	A072036 001	Feb 16, 1988
J AND J CONSUMER INC	200MG	N019012 001	May 18, 1984
	200MG	N019012 002	Jul 29, 1987
<b>RUFEN</b>			
BASF	600MG	N018197 002	Mar 05, 1984
<b>TABLET, CHEWABLE;ORAL</b>			
<b>MOTRIN</b>			
MCNEIL PED	50MG	N020135 001	Nov 16, 1994
	100MG	N020135 002	Nov 16, 1994

**DISCONTINUED DRUG PRODUCT LIST**

6-201(of 375)

\*\* See List Footnote

IBUPROFEN; OXYCODONE HYDROCHLORIDE

TABLET;ORAL

COMBUNOX

FOREST LABS 400MG;5MG \*\*

N021378 001 Nov 26, 2004

OXYCODONE HYDROCHLORIDE AND IBUPROFEN

WATSON LABS 400MG;5MG

A078394 001 Nov 26, 2007

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET;ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

CONTRACT PHARMACAL 200MG;30MG

A075588 001 Apr 08, 2002

IDARUBICIN HYDROCHLORIDE

INJECTABLE;INJECTION

IDAMYCIN

PHARMACIA AND UPJOHN 5MG/VIAL

N050661 002 Sep 27, 1990

10MG/VIAL

N050661 001 Sep 27, 1990

20MG/VIAL

N050661 003 Apr 25, 1995

IDARUBICIN HYDROCHLORIDE

SANDOZ 1MG/ML

A091293 001 Mar 29, 2011

TEVA PARENTERAL 5MG/VIAL

A065037 003 May 01, 2002

10MG/VIAL

A065037 002 May 01, 2002

20MG/VIAL

A065037 001 May 01, 2002

IDOXURIDINE

OINTMENT;OPHTHALMIC

STOXIL

GLAXOSMITHKLINE 0.5%

N015868 001

SOLUTION/DROPS;OPHTHALMIC

DENDRID

+ ALCON 0.1%

N014169 001

HERPLEX

ALLERGAN 0.1%

N013935 002

STOXIL

GLAXOSMITHKLINE 0.1%

N013934 001

IFOSFAMIDE; MESNA

INJECTABLE;INJECTION

IFEX/MESNEX KIT

BAXTER HLTHCARE

1GM/VIAL;100MG/ML

N019763 003 Oct 10, 1992

3GM/VIAL;100MG/ML

N019763 004 Oct 10, 1992

INJECTABLE;INTRAVENOUS

IFOSFAMIDE/MESNA KIT

TEVA PHARMS USA

1GM/20ML;1GM/10ML (50MG/ML;100MG/ML)

A075874 001 Feb 26, 2002

3GM/60ML;1GM/10ML (50MG/ML;100MG/ML)

A075874 002 Feb 26, 2002

ILOPROST

SOLUTION;INHALATION

VENTAVIS

ACTELION PHARMS LTD 20MCG/2ML (10MCG/ML)

N021779 001 Dec 29, 2004

IMATINIB MESYLATE

CAPSULE;ORAL

GLEEVEC

+ NOVARTIS

EQ 50MG BASE \*\*

N021335 001 May 10, 2001

+

EQ 100MG BASE \*\*

N021335 002 May 10, 2001

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE;ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS 25MG/ML

A086765 001

INJECTABLE;INJECTION

TOFRANIL

NOVARTIS 12.5MG/ML

N011838 002

TABLET;ORAL

IMIPRAMINE HYDROCHLORIDE

LEDERLE 10MG

A086269 001

25MG

A086267 001

50MG

A086268 001

OXFORD PHARMS 50MG

A040751 001 Feb 28, 2008

PAR PHARM 10MG

A089422 001 Jul 14, 1987

25MG

A089497 001 Jul 14, 1987

ROXANE 10MG

A083799 001

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-202(of 375)

\*\* See List Footnote

IMIPRAMINE HYDROCHLORIDE

TABLET;ORAL

IMIPRAMINE HYDROCHLORIDE

	25MG	A083799 002
	50MG	A083799 003
SANDOZ	10MG	A085200 001
	25MG	A084869 002
	50MG	A085133 001
TEVA	10MG	A083729 001
	25MG	A083729 004
	50MG	A083729 003
USL PHARMA	25MG	A087776 001 Feb 10, 1982
VANGARD	10MG	A088036 001 Nov 03, 1982
	25MG	A087619 001 Feb 09, 1982
	50MG	A087631 001 Jan 04, 1982
WATSON LABS	10MG	A085220 001
	10MG	A085875 001
	25MG	A084252 002
	25MG	A085878 001
	50MG	A085221 001
	50MG	A085877 001
WEST WARD	25MG	A088222 001 May 26, 1983
	50MG	A088223 001 May 26, 1983
JANIMINE		
ABBOTT	10MG	N017895 001
	25MG	N017895 002
	50MG	N017895 003
PRAMINE		
ALRA	10MG	A083827 001
	25MG	A083827 002
	50MG	A083827 003
PRESAMINE		
SANOFI AVENTIS US	10MG	N011836 006
	25MG	N011836 003
	50MG	N011836 007

IMIPRAMINE PAMOATE

CAPSULE;ORAL

IMIPRAMINE PAMOATE

MYLAN PHARMS INC	EQ 75MG HCL	A202338 001 Jun 28, 2013
	EQ 100MG HCL	A202338 002 Jun 28, 2013
	EQ 125MG HCL	A202338 003 Jun 28, 2013
	EQ 150MG HCL	A202338 004 Jun 28, 2013
TOFRANIL-PM		
+ SPECGX LLC	EQ 75MG HCL **	N017090 001
+	EQ 100MG HCL **	N017090 004
+	EQ 125MG HCL **	N017090 003
+	EQ 150MG HCL **	N017090 002

INAMRINONE LACTATE

INJECTABLE;INJECTION

AMRINONE LACTATE

BAXTER HLTHCARE CORP	EQ 5MG BASE/ML	A075542 001 May 10, 2000
HOSPIRA	EQ 5MG BASE/ML	A074616 001 Aug 03, 1998
INOCOR		
SANOFI AVENTIS US	EQ 5MG BASE/ML	N018700 001 Jul 31, 1984

INDAPAMIDE

TABLET;ORAL

INDAPAMIDE

ANI PHARMS INC	1.25MG	A074498 002 Feb 12, 1998
	2.5MG	A074498 001 Oct 31, 1996
FOSUN PHARMA	1.25MG	A074594 001 May 23, 1996
	2.5MG	A074594 002 May 23, 1996
MYLAN PHARMS INC	1.25MG	A075105 001 Jul 23, 1998
	2.5MG	A075105 002 Jul 23, 1998
TEVA	1.25MG	A074665 001 Apr 04, 1997
	2.5MG	A074665 002 Apr 04, 1997
WATSON LABS	1.25MG	A074585 001 Sep 26, 1996
	2.5MG	A074585 002 Sep 26, 1996

**DISCONTINUED DRUG PRODUCT LIST**

6-203(of 375)

\*\* See List Footnote

INDAPAMIDE

TABLET;ORAL

LOZOL

+ SANOFI AVENTIS US	1.25MG **
+	2.5MG **

N018538 002	Apr 29, 1993
N018538 001	Jul 06, 1983

INDECAINIDE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

DECABID

LILLY	EQ 50MG BASE
	EQ 75MG BASE
	EQ 100MG BASE

N019693 001	Dec 29, 1989
N019693 002	Dec 29, 1989
N019693 003	Dec 29, 1989

INDINAVIR SULFATE

CAPSULE;ORAL

CRIXIVAN

MERCK SHARP DOHME	EQ 100MG BASE
	EQ 333MG BASE

N020685 006	Apr 19, 2000
N020685 005	Dec 17, 1998

INDOCYANINE GREEN

INJECTABLE;INJECTION

IC-GREEN

AKORN	10MG/VIAL
	40MG/VIAL
	50MG/VIAL

N011525 003	
N011525 004	
N011525 002	

INDOMETHACIN

CAPSULE;ORAL

INDO-LEMMON

TEVA	25MG
	50MG

A070266 001	Nov 07, 1985
A070267 001	Nov 07, 1985

INDOCIN

+ IROKO PHARMS LLC	25MG **
+	50MG **

N016059 001	
N016059 002	

INDOMETHACIN

ABLE	25MG
	50MG

A076666 001	Dec 17, 2003
A076666 002	Dec 17, 2003

CHARTWELL MOLECULES	25MG
	50MG

N018829 002	Aug 06, 1984
A070651 001	Mar 05, 1986

CYCLE PHARMS LTD	25MG
	50MG

N018829 001	Aug 06, 1984
A070353 001	Jun 18, 1985

DURAMED PHARMS BARR	25MG
	50MG

A070354 001	Jun 18, 1985
A070326 001	Oct 18, 1985

HALSEY	25MG
	50MG

A070327 001	Oct 18, 1985
A070782 001	Jun 03, 1987

IVAX SUB TEVA PHARMS	25MG
	50MG

A070635 001	Jun 03, 1987
N018730 001	May 04, 1984

MUTUAL PHARM	25MG
	50MG

N018806 001	Oct 03, 1986
A070067 001	Oct 03, 1986

MYLAN	50MG
PARKE DAVIS	25MG

A070068 001	Oct 03, 1986
N018858 002	Apr 20, 1984

PIONEER PHARMS	25MG
	50MG

A070813 001	Aug 11, 1986
A070592 001	Aug 11, 1986

PLIVA	25MG
	50MG

A071148 001	Mar 18, 1987
A071149 001	Mar 18, 1987

SUN PHARM INDUSTRIES	25MG
	50MG

A070900 002	Feb 09, 1987
A070900 001	Feb 09, 1987

SUPERPHARM	25MG
	50MG

A070487 001	Oct 10, 1986
A070488 001	Oct 10, 1986

TEVA	25MG
	50MG

A071342 001	Apr 18, 1988
A071343 001	Apr 18, 1988

WATSON LABS	25MG
	25MG

A070529 001	Oct 18, 1985
A070784 001	Aug 20, 1986

	25MG
	50MG

A072996 001	Jul 31, 1991
N018690 001	Jul 31, 1984

	50MG
	50MG

A070530 001	Oct 18, 1985
A070785 001	Aug 20, 1986

	50MG
	50MG

A071635 001	May 18, 1987
A072997 001	Jul 31, 1991

**DISCONTINUED DRUG PRODUCT LIST**

6-204(of 375)

\*\* See List Footnote

INDOMETHACIN

CAPSULE;ORAL INDOMETHACIN	50MG	N018690 002 Jul 31, 1984
CAPSULE, EXTENDED RELEASE;ORAL INDOCIN SR + IROKO PHARMS	75MG **	N018185 001 Feb 23, 1982
INDOMETHACIN ABLE INWOOD LABS WATSON LABS INC	75MG 75MG 75MG	A076114 001 Feb 06, 2002 A072410 001 Mar 15, 1989 A202572 001 Dec 09, 2013
SUPPOSITORY;RECTAL INDOCIN + IROKO PHARMS	50MG **	N017814 001 Aug 13, 1984
SUSPENSION;ORAL INDOMETHACIN CYCLE PHARMS LTD	25MG/5ML	A071412 001 Mar 18, 1987

INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE;SUBCUTANEOUS NOVOLOG MIX 50/50 NOVO NORDISK INC	50 UNITS/ML;50 UNITS/ML	N021810 001 Aug 26, 2008
NOVOLOG MIX 70/30 PENFILL NOVO NORDISK INC	210 UNITS/3ML;90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML) 210 UNITS/3ML;90 UNITS/3ML (70 UNITS/ML; 30 UNITS/ML)	N021172 002 Nov 01, 2001 N021172 003 Nov 01, 2001

INSULIN ASPART RECOMBINANT

INJECTABLE;SUBCUTANEOUS NOVOLOG INNOLET NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N020986 004 Apr 23, 2004
--	------------------------------	--------------------------

INSULIN DETEMIR RECOMBINANT

INJECTABLE;SUBCUTANEOUS LEVEMIR FLEXPEN NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N021536 002 Jun 16, 2005
LEVEMIR INNOLET NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N021536 003 Jun 16, 2005
LEVEMIR PENFILL NOVO NORDISK INC	300 UNITS/3ML (100 UNITS/ML)	N021536 004 Jun 16, 2005

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION HUMALOG MIX 50/50 PEN LILLY	50 UNITS/ML;50 UNITS/ML	N021018 003 Dec 22, 1999
HUMALOG MIX 75/25 PEN LILLY	75 UNITS/ML;25 UNITS/ML	N021017 003 Dec 22, 1999

INSULIN LISPRO RECOMBINANT

INJECTABLE;INJECTION HUMALOG PEN LILLY	100 UNITS/ML	N020563 002 Aug 06, 1998
--	--------------	--------------------------

INSULIN PORK

INJECTABLE;INJECTION ILETIN I LILLY	500 UNITS/ML	N017931 001
INSULIN NOVO NORDISK INC	40 UNITS/ML	N017926 001
REGULAR INSULIN NOVO NORDISK INC	100 UNITS/ML	N017926 003

INSULIN PURIFIED BEEF

INJECTABLE;INJECTION REGULAR ILETIN II LILLY	100 UNITS/ML	N018478 001
--	--------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-205(of 375)

\*\* See List Footnote

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

ILETIN II

LILLY 500 UNITS/ML N018344 002

REGULAR ILETIN II (PORK)

LILLY 100 UNITS/ML N018344 001

REGULAR PURIFIED PORK INSULIN

NOVO NORDISK INC 100 UNITS/ML N018381 001

VELOSULIN

NOVO NORDISK INC 100 UNITS/ML N018193 001

INSULIN PURIFIED PORK; INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN NORDISK MIXTARD (PORK)

NOVO NORDISK INC 30 UNITS/ML; 70 UNITS/ML N018195 001

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN BR

LILLY 100 UNITS/ML N019529 001 Apr 28, 1986

VELOSULIN BR

NOVO NORDISK INC 100 UNITS/ML N021028 001 Jul 19, 1999

POWDER; INHALATION

EXUBERA

PFIZER 1MG/INH N021868 001 Jan 27, 2006

3MG/INH N021868 002 Jan 27, 2006

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN 50/50

LILLY 50 UNITS/ML; 50 UNITS/ML N020100 001 Apr 29, 1992

INSULIN RECOMBINANT PURIFIED HUMAN

INJECTABLE; INJECTION

NOVOLIN R

NOVO NORDISK INC 100 UNITS/ML N018778 001 Aug 30, 1983

VELOSULIN BR HUMAN

NOVO NORDISK INC 100 UNITS/ML N019450 001 May 30, 1986

INSULIN RECOMBINANT PURIFIED HUMAN; INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

MIXTARD HUMAN 70/30

BAYER PHARMS 30 UNITS/ML; 70 UNITS/ML N019585 001 Mar 11, 1988

NOVOLIN 70/30

NOVO NORDISK INC 30 UNITS/ML; 70 UNITS/ML N019441 001 Jul 11, 1986

INSULIN SUSP ISOPHANE BEEF

INJECTABLE; INJECTION

NPH INSULIN

NOVO NORDISK INC 40 UNITS/ML N017929 001

100 UNITS/ML N017929 003

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION

NPH ILETIN I (BEEF-PORK)

LILLY 40 UNITS/ML N017936 001

100 UNITS/ML N017936 002

INSULIN SUSP ISOPHANE PURIFIED BEEF

INJECTABLE; INJECTION

NPH ILETIN II

LILLY 100 UNITS/ML N018479 001

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN INSULATARD NPH NORDISK

NOVO NORDISK INC 100 UNITS/ML N018194 001

NPH ILETIN II (PORK)

LILLY 100 UNITS/ML N018345 001

NPH PURIFIED PORK ISOPHANE INSULIN

NOVO NORDISK INC 100 UNITS/ML N018623 001

**DISCONTINUED DRUG PRODUCT LIST**

6-206(of 375)

\*\* See List Footnote

INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

INSULATARD NPH HUMAN

NOVO NORDISK INC 100 UNITS/ML

N019449 001 May 30, 1986

NOVOLIN N

NOVO NORDISK INC 100 UNITS/ML

N019065 001 Jan 23, 1985

INSULIN SUSP PROTAMINE ZINC BEEF/PORK

INJECTABLE; INJECTION

PROTAMINE ZINC &amp; ILETIN I (BEEF-PORK)

LILLY 40 UNITS/ML  
100 UNITS/ML

N017932 001

N017932 002

INSULIN SUSP PROTAMINE ZINC PURIFIED BEEF

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II

LILLY 100 UNITS/ML

N018476 001

PROTAMINE ZINC INSULIN

BRISTOL MYERS SQUIBB 40 UNITS/ML  
100 UNITS/ML

N017928 001

N017928 003

INSULIN SUSP PROTAMINE ZINC PURIFIED PORK

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II (PORK)

LILLY 100 UNITS/ML

N018346 001

INSULIN ZINC SUSP BEEF

INJECTABLE; INJECTION

LENTE INSULIN

NOVO NORDISK INC 40 UNITS/ML  
100 UNITS/ML

N017998 001

N017998 003

INSULIN ZINC SUSP EXTENDED BEEF

INJECTABLE; INJECTION

ULTRALENTE INSULIN

NOVO NORDISK INC 100 UNITS/ML

N017997 003

INSULIN ZINC SUSP EXTENDED PURIFIED BEEF

INJECTABLE; INJECTION

ULTRALENTE

NOVO NORDISK INC 100 UNITS/ML

N018385 001

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN U

LILLY 40 UNITS/ML  
100 UNITS/ML

N019571 001 Jun 10, 1987

N019571 002 Jun 10, 1987

INSULIN ZINC SUSP PROMPT BEEF

INJECTABLE; INJECTION

SEMILENTE INSULIN

NOVO NORDISK INC 100 UNITS/ML

N017996 003

INSULIN ZINC SUSP PROMPT PURIFIED PORK

INJECTABLE; INJECTION

SEMILENTE

NOVO NORDISK INC 100 UNITS/ML

N018382 001

INSULIN ZINC SUSP PURIFIED BEEF

INJECTABLE; INJECTION

LENTE ILETIN II

LILLY 100 UNITS/ML

N018477 001

INSULIN ZINC SUSP PURIFIED BEEF/PORK

INJECTABLE; INJECTION

LENTARD

NOVO NORDISK INC 100 UNITS/ML

N018384 001

INSULIN ZINC SUSP PURIFIED PORK

INJECTABLE; INJECTION

LENTE

NOVO NORDISK INC 100 UNITS/ML

N018383 001

LENTE ILETIN II (PORK)

LILLY 100 UNITS/ML

N018347 001

**DISCONTINUED DRUG PRODUCT LIST**

6-207(of 375)

\*\* See List Footnote

INSULIN ZINC SUSP RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN L

LILLY

100 UNITS/ML

N019377 002 Sep 30, 1985

NOVOLIN L

NOVO NORDISK INC

100 UNITS/ML

N019965 001 Jun 25, 1991

INSULIN ZINC SUSP SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION

NOVOLIN L

NOVO NORDISK INC

100 UNITS/ML

N018777 001 Aug 30, 1983

INULIN

INJECTABLE; INJECTION

INULIN AND SODIUM CHLORIDE

ISO TEX

100MG/ML

N002282 001

INVERT SUGAR

INJECTABLE; INJECTION

TRAVERT 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE

10GM/100ML

N016717 001

IOBENGUANE SULFATE I-131

INJECTABLE; INJECTION

IOBENGUANE SULFATE I 131

PHARMALUCENCE

2.3mCi/ML

N020084 001 Mar 25, 1994

IOCETAMIC ACID

TABLET; ORAL

CHOLEBRINE

MALLINCKRODT

750MG

N017129 001

IODAMIDE MEGLUMINE

INJECTABLE; INJECTION

RENOVUE-65

BRACCO

65%

N017902 001

RENOVUE-DIP

BRACCO

24%

N017903 001

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION

CHOLOGRAFIN MEGLUMINE

BRACCO

10.3%

N009321 007

+

52%

N009321 003

IODIPAMIDE SODIUM

INJECTABLE; INJECTION

CHOLOGRAFIN SODIUM

BRACCO

20%

N009321 001

IODIXANOL

INJECTABLE; INJECTION

VISIPAQUE 270

GE HEALTHCARE

55%

N020808 001 Aug 29, 1997

ODOHIPPURATE SODIUM I-123

INJECTABLE; INJECTION

NEPHROFLOW

GE HEALTHCARE

1mCi/ML

N018289 001 Dec 28, 1984

ODOHIPPURATE SODIUM I-131

INJECTABLE; INJECTION

HIPPURAN I 131

MALLINCKRODT

0.25mCi/ML

N016666 001

HIPPUTOPE

BRACCO

1-2mCi/VIAL

N015419 002

ODOHIPPURATE SODIUM I 131

PHARMALUCENCE

0.2mCi/ML

N017313 001

**DISCONTINUED DRUG PRODUCT LIST**

6-208(of 375)

\*\* See List Footnote

IODOXAMATE MEGLUMINEINJECTABLE; INJECTION  
CHOLOVUE

BRACCO	9.9%	N018077 001
	40.3%	N018076 001

IOFETAMINE HYDROCHLORIDE I-123INJECTABLE; INJECTION  
SPECTAMINE

IMP	1mCi/ML	N019432 001 Dec 24, 1987
-----	---------	--------------------------

IOHESOLINJECTABLE; INJECTION  
OMNIPAQUE 210

GE HEALTHCARE	45.3%	N018956 006 Jun 30, 1989
---------------	-------	--------------------------

SOLUTION; URETHRAL

OMNIPAQUE 70		N018956 007 Jun 01, 1994
--------------	--	--------------------------

GE HEALTHCARE	15.1%
---------------	-------

IOPAMIDOLINJECTABLE; INJECTION  
IOPAMIDOL

BAXTER HLTHCARE	41%	A074629 001 Nov 06, 1996
	51%	A074629 004 Mar 31, 1998
	61%	A074629 002 Nov 06, 1996
	76%	A074629 003 Nov 06, 1996
HOSPIRA	61%	A074734 001 Dec 10, 1996
	76%	A074734 002 Dec 10, 1996

IOPAMIDOL-200

COOK IMAGING	41%	A074881 001 Jul 28, 2000
HOSPIRA	41%	A074898 001 Dec 30, 1997

IOPAMIDOL-200 IN PLASTIC CONTAINER

HOSPIRA	41%	A074636 001 Dec 30, 1997
---------	-----	--------------------------

IOPAMIDOL-250

COOK IMAGING	51%	A074881 002 Jul 28, 2000
FRESENIUS KABI USA	51%	A074679 001 Apr 02, 1997
HOSPIRA	51%	A074898 002 Dec 30, 1997
	51%	A075005 001 Feb 24, 1998

IOPAMIDOL-250 IN PLASTIC CONTAINER

HOSPIRA	51%	A074636 002 Dec 30, 1997
---------	-----	--------------------------

IOPAMIDOL-300

ABBVIE	61%	A074638 001 Apr 30, 1997
COOK IMAGING	61%	A074881 003 Jul 28, 2000
FRESENIUS KABI USA	61%	A074679 002 Apr 02, 1997
HOSPIRA	61%	A074898 003 Dec 30, 1997
	61%	A075005 002 Feb 24, 1998

IOPAMIDOL-300 IN PLASTIC CONTAINER

HOSPIRA	61%	A074636 003 Dec 30, 1997
	61%	A074637 001 Apr 03, 1997

IOPAMIDOL-370

COOK IMAGING	76%	A074881 004 Jul 28, 2000
FRESENIUS KABI USA	76%	A074679 003 Apr 02, 1997
HOSPIRA	76%	A074898 004 Dec 30, 1997
	76%	A075005 003 Feb 24, 1998

IOPAMIDOL-370 IN PLASTIC CONTAINER

HOSPIRA	76%	A074636 004 Dec 30, 1997
---------	-----	--------------------------

ISOVUE-128

BRACCO	26%	N018735 005 Oct 21, 1986
--------	-----	--------------------------

ISOVUE-200

BRACCO	41%	N020327 001 Oct 12, 1994
--------	-----	--------------------------

IOPANOIC ACIDTABLET; ORAL  
TELEPAQUE

GE HEALTHCARE	500MG	N008032 001
---------------	-------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-209(of 375)

\*\* See List Footnote

**IOPHENIDYLATE**

INJECTABLE; INJECTION  
 PANTOPAQUE  
 ALCON 100% N005319 001

**IOPROMIDE**

INJECTABLE; INJECTION  
 ULTRAVIST 150  
 + BAYER HLTHCARE 31.2% N020220 004 May 10, 1995  
 ULTRAVIST 300 IN PLASTIC CONTAINER  
 + BAYER HLTHCARE 62.3% N020220 005 Nov 18, 2008

**IOTHALAMATE MEGLUMINE; IOTHALAMATE SODIUM**

INJECTABLE; INJECTION  
 VASCORAY  
 MALLINCKRODT 52%;26% N016783 001

**IOTHALAMATE SODIUM**

INJECTABLE; INJECTION  
 ANGIO-CONRAY  
 MALLINCKRODT 80% N013319 001  
 CONRAY 325  
 MALLINCKRODT 54.3% N017685 001  
 CONRAY 400  
 MALLINCKRODT 66.8% N014295 001

**IOTROLAN**

INJECTABLE; INTRATHECAL  
 OSMOVIST 190  
 BAYER HLTHCARE 40.6% N019580 001 Dec 07, 1989  
 OSMOVIST 240  
 BAYER HLTHCARE 51.3% N019580 002 Dec 07, 1989

**IOVERSOL**

INJECTABLE; INJECTION  
 OPTIRAY 160  
 LIEBEL-FLARSHEIM 34% N019710 003 Dec 30, 1988  
 OPTIRAY 240  
 LIEBEL-FLARSHEIM 51% N020923 001 May 28, 1998

**IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM**

INJECTABLE; INJECTION  
 HEXABRIX  
 GUERBET 39.3%;19.6% N018905 002 Jul 26, 1985

**OXILAN**

INJECTABLE; INJECTION  
 OXILAN-300  
 GUERBET 62% N020316 001 Dec 21, 1995  
 OXILAN-350  
 GUERBET 73% N020316 002 Dec 21, 1995

**IPODATE CALCIUM**

GRANULE; ORAL  
 ORAGRAFIN CALCIUM  
 BRACCO 3GM/PACKET N012968 001

**IPODATE SODIUM**

CAPSULE; ORAL  
 BILIVIST  
 BAYER HLTHCARE 500MG A087768 001 Aug 11, 1982  
 ORAGRAFIN SODIUM  
 BRACCO 500MG N012967 001

**IPRATROPIUM BROMIDE**

AEROSOL, METERED; INHALATION  
 ATROVENT  
 BOEHRINGER INGELHEIM 0.018MG/INH N019085 001 Dec 29, 1986  
 SOLUTION; INHALATION  
 ATROVENT  
 + BOEHRINGER INGELHEIM 0.02% \*\* N020228 001 Sep 29, 1993  
 IPRATROPIUM BROMIDE  
 ACTAVIS MID ATLANTIC 0.02% A075111 001 Apr 22, 1999  
 APOTEX INC 0.02% A075441 001 Mar 28, 2001

**DISCONTINUED DRUG PRODUCT LIST**

6-210(of 375)

\*\* See List Footnote

IPRATROPIUM BROMIDE

SOLUTION; INHALATION

IPRATROPIUM BROMIDE

BAUSCH AND LOMB INC	0.02%	A075835 001 Oct 15, 2001
MYLAN SPECIALITY LP	0.02%	A074755 001 Jan 10, 1997
PHARMASCIENCE INC	0.02%	A075507 001 Jan 19, 2001
ROXANE	0.02%	A075867 001 Jul 22, 2002
TEVA PHARMS USA	0.02%	A075313 001 Feb 07, 2000
SPRAY, METERED; NASAL		
IPRATROPIUM BROMIDE		
APOTEX INC	0.021MG/SPRAY	A076156 001 Apr 18, 2003
	0.042MG/SPRAY	A076155 001 Apr 18, 2003

IRBESARTAN

TABLET; ORAL

IRBESARTAN

AJANTA PHARMA LTD	75MG	A203685 001 Dec 10, 2015
	150MG	A203685 002 Dec 10, 2015
	300MG	A203685 003 Dec 10, 2015
MYLAN PHARMS INC	75MG	A200461 001 Sep 27, 2012
	150MG	A200461 002 Sep 27, 2012
	300MG	A200461 003 Sep 27, 2012
WATSON LABS INC	75MG	A090720 001 Oct 12, 2012
	150MG	A090720 002 Oct 12, 2012
	300MG	A090720 003 Oct 12, 2012

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

IRINOTECAN HYDROCHLORIDE

SANDOZ	40MG/2ML (20MG/ML)	A077994 001 Feb 27, 2008
	100MG/5ML (20MG/ML)	A077994 002 Feb 27, 2008
SANDOZ INC	40MG/2ML (20MG/ML)	A090137 001 Nov 12, 2009
	100MG/5ML (20MG/ML)	A090137 002 Nov 12, 2009

IRON DEXTRAN

INJECTABLE; INJECTION

IRON DEXTRAN

SANOFI AVENTIS US	EQ 50MG IRON/ML	N010787 002
-------------------	-----------------	-------------

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

LUITPOLD	EQ 65MG BASE/3.25ML (EQ 20MG BASE/ML)	N021135 005 Mar 29, 2013
	EQ 75MG BASE/3.75ML (EQ 20MG BASE/ML)	N021135 003 Mar 29, 2005

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

BETA-2

NEPHRON	1%	A086711 001
BRONKOSOL		
SANOFI AVENTIS US	0.25%	N012339 009
	1%	N012339 008
ISOETHARINE HYDROCHLORIDE		
ALPHARMA US PHARMS	1%	A087101 001
ASTRAZENECA	0.062%	A087937 001 Nov 15, 1982
	0.062%	A089614 001 Jun 13, 1991
	0.125%	A087938 001 Nov 15, 1982
	0.125%	A089615 001 Jun 13, 1991
	0.167%	A088470 001 Mar 14, 1984
	0.167%	A089616 001 Jun 13, 1991
	0.2%	A088471 001 Mar 14, 1984
	0.2%	A089617 001 Jun 13, 1991
	0.25%	A088472 001 Mar 14, 1984
	0.25%	A089618 001 Jun 13, 1991
BAXTER HLTHCARE	0.08%	A088144 001 Jul 29, 1983
	0.14%	A088145 001 Mar 26, 1984
	0.25%	A088146 001 Aug 01, 1983
DEY	0.08%	A088187 001 Dec 03, 1982
	0.1%	A087389 001
	0.17%	A087390 001
	0.25%	A088188 001 Dec 03, 1982
	1%	A086763 001

**DISCONTINUED DRUG PRODUCT LIST**

6-211(of 375)

\*\* See List Footnote

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION

ISOETHARINE HYDROCHLORIDE

INTL MEDICATION	0.077%	A086651 001
	0.08%	A086651 002
	0.1%	A086651 003
	0.143%	A086651 004
	0.167%	A086651 005
	0.2%	A086651 006
	0.25%	A086651 007
	1%	A086651 008
PARKE DAVIS	0.5%	A085997 001
	1%	A085889 001
ROXANE	0.1%	A087396 001
	0.125%	A087025 001
	0.167%	A088226 001 Sep 16, 1983
	0.2%	A087324 001
	0.25%	A088275 001 Jun 03, 1983
	1%	A086899 001
ISOETHARINE HYDROCHLORIDE S/F		
DEY	0.08%	A089817 001 Nov 22, 1988
	0.1%	A089818 001 Nov 22, 1988
	0.17%	A089819 001 Nov 22, 1988
	0.25%	A089820 001 Nov 22, 1988
	1%	A089252 001 Sep 15, 1986

ISOETHARINE MESYLATE

AEROSOL, METERED; INHALATION

BRONKOMETER

SANOFI AVENTIS US	0.34MG/INH	N012339 007
ISOETHARINE MESYLATE ALPHARMA US PHARMS	0.34MG/INH	A087858 001 Aug 21, 1984

ISOFLURANE

LIQUID; INHALATION

ISOFLURANE

WATSON LABS INC	99.9%	A074393 001 May 12, 1995
-----------------	-------	--------------------------

ISOFLUROPHATE

OINTMENT; OPHTHALMIC

FLOROPRYL

MERCK	0.025%	N010656 001
-------	--------	-------------

ISONIAZID

INJECTABLE; INJECTION

NYDRAZID

SANDOZ	100MG/ML **	N008662 001
RIMIFON		
ROCHE	25MG/ML	N008420 002

SYRUP; ORAL

ISONIAZID

MIKART	50MG/5ML	A081118 001 Jul 21, 1997
LANIAZID		
LANNETT	50MG/5ML	A089243 001 Feb 03, 1986

RIMIFON

ROCHE	50MG/5ML	N008420 001
-------	----------	-------------

TABLET; ORAL

DOW-ISONIAZID

DOW PHARM	300MG	A080330 002
HYZYD		

MEDPOINTE PHARM HLC	100MG	A080134 003
	300MG	A080134 004

INH

NOVARTIS	300MG	A080935 001
----------	-------	-------------

ISONIAZID		
DURAMED PHARMS BARR	100MG	A088231 001 Mar 17, 1983
	300MG	A088119 001 Mar 17, 1983

HALSEY	50MG	A083632 001
HIKMA INTL PHARMS	100MG	A080212 001

	300MG	A087425 001
--	-------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-212(of 375)

\*\* See List Footnote

ISONIAZID

TABLET;ORAL

ISONIAZID

IMPAX LABS	100MG	A080153 001
IVAX SUB TEVA PHARMS	100MG	A080270 001
	300MG	A083610 001
LILLY	100MG	N008499 002
	300MG	N008499 003
MK LABS	100MG	A080941 001
NEXGEN PHARMA INC	100MG	A084050 001
PANRAY	50MG	N008428 001
	100MG	N008428 002
	300MG	N008428 003
PERRIGO	100MG	A083060 001
PHARMAVITE	100MG	A085091 001
PHOENIX LABS NY	50MG	A080368 001
	100MG	A080368 002
PUREPAC PHARM	50MG	A080132 003 Jul 14, 1982
	100MG	A080132 004 Jul 14, 1982
SUN PHARM INDUSTRIES	100MG	A080136 001
	300MG	A083633 001
WATSON LABS	50MG	A080522 001
	100MG	A080401 001
	100MG	A080523 001
	100MG	A085790 001
	300MG	A080521 001
	300MG	A083178 001
	300MG	A085784 001
WHITEWORTH TOWN PLSN	100MG	A080120 002
LANIAZID		
LANNETT	50MG	A080140 001
	100MG	A080140 002
NYDRAZID		
BRISTOL MYERS SQUIBB	100MG	N008392 003
STANOZIDE		
EVERYLIFE	100MG	A080126 001
	300MG	A080126 002

ISONIAZID; RIFAMPIN

CAPSULE;ORAL

RIFAMPIN AND ISONIAZID

HIKMA INTL PHARMS 150MG;300MG

A065221 001 Jul 29, 2005

ISOPROPAMIDE IODIDE

TABLET;ORAL

DARBID

GLAXOSMITHKLINE EQ 5MG BASE

N010744 001

ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED;INHALATION

ISOPROTERENOL HYDROCHLORIDE

3M 0.12MG/INH

N010375 004

ALPHARMA US PHARMS 0.12MG/INH

A085904 001

ISUPREL

SANOFI AVENTIS US 0.103MG/INH

N011178 001

DISC;INHALATION

NORISODRINE AEROTROL

ABBOTT 0.25%

N016814 001

INJECTABLE;INJECTION

ISOPROTERENOL HYDROCHLORIDE

ABRAXIS PHARM 0.2MG/ML

A083431 001

BAXTER HLTHCARE 0.2MG/ML

A083486 001

HOSPIRA 0.02MG/ML

A083283 001

0.2MG/ML

A083346 001

INTL MEDICATION 0.2MG/ML

A083724 001

SOLUTION;INHALATION

AEROLONE

LILLY 0.25%

N007245 001

ISOPROTERENOL HYDROCHLORIDE

ARMOUR PHARM 0.031%

A087935 001 Nov 18, 1982

0.062%

A087936 001 Nov 18, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-213(of 375)

\*\* See List Footnote

ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

ISOPROTERENOL HYDROCHLORIDE

DEY	0.5%	A086764 001 Jan 04, 1982
PARKE DAVIS	0.25%	A085994 001
	0.5%	A085540 001
ISUPREL		
SANOFI AVENTIS US	0.5%	N006327 002
	1%	N006327 003
VAPO-ISO		
FISONS	0.5%	N016813 001
TABLET; RECTAL, SUBLINGUAL		
ISUPREL		
SANOFI AVENTIS US	10MG	N006328 001
	15MG	N006328 002

ISOPROTERENOL HYDROCHLORIDE; PHENYLEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

DUO-MEDIHALER

3M	0.16MG/INH; 0.24MG/INH	N013296 001
----	------------------------	-------------

ISOPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

MEDIHALER-ISO

3M	0.08MG/INH	N010375 003
----	------------	-------------

POWDER; INHALATION

NORISODRINE

ABBVIE	10%	N006905 003
	25%	N006905 002

ISOSORBIDE

SOLUTION; ORAL

ISMOTIC

ALCON	100GM/220ML	N017063 001
-------	-------------	-------------

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

ISORDIL

WYETH AYERST	40MG	N012882 002 Jul 29, 1988
TABLET; ORAL		
ISORDIL		
+ VALEANT PHARMS NORTH	10MG **	N012093 002 Jul 29, 1988

+	20MG **	N012093 006 Jul 29, 1988
+	30MG **	N012093 005 Jul 29, 1988

ISOSORBIDE DINITRATE

SUN PHARM INDUSTRIES	5MG	A086166 002 Sep 19, 1986
	10MG	A086169 001 Sep 19, 1986
	20MG	A086167 001 Sep 19, 1986
	30MG	A087564 001 Sep 18, 1986
SUPERPHARM	5MG	A089190 001 Feb 17, 1987
	10MG	A089191 001 Feb 17, 1987
	20MG	A089192 001 Feb 17, 1987
WATSON LABS	5MG	A086034 001 Jan 06, 1988
	10MG	A086032 001 Jan 07, 1988

SORBITRATE

ASTRAZENECA	5MG	N016192 001 Apr 01, 1996
	10MG	N016192 002 Apr 01, 1996
	20MG	A086405 002 Aug 21, 1990
	30MG	A088124 001 Aug 21, 1990
	40MG	A088125 001 Aug 21, 1990

TABLET; SUBLINGUAL

ISORDIL

+ BIOVAIL	2.5MG **	N012940 004 Jul 29, 1988
+	5MG **	N012940 003 Jul 29, 1988
+	10MG **	N012940 005 Jul 29, 1988

ISOSORBIDE DINITRATE

HIKMA INTL PHARMS	2.5MG	A086054 001 Oct 29, 1987
	5MG	A086055 001 Nov 02, 1987
SANDOZ	2.5MG	A086225 001 Feb 19, 1988
	5MG	A086222 001 Feb 19, 1988
SUN PHARM INDUSTRIES	2.5MG	A084204 001 Sep 18, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-214(of 375)

\*\* See List Footnote

ISOSORBIDE DINITRATE

TABLET; SUBLINGUAL

ISOSORBIDE DINITRATE

	5MG	A086168 001	Sep 18, 1986
	10MG	A087545 001	Sep 18, 1986
WATSON LABS	2.5MG **	A086033 001	Feb 26, 1988
WATSON LABS TEVA	5MG **	A086031 001	Sep 29, 1987
SORBITRATE			
ASTRAZENECA	2.5MG	N016191 002	Apr 01, 1996
	5MG	N016191 001	Apr 01, 1996
TABLET, CHEWABLE; ORAL			
SORBITRATE			
ASTRAZENECA	5MG	N016776 002	Apr 01, 1996
	10MG	N016776 003	Apr 01, 1996
TABLET, EXTENDED RELEASE; ORAL			
ISORDIL			
WYETH AYERST	40MG	N012882 001	Jul 29, 1988
ISOSORBIDE DINITRATE			
IMPAKX LABS INC	40MG	A040723 001	Mar 17, 2008

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISMO

PROMIUS PHARMA	20MG	N019091 001	Dec 30, 1991
TABLET, EXTENDED RELEASE; ORAL			
IMDUR			
+ SCHERING PLOUGH	30MG **	N020225 001	Aug 12, 1993
+	60MG **	N020225 002	Aug 12, 1993
+	120MG **	N020225 003	Mar 30, 1995
ISOSORBIDE MONONITRATE			
ACTAVIS ELIZABETH	30MG	A075306 001	Dec 31, 1998
	60MG	A075306 002	Dec 31, 1998
ALKERMES GAINESVILLE	60MG	A075041 001	Sep 22, 1998
IVAX SUB TEVA PHARMS	30MG	A075448 002	Aug 07, 2001
	60MG	A075448 001	Jun 19, 2000
	120MG	A075448 003	Aug 07, 2001
SKYEPHARMA AG	60MG	A075166 001	Oct 07, 1999

ISOSULFAN BLUE

INJECTABLE; INJECTION

LYMPHAZURIN

+ COVIDIEN

1% \*\*

N018310 001

ISOTRETINOIN

CAPSULE; ORAL

ACCUTANE

+ HOFFMANN LA ROCHE	10MG **	N018662 002	May 07, 1982
+	20MG **	N018662 004	Mar 28, 1983
+	40MG **	N018662 003	May 07, 1982

SOTRET

SUN PHARM INDs LTD	10MG	A076041 001	Dec 24, 2002
	20MG	A076041 002	Dec 24, 2002
	30MG	A076503 001	Jun 20, 2003
	40MG	A076041 003	Dec 24, 2002

ISRADIPINE

CAPSULE; ORAL

DYNACIRC

+ SMITHKLINE BEECHAM	2.5MG	N019546 001	Dec 20, 1990
+	5MG	N019546 002	Dec 20, 1990

TABLET, EXTENDED RELEASE; ORAL

DYNACIRC CR			
+ GLAXOSMITHKLINE LLC	5MG **	N020336 001	Jun 01, 1994
+	10MG **	N020336 002	Jun 01, 1994

ISRADIPINE

MYLAN PHARMS INC	5MG	A201067 001	Nov 27, 2015
	10MG	A201067 002	Nov 27, 2015

**DISCONTINUED DRUG PRODUCT LIST**

6-215(of 375)

\*\* See List Footnote

**ITRACONAZOLE**

INJECTABLE; INJECTION  
 SPORANOX  
 JANSSEN PHARMS 10MG/ML N020966 001 Mar 30, 1999  
 SOLUTION; ORAL  
 ITRACONAZOLE  
 AMNEAL PHARMS 10MG/ML A205573 001 Oct 30, 2015

**IVERMECTIN**

TABLET; ORAL  
 STROMECTOL  
 MERCK SHARP DOHME 6MG N050742 001 Nov 22, 1996

**KANAMYCIN SULFATE**

CAPSULE; ORAL  
 KANTREX  
 APOTHECON EQ 500MG BASE A060516 001  
 EQ 500MG BASE A061911 001  
 EQ 500MG BASE A062726 001 Mar 06, 1987  
 INJECTABLE; INJECTION  
 KANAMYCIN  
 WEST-WARD PHARMS INT EQ 75MG BASE/2ML A062324 001  
 EQ 500MG BASE/2ML A062324 002  
 EQ 1GM BASE/3ML A062324 003  
 KANAMYCIN SULFATE  
 ABRAXIS PHARM EQ 75MG BASE/2ML A062504 001 Apr 05, 1984  
 EQ 500MG BASE/2ML A062504 002 Apr 05, 1984  
 EQ 1GM BASE/3ML A062504 003 Apr 05, 1984  
 INTL MEDICATION EQ 500MG BASE/2ML A062466 001 Sep 30, 1983  
 EQ 1GM BASE/3ML A062466 002 Sep 30, 1983  
 LOCH EQ 75MG BASE/2ML A063021 001 Jul 31, 1992  
 EQ 500MG BASE/2ML A063022 001 Jul 31, 1992  
 EQ 1GM BASE/3ML A063025 001 Jul 31, 1992  
 PHARMAFAIR EQ 75MG BASE/2ML A062668 001 May 07, 1987  
 EQ 500MG BASE/2ML A062672 001 May 07, 1987  
 EQ 1GM BASE/3ML A062669 001 May 07, 1987  
 SOLOPAK EQ 75MG BASE/2ML A062605 003 Feb 26, 1986  
 EQ 500MG BASE/2ML A062605 001 Feb 26, 1986  
 EQ 1GM BASE/3ML A062605 002 Feb 26, 1986  
 WARNER CHILCOTT EQ 1GM BASE/3ML A063092 001 Oct 11, 1989  
 WATSON LABS EQ 1GM BASE/3ML A062520 003 May 09, 1985  
 KANTREX  
 APOTHECON EQ 75MG BASE/2ML A061655 003  
 EQ 75MG BASE/2ML A061901 003  
 EQ 75MG BASE/2ML A062564 001 Sep 21, 1984  
 EQ 500MG BASE/2ML A061655 001  
 EQ 500MG BASE/2ML A061901 001  
 EQ 500MG BASE/2ML A062564 002 Sep 21, 1984  
 EQ 1GM BASE/3ML A061655 002  
 EQ 1GM BASE/3ML A061901 002  
 EQ 1GM BASE/3ML A062564 003 Sep 21, 1984  
 KLEBCIL  
 KING PHARMS EQ 75MG BASE/2ML A062170 001  
 EQ 500MG BASE/2ML A062170 002  
 EQ 1GM BASE/3ML A062170 003

**KETOCONAZOLE**

CREAM; TOPICAL  
 NIZORAL  
 + JANSSEN PHARMA 2% N019084 001 Dec 31, 1985  
 SUSPENSION; ORAL  
 NIZORAL  
 JANSSEN PHARMA 100MG/5ML A070767 001 Nov 07, 1986  
 TABLET; ORAL  
 KETOCONAZOLE  
 AAIPHARMA LLC 200MG A075341 001 Jul 27, 1999  
 APOTEX 200MG A075912 001 Jan 10, 2002  
 PLIVA 200MG A075362 001 Jun 15, 1999  
 SUN PHARM INDUSTRIES 200MG A075314 001 Jun 15, 1999  
 TEVA 200MG A074971 001 Jun 15, 1999

**DISCONTINUED DRUG PRODUCT LIST**

6-216(of 375)

\*\* See List Footnote

KETOCONAZOLE

TABLET;ORAL

NIZORAL

+ JANSSEN PHARMS

200MG \*\*

N018533 001

KETOPROFEN

CAPSULE;ORAL

KETOPROFEN

AUROLIFE PHARMA LLC 50MG

A074024 001 Dec 29, 1995

75MG

A074024 002 Dec 29, 1995

TEVA 25MG

A073515 001 Dec 22, 1992

ORUDIS

+ WYETH AYERST 25MG \*\*

N018754 001 Jul 31, 1987

+ 50MG \*\*

N018754 002 Jan 09, 1986

+ 75MG \*\*

N018754 003 Jan 09, 1986

CAPSULE, EXTENDED RELEASE;ORAL

KETOPROFEN

ALKERMES GAINESVILLE 200MG

A074879 001 Dec 10, 1997

ORUVAIL

+ WYETH PHARMS INC 100MG \*\*

N019816 003 Feb 08, 1995

+ 150MG \*\*

N019816 002 Feb 08, 1995

+ 200MG \*\*

N019816 001 Sep 24, 1993

FILM;ORAL

NEXCDE

NOVARTIS 12.5MG

N022470 001 Nov 25, 2009

TABLET;ORAL

ACTRON

BAYER 12.5MG

N020499 001 Oct 06, 1995

KETOPROFEN

PERRIGO 12.5MG

A075364 001 Feb 07, 2002

ORUDIS KT

+ WYETH CONS 12.5MG \*\*

N020429 001 Oct 06, 1995

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

APOTEX INC 30MG/ML

A075626 001 Jul 24, 2001

30MG/ML

A077201 001 Oct 14, 2005

APOTHECON 15MG/ML

A075348 001 Nov 28, 2000

30MG/ML

A075348 002 Nov 28, 2000

BAXTER HLTHCARE CORP 15MG/ML

A075631 002 Jun 29, 2001

30MG/ML

A075631 001 Jun 29, 2001

BEDFORD 15MG/ML

A075230 002 Oct 25, 1999

30MG/ML

A075230 001 Oct 25, 1999

GLAND PHARMA LTD 15MG/ML

A076722 001 Jul 27, 2004

30MG/ML

A076722 002 Jul 27, 2004

HOSPIRA 15MG/ML

A074801 001 Jun 05, 1997

30MG/ML

A074801 002 Jun 05, 1997

LUITPOLD 15MG/ML

A078145 001 Jan 14, 2008

30MG/ML

A078145 002 Jan 14, 2008

MYLAN LABS LTD 15MG/ML

A078299 001 Jul 16, 2007

15MG/ML

A201155 001 Aug 04, 2014

30MG/ML

A078299 002 Jul 16, 2007

SANDOZ INC 15MG/ML

A201155 002 Aug 04, 2014

SUN PHARMA GLOBAL 15MG/ML

A076271 001 Oct 06, 2004

30MG/ML

A078737 001 Oct 06, 2008

WEST-WARD PHARMS INT 15MG/ML \*\*

A078737 002 Oct 06, 2008

15MG/ML

A075222 001 Apr 26, 1999

15MG/ML

A075299 001 Nov 03, 1999

30MG/ML \*\*

A075772 001 Jul 21, 2004

30MG/ML \*\*

A075222 002 Apr 26, 1999

30MG/ML

A075228 001 Apr 26, 1999

30MG/ML

A075299 002 Nov 03, 1999

30MG/ML

A075772 002 Jul 21, 2004

WOCKHARDT 30MG/ML

A077943 001 Mar 27, 2007

TORADOL

+ ROCHE PALO 15MG/ML \*\*

N019698 001 Nov 30, 1989

+ 30MG/ML \*\*

N019698 002 Nov 30, 1989

**DISCONTINUED DRUG PRODUCT LIST**

6-217(of 375)

\*\* See List Footnote

KETOROLAC TROMETHAMINE

SOLUTION/DROPS;OPHTHALMIC ACULAR PRESERVATIVE FREE ALLERGAN	0.5%	N020811 001 Nov 03, 1997
KETOROLAC TROMETHAMINE AKORN	0.45%	A203376 001 Feb 10, 2014
TABLET;ORAL KETOROLAC TROMETHAMINE CYCLE PHARMS LTD	10MG	A074790 001 Jun 26, 1997
WATSON LABS	10MG	A074955 001 Sep 19, 1997
TORADOL + ROCHE PALO	10MG **	N019645 001 Dec 20, 1991

KETOTIFEN FUMARATE

SOLUTION/DROPS;OPHTHALMIC KETOTIFEN FUMARATE APOTEX INC	EQ 0.025% BASE	A077354 001 May 09, 2006
ZADITOR + ALCON PHARMA	EQ 0.025% BASE **	N021066 002 Oct 19, 2006

KRYPTON, KR-81M

GAS;INHALATION MPI KRYPTON 81M GENERATOR GE HEALTHCARE	N/A	N018088 001
--	-----	-------------

LABETALOL HYDROCHLORIDE

INJECTABLE;INJECTION LABETALOL HYDROCHLORIDE APOTHECON	5MG/ML	A075355 001 Nov 29, 1999
BAXTER HLTHCARE CORP	5MG/ML	A076051 001 Jul 05, 2002
HOSPIRA	5MG/ML	A075242 001 Sep 30, 1999
NORMODYNE SCHERING	5MG/ML	N018686 001 Aug 01, 1984
TRANDATE + SEBELA IRELAND LTD	5MG/ML **	N019425 001 Dec 31, 1985
TABLET;ORAL LABETALOL HYDROCHLORIDE APOTHECON	100MG	A075223 001 Nov 20, 1998
	200MG	A075223 002 Nov 20, 1998
	300MG	A075223 003 Nov 20, 1998
TEVA	100MG	A074989 001 Sep 30, 1998
	200MG	A074989 002 Sep 30, 1998
	300MG	A074989 003 Sep 30, 1998
NORMODYNE + SCHERING	100MG **	N018687 001 Aug 31, 1987
+	200MG **	N018687 002 Aug 01, 1984
+	300MG **	N018687 003 Aug 01, 1984
+	400MG **	N018687 004 Aug 01, 1984
TRANDATE + CNTY LINE PHARMS	400MG **	N018716 004 Aug 01, 1984

LACTULOSE

SOLUTION;ORAL CHRONULAC + SANOFI AVENTIS US	10GM/15ML **	N017884 001
CONSTULOSE ACTAVIS MID ATLANTIC	10GM/15ML	A070288 001 Aug 15, 1988
DUPHALAC SOLVAY	10GM/15ML	A072372 001 Mar 22, 1989
EVALOSE TEVA PHARMS	10GM/15ML	A073497 001 May 28, 1993
LACTULOSE APOTEX INC	10GM/15ML	A075911 001 Feb 21, 2002
MORTON GROVE	10GM/15ML	A071841 001 Sep 22, 1988
PACO	10GM/15ML	A073160 001 Aug 25, 1992
LAXILOSE NOSTRUM LABS	10GM/15ML	A073686 001 May 28, 1993
SOLUTION;ORAL, RECTAL ACILAC NOSTRUM LABS	10GM/15ML	A073685 001 May 28, 1993

**DISCONTINUED DRUG PRODUCT LIST**

6-218(of 375)

\*\* See List Footnote

LACTULOSE

SOLUTION;ORAL, RECTAL

CEPHULAC

+ SANOFI AVENTIS US 10GM/15ML \*\*

N017657 001

GENERLAC

MORTON GROVE 10GM/15ML

A071842 001 Sep 27, 1988

HEPTALAC

TEVA PHARMS 10GM/15ML

A073504 001 May 28, 1993

LACTULOSE

APOTEX INC 10GM/15ML

A076645 001 Jul 28, 2003

PACO 10GM/15ML

A072029 001 Aug 25, 1992

ROXANE 10GM/15ML

A073590 001 May 29, 1992

SOLVAY 10GM/15ML

N017906 001

PORTALAC

SOLVAY 10GM/15ML

A072374 001 Mar 22, 1989

LAMIVUDINE; Raltegravir POTASSIUM

TABLET;ORAL

DUTREBIS

MERCK SHARP DOHME 150MG;EQ 300MG BASE

N206510 001 Feb 06, 2015

LAMOTRIGINE

TABLET;ORAL

LAMICTAL

+ GLAXOSMITHKLINE LLC 50MG \*\*

N020241 006 Dec 27, 1994

+ 250MG \*\*

N020241 004 Dec 27, 1994

LAMOTRIGINE

ACTAVIS TOTOWA 25MG

A078669 001 Apr 08, 2011

100MG

A078669 002 Apr 08, 2011

150MG

A078669 003 Apr 08, 2011

200MG

A078669 004 Apr 08, 2011

HIKMA PHARMS 25MG

A078134 001 Apr 19, 2011

100MG

A078134 002 Apr 19, 2011

150MG

A078134 003 Apr 19, 2011

200MG

A078134 004 Apr 19, 2011

MYLAN 25MG

A077428 001 Jan 27, 2009

100MG

A077428 002 Jan 27, 2009

150MG

A077428 003 Jan 27, 2009

200MG

A077428 004 Jan 27, 2009

MYLAN LABS LTD 25MG

A078443 001 Feb 11, 2009

100MG

A078443 002 Feb 11, 2009

150MG

A078443 003 Feb 11, 2009

200MG

A078443 004 Feb 11, 2009

PHARMASCIENCE INC 25MG

A078310 001 Feb 04, 2009

100MG

A078310 002 Feb 04, 2009

150MG

A078310 003 Feb 04, 2009

200MG

A078310 004 Feb 04, 2009

ROXANE 25MG

A077392 001 Jan 27, 2009

100MG

A077392 002 Jan 27, 2009

150MG

A077392 003 Jan 27, 2009

200MG

A077392 004 Jan 27, 2009

SANDOZ 25MG

A078645 001 Jan 27, 2009

100MG

A078645 002 Jan 27, 2009

150MG

A078645 003 Jan 27, 2009

200MG

A078645 004 Jan 27, 2009

WOCKHARDT 25MG

A078982 001 Jan 27, 2009

100MG

A078982 002 Jan 27, 2009

150MG

A078982 003 Jan 27, 2009

200MG

A078982 004 Jan 27, 2009

TABLET, CHEWABLE;ORAL

LAMICTAL CD

GLAXOSMITHKLINE LLC 100MG

N020764 003 Aug 24, 1998

LAMOTRIGINE

MYLAN 5MG

A076630 001 Jan 22, 2009

25MG

A076630 002 Jan 22, 2009

SANDOZ 5MG

A078409 002 Jan 22, 2009

25MG

A078409 003 Jan 22, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-219(of 375)

\*\* See List Footnote

**LANSOPRAZOLE**

FOR SUSPENSION, DELAYED RELEASE;ORAL  
PREVACID

TAKEDA PHARMS NA      15MG/PACKET  
                          30MG/PACKET

N021281 001 May 03, 2001  
N021281 002 May 03, 2001

INJECTABLE;INTRAVENOUS

PREVACID IV

+ TAKEDA PHARMS NA      30MG/VIAL \*\*

N021566 001 May 27, 2004

TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE;ORAL

LANSOPRAZOLE

ANI PHARMS INC      15MG  
                          30MG

A078730 001 Oct 15, 2010  
A078730 002 Oct 15, 2010

**LANSOPRAZOLE; NAPROXEN**

CAPSULE, DELAYED REL PELLETS, TABLET;ORAL

PREVACID NAPRAPAC 250 (COPACKAGED)

+ TAKEDA PHARMS NA      15MG,N/A;N/A,250MG \*\*

N021507 002 Nov 14, 2003

PREVACID NAPRAPAC 375 (COPACKAGED)

TAKEDA PHARMS NA      15MG,N/A;N/A,375MG

N021507 003 Nov 14, 2003

PREVACID NAPRAPAC 500 (COPACKAGED)

TAKEDA PHARMS NA      15MG,N/A;N/A,500MG

N021507 004 Nov 14, 2003

**LANTHANUM CARBONATE**

TABLET, CHEWABLE;ORAL

FOSRENOL

SHIRE LLC      EQ 250MG BASE

N021468 001 Oct 26, 2004

**LAPYRUM CHLORIDE; UNDECYLUM CHLORIDE IODINE COMPLEX**

SOLUTION;TOPICAL

VIRAC REX

CHESEBROUGH PONDS      0.5%;1.8%

N011914 001

**LATANOPROST**

SOLUTION/DROPS;OPHTHALMIC

LATANOPROST

APOTEX INC      0.005%

A077697 001 Mar 22, 2011

**LEFLUNOMIDE**

TABLET;ORAL

LEFLUNOMIDE

SANDOZ      10MG  
                          10MG  
                          20MG  
                          20MG

A077085 001 Sep 13, 2005  
A077087 001 Sep 13, 2005  
A077085 002 Sep 13, 2005  
A077087 002 Sep 13, 2005

**LEPIRUDIN RECOMBINANT**

INJECTABLE;INJECTION

REFLUDAN

BAYER HLTHCARE      50MG/VIAL

N020807 001 Mar 06, 1998

**LETROZOLE**

TABLET;ORAL

LETROZOLE

ACTAVIS TOTOWA      2.5MG  
IMPAX LABS      2.5MG  
KREMERS URBAN PHARMS      2.5MG  
LANNETT HOLDINGS INC      2.5MG  
SUN PHARM IND LTD      2.5MG  
SYNTTHON PHARMS      2.5MG

A090292 001 Jul 13, 2011  
A091638 001 Jun 03, 2011  
A091098 001 Jun 03, 2011  
A202048 001 Oct 29, 2014  
A091466 001 Jun 03, 2011  
A090196 001 Jun 03, 2011

**LEUCOVORIN CALCIUM**

FOR SOLUTION;ORAL

LEUCOVORIN CALCIUM

HOSPIRA      EQ 60MG BASE/VIAL

N008107 003 Jan 30, 1987

INJECTABLE;INJECTION

LEUCOVORIN CALCIUM

ABIC      EQ 3MG BASE/ML  
                          EQ 50MG BASE/VIAL  
ABRAXIS PHARM      EQ 50MG BASE/VIAL  
ELKINS SINK      EQ 50MG BASE/VIAL  
EQ 100MG BASE/VIAL  
+ HOSPIRA      EQ 3MG BASE/ML \*\*  
+                            EQ 50MG BASE/VIAL \*\*

A089352 001 Jun 01, 1988  
A089353 001 Jun 01, 1988  
A088939 001 Dec 01, 1986  
A070480 001 Jan 02, 1987  
A081224 001 Jun 03, 1994  
N008107 001  
N008107 002

**DISCONTINUED DRUG PRODUCT LIST**

6-220(of 375)

\*\* See List Footnote

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

+	EQ 100MG BASE/VIAL **	N008107 004	May 23, 1988
+	EQ 350MG BASE/VIAL **	N008107 005	Apr 05, 1989
PHARMACHEMIE	EQ 350MG BASE/VIAL	A040262 001	Dec 15, 1999
PHARMACHEMIE USA	EQ 50MG BASE/VIAL	A089628 001	Apr 17, 1997
	EQ 100MG BASE/VIAL	A089915 001	Apr 17, 1997
TEVA PARENTERAL	EQ 50MG BASE/VIAL	A081278 001	Sep 28, 1993
LEUCOVORIN CALCIUM PRESERVATIVE FREE			
HOSPIRA	EQ 10MG BASE/ML **	A040147 001	Jun 25, 1997
LUITPOLD	EQ 50MG BASE/VIAL	A040338 001	Jan 31, 2001
TEVA PARENTERAL	EQ 10MG BASE/ML	A040332 001	Jun 28, 1999
WELLCOVORIN			
GLAXOSMITHKLINE	EQ 5MG BASE/ML	A087439 001	Oct 19, 1982
	EQ 25MG BASE/VIAL	A089833 001	Jan 23, 1989
	EQ 50MG BASE/VIAL	A089465 001	Jan 23, 1989
	EQ 100MG BASE/VIAL	A089834 001	Jan 23, 1989
TABLET; ORAL			
LEUCOVORIN CALCIUM			
EPIC PHARMA LLC	EQ 5MG BASE	A074544 001	Aug 28, 1997
	EQ 25MG BASE	A074544 002	Aug 28, 1997
IDT AUSTRALIA LTD	EQ 15MG BASE	A075327 001	Mar 24, 1999
PAR PHARM	EQ 5MG BASE	A071600 001	Oct 14, 1987
	EQ 25MG BASE	A071598 001	Oct 14, 1987
PHARMACHEMIE	EQ 5MG BASE	A073099 001	Mar 28, 1997
	EQ 25MG BASE	A073101 001	Mar 28, 1997
XANODYNE PHARM	EQ 5MG BASE	N018459 001	Jan 30, 1986
	EQ 10MG BASE	A071962 001	Nov 19, 1987
	EQ 15MG BASE	A071104 001	Mar 04, 1987
WELLCOVORIN			
+ GLAXOSMITHKLINE	EQ 5MG BASE **	N018342 001	Jul 08, 1983
+	EQ 25MG BASE **	N018342 002	Jul 08, 1983

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION

VIADUR

ORTHO MCNEIL JANSSEN	EQ 65MG BASE	N021088 001	Mar 03, 2000
----------------------	--------------	-------------	--------------

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

GENZYME	1MG/0.2ML	A075721 001	Nov 29, 2001
---------	-----------	-------------	--------------

LUPRON

ABBVIE ENDOCRINE INC	1MG/0.2ML	N019010 001	Apr 09, 1985
----------------------	-----------	-------------	--------------

LUPRON DEPOT

+	ABBVIE ENDOCRINE INC	3.75MG/VIAL **	N020011 001	Oct 22, 1990
---	----------------------	----------------	-------------	--------------

LUPRON DEPOT-PED

+	ABBVIE ENDOCRINE INC	3.75MG/VIAL, 7.5MG/VIAL **	N020263 003	Apr 16, 1993
---	----------------------	----------------------------	-------------	--------------

+	ABBVIE ENDOCRINE INC	7.5MG/VIAL, 7.5MG/VIAL **	N020263 004	Apr 16, 1993
---	----------------------	---------------------------	-------------	--------------

LEVALLORPHAN TARTRATE

INJECTABLE; INJECTION

LORFAN

ROCHE	1MG/ML	N010423 001	
-------	--------	-------------	--

LEVAMISOLE HYDROCHLORIDE

TABLET; ORAL

ERGAMISOL

JANSSEN PHARMA	EQ 50MG BASE	N020035 001	Jun 18, 1990
----------------	--------------	-------------	--------------

LEVETIRACETAM

SOLUTION; ORAL

LEVETIRACETAM

APOTEX INC	100MG/ML	A090187 001	Aug 05, 2011
------------	----------	-------------	--------------

TABLET; ORAL

LEVETIRACETAM

ACTAVIS LABS FL INC	250MG	A077408 001	Mar 02, 2009
---------------------	-------	-------------	--------------

	500MG	A077408 002	Mar 02, 2009
--	-------	-------------	--------------

	750MG	A077408 003	Mar 02, 2009
--	-------	-------------	--------------

MYLAN	250MG	A078731 001	Feb 10, 2009
-------	-------	-------------	--------------

	500MG	A078731 002	Feb 10, 2009
--	-------	-------------	--------------

	750MG	A078731 003	Feb 10, 2009
--	-------	-------------	--------------

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-221(of 375)

\*\* See List Footnote

LEVETIRACETAM

TABLET;ORAL

LEVETIRACETAM

SANDOZ	1GM 1GM 250MG 500MG 750MG	A078731 004 Feb 10, 2009 A077324 004 Jan 15, 2009 A077324 001 Jan 15, 2009 A077324 002 Jan 15, 2009 A077324 003 Jan 15, 2009
WATSON LABS INC	250MG 500MG 750MG 1GM	A078797 002 Jan 15, 2009 A078797 003 Jan 15, 2009 A078797 004 Jan 15, 2009 A078797 001 Jan 15, 2009

TABLET, EXTENDED RELEASE;ORAL

LEVETIRACETAM

MYLAN PHARMS INC	500MG 750MG 1GM	A200475 001 Dec 19, 2011 A200475 002 Dec 19, 2011 A200475 003 Dec 07, 2015
SANDOZ	500MG 750MG	A091668 001 Nov 01, 2012 A091668 002 Nov 01, 2012

LEVOBETAXOLOL HYDROCHLORIDESUSPENSION/DROPS;OPHTHALMIC  
BETAXON

ALCON PHARMS LTD	EQ 0.5% BASE	N021114 001 Feb 23, 2000
------------------	--------------	--------------------------

LEVOBUNOLOL HYDROCHLORIDESOLUTION/DROPS;OPHTHALMIC  
LEVOBUNOLOL HYDROCHLORIDE

ALCON LABS INC	0.25%	A074851 001 Oct 28, 1996
APOTEX INC	0.25%	A075473 001 Aug 03, 2000
	0.5%	A075475 001 Aug 03, 2000
BAUSCH AND LOMB	0.25%	A074307 001 Mar 04, 1994

LEVOBUPIVACAINE HYDROCHLORIDEINJECTABLE; INJECTION  
CHIROCAINE

PURDUE PHARMA LP	EQ 2.5MG BASE/ML EQ 5MG BASE/ML EQ 7.5MG BASE/ML	N020997 001 Aug 05, 1999 N020997 002 Aug 05, 1999 N020997 003 Aug 05, 1999
------------------	--	--

LEVOCABASTINE HYDROCHLORIDESUSPENSION/DROPS;OPHTHALMIC  
LIVOSTIN

NOVARTIS	EQ 0.05% BASE	N020219 001 Nov 10, 1993
----------	---------------	--------------------------

LEVOCARNITINEINJECTABLE; INJECTION  
LEVOCARNITINE

TEVA PHARMS USA	200MG/ML	A075881 001 Mar 29, 2001
SOLUTION;ORAL CARNITOR		

LEADIANT BIOSCI INC	1GM/10ML	N018948 002 Apr 27, 1988
---------------------	----------	--------------------------

LEVO CETIRIZINE DIHYDROCHLORIDE

TABLET;ORAL

LEVO CETIRIZINE DIHYDROCHLORIDE

SANDOZ	5MG	A090486 001 Mar 26, 2013
--------	-----	--------------------------

LEVODOPA

CAPSULE;ORAL

BENDOPA

VALEANT PHARM INTL	100MG 250MG 500MG	N016948 003 N016948 001 N016948 002
--------------------	-------------------------	---

DOPAR

SHIRE	100MG 250MG 500MG	N016913 003 N016913 001 N016913 002
-------	-------------------------	---

LARODOPA

ROCHE	100MG 250MG 500MG	N016912 002 N016912 001 N016912 006
-------	-------------------------	---

**DISCONTINUED DRUG PRODUCT LIST**

6-222(of 375)

\*\* See List Footnote

LEVODOPA

TABLET;ORAL

DOPAR

SHIRE

250MG

N016913 004

500MG

N016913 005

LARODOPA

ROCHE

100MG

N016912 005

250MG

N016912 003

500MG

N016912 004

LEVOFLOXACIN

INJECTABLE;INJECTION

LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ JANSSEN PHARMS	EQ 250MG/50ML (EQ 5MG/ML) **	N020635 002	Dec 20, 1996
+	EQ 500MG/100ML (EQ 5MG/ML) **	N020635 003	Dec 20, 1996
+	EQ 750MG/150ML (EQ 5MG/ML) **	N020635 005	Dec 20, 1996

LEVOFLOXACIN

AKORN

EQ 500MG/20ML (EQ 25MG/ML)

A091644 001 Jun 20, 2011

EQ 750MG/30ML (EQ 25MG/ML)

A091644 002 Jun 20, 2011

HOSPIRA INC

EQ 500MG/20ML (EQ 25MG/ML)

A078577 001 Aug 12, 2015

EQ 750MG/30ML (EQ 25MG/ML)

A078577 002 Aug 12, 2015

SOLUTION;ORAL

LEVAQUIN

+ JANSSEN PHARMS	250MG/10ML	N021721 001	Oct 21, 2004
------------------	------------	-------------	--------------

SOLUTION/DROPS;OPHTHALMIC

IQUIX

+ SANTEN	1.5% **	N021571 001	Mar 01, 2004
----------	---------	-------------	--------------

LEVOFLOXACIN

APOTEX INC

0.5%

A078282 001 Dec 20, 2010

QUIXIN

+ SANTEN	0.5% **	N021199 001	Aug 18, 2000
----------	---------	-------------	--------------

TABLET;ORAL

LEVOFLOXACIN

MYLAN

250MG

A076276 001 Jun 20, 2011

500MG

A076276 002 Jun 20, 2011

750MG

A077097 001 Jun 20, 2011

WATSON LABS INC

250MG

A201484 001 Nov 22, 2013

500MG

A201484 002 Nov 22, 2013

750MG

A201484 003 Nov 22, 2013

LEVOLEUCOVORIN CALCIUM

SOLUTION;IV (INFUSION)

FUSILEV

+ SPECTRUM PHARMS	EQ 175MG BASE/17.5ML (EQ 10MG BASE/ML) **	N020140 002	Apr 29, 2011
+	EQ 250MG BASE/25ML (EQ 10MG BASE/ML) **	N020140 003	Apr 29, 2011

LEVOMEPROMAZINE

INJECTABLE;INJECTION

LEVOPROME

IMMUNEX

20MG/ML

N015865 001

LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE;ORAL

ORLAAM

+	ROXANE	10MG/ML **	N020315 001	Jul 09, 1993
---	--------	------------	-------------	--------------

LEVONORDEFRIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE;INJECTION

ARESTOCAIN HYDROCHLORIDE W/ LEVONORDEFRIN

SOLVAY 0.05MG/ML;2%

A085010 001

CARBOCAINE W/ NEO-COBEFRIN

EASTMAN KODAK 0.05MG/ML;2%

N012125 002

ISOCAINE HYDROCHLORIDE W/ LEVONORDEFRIN

SEPTODONT INC 0.05MG/ML;2%

A084697 001

MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFRIN

BELMORA LLC 0.05MG/ML;2%

A084850 002 Oct 21, 1983

POLOCAINE W/ LEVONORDEFRIN

DENTSPLY PHARM 0.05MG/ML;2%

A089517 001 Apr 14, 1988

**DISCONTINUED DRUG PRODUCT LIST**

6-223(of 375)

\*\* See List Footnote

LEVONORDEFRIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAIN AND NOVOCAIN W/ NEO-COBESTRIN

EASTMAN KODAK 0.05MG/ML; 2%; 0.4%

N008592 007

LEVONORGESTREL

IMPLANT; IMPLANTATION

JADELLE

+ POPULATION COUNCIL 75MG/IMPLANT \*\*

N020544 001 Nov 01, 1996

LEVONORGESTREL

WYETH PHARMS INC 75MG/IMPLANT

N020627 001 Aug 15, 1996

NORPLANT

POPULATION COUNCIL 36MG/IMPLANT

N019897 001 Dec 10, 1990

NORPLANT SYSTEM IN PLASTIC CONTAINER

WYETH PHARMS INC 36MG/IMPLANT

N020088 001 Dec 10, 1990

TABLET; ORAL

LEVONORGESTREL

FDN CONSUMER 0.75MG \*\*

A078665 001 Aug 28, 2009

LUPIN LTD 0.75MG

A091328 001 Jan 23, 2013

WATSON LABS 0.75MG

A078666 001 Jun 24, 2009

PLAN B

+ FDN CONSUMER 0.75MG \*\*

N021045 001 Jul 28, 1999

+ 0.75MG \*\*

N021045 002 Aug 24, 2006

LEVOPOPOXYPHENE NAPSYLATE ANHYDROUS

CAPSULE; ORAL

NOVRAD

LILLY EQ 50MG BASE

N012928 006

EQ 100MG BASE

N012928 004

SUSPENSION; ORAL

NOVRAD

LILLY EQ 50MG BASE/5ML

N012928 002

LEVORPHANOL TARTRATE

INJECTABLE; INJECTION

LEVO-DROMORAN

VALEANT PHARM INTL 2MG/ML

N008719 001 Dec 19, 1991

TABLET; ORAL

LEVO-DROMORAN

+ VALEANT PHARM INTL 2MG \*\*

N008720 001 Dec 19, 1991

LEVOTHYROXINE SODIUM

SOLUTION; ORAL

TIROSINT-SOL

+ INSTITUT BIOCHIMIQUE 13MCG/ML

N206977 001 Dec 15, 2016

+ 25MCG/ML

N206977 002 Dec 15, 2016

+ 50MCG/ML

N206977 003 Dec 15, 2016

+ 75MCG/ML

N206977 004 Dec 15, 2016

+ 88MCG/ML

N206977 005 Dec 15, 2016

+ 100MCG/ML

N206977 006 Dec 15, 2016

+ 112MCG/ML

N206977 007 Dec 15, 2016

+ 125MCG/ML

N206977 008 Dec 15, 2016

+ 137MCG/ML

N206977 009 Dec 15, 2016

+ 150MCG/ML

N206977 010 Dec 15, 2016

+ 175MCG/ML

N206977 011 Dec 15, 2016

+ 200MCG/ML

N206977 012 Dec 15, 2016

TABLET; ORAL

LEVOLET

GENUS LIFESCIENCES 0.025MG

N021137 001 Jun 06, 2003

0.05MG

N021137 002 Jun 06, 2003

0.075MG

N021137 003 Jun 06, 2003

0.088MG

N021137 004 Jun 06, 2003

0.1MG

N021137 005 Jun 06, 2003

0.112MG

N021137 006 Jun 06, 2003

0.125MG

N021137 007 Jun 06, 2003

0.137MG

N021137 008 Jun 06, 2003

0.15MG

N021137 009 Jun 06, 2003

0.175MG

N021137 010 Jun 06, 2003

0.2MG

N021137 011 Jun 06, 2003

0.3MG

N021137 012 Jun 06, 2003

**DISCONTINUED DRUG PRODUCT LIST**

6-224(of 375)

\*\* See List Footnote

LEVOOTHYROXINE SODIUM

TABLET;ORAL

LEVOOTHYROXINE SODIUM

MERCK KGAA

0.025MG  
0.05MG  
0.075MG  
0.088MG  
0.1MG  
0.112MG  
0.125MG  
0.15MG  
0.175MG  
0.2MG  
0.3MG

A076752 001 Jun 16, 2005  
A076752 002 Jun 16, 2005  
A076752 003 Jun 16, 2005  
A076752 004 Jun 16, 2005  
A076752 005 Jun 16, 2005  
A076752 006 Jun 16, 2005  
A076752 007 Jun 16, 2005  
A076752 008 Jun 16, 2005  
A076752 009 Jun 16, 2005  
A076752 010 Jun 16, 2005  
A076752 011 Jun 16, 2005

LEVOXYL

+ KING PHARMS

0.3MG \*\*

N021301 012 May 25, 2001

NOVOTHYROX

MERCK KGAA

0.025MG  
0.05MG  
0.075MG  
0.088MG  
0.1MG  
0.112MG  
0.125MG  
0.137MG  
0.15MG  
0.175MG  
0.2MG  
0.3MG

N021292 001 May 31, 2002  
N021292 002 May 31, 2002  
N021292 003 May 31, 2002  
N021292 004 May 31, 2002  
N021292 005 May 31, 2002  
N021292 006 May 31, 2002  
N021292 007 May 31, 2002  
N021292 008 May 31, 2002  
N021292 009 May 31, 2002  
N021292 010 May 31, 2002  
N021292 011 May 31, 2002  
N021292 012 May 31, 2002

THYRO-TABS

+ LLOYD

0.025MG \*\*  
0.05MG \*\*  
0.075MG \*\*  
0.088MG \*\*  
0.1MG \*\*  
0.112MG \*\*  
0.125MG \*\*  
0.137MG \*\*  
0.15MG \*\*  
0.175MG \*\*  
0.2MG \*\*  
0.3MG \*\*

N021116 001 Oct 24, 2002  
N021116 002 Oct 24, 2002  
N021116 003 Oct 24, 2002  
N021116 010 Oct 24, 2002  
N021116 004 Oct 24, 2002  
N021116 011 Oct 24, 2002  
N021116 005 Oct 24, 2002  
N021116 012 Dec 07, 2004  
N021116 006 Oct 24, 2002  
N021116 007 Oct 24, 2002  
N021116 008 Oct 24, 2002  
N021116 009 Oct 24, 2002

LIDOCAINE

AEROSOL;ORAL

XYLOCAINE

ASTRAZENECA

10%

N014394 001

FILM, EXTENDED RELEASE;BUCCAL

DENTIPATCH

NOVEN

23MG/PATCH

N020575 001 May 21, 1996

OINTMENT;TOPICAL

ALPHACAIN

CARLISLE

5%

A084944 001

5%

A084946 001

5%

A084947 001

LIDOCAINE

BELMORA LLC

5%

A080210 001

XYLOCAINE

+ ASTRAZENECA

5% \*\*

N008048 001

PATCH;TOPICAL

DENTIPATCH

NOVEN

46.1MG/PATCH

N020575 002 May 21, 1996

SOLUTION;TOPICAL

XYLOCAINE

ASTRAZENECA

5%

N014127 001

SUPPOSITORY;RECTAL

XYLOCAINE

ASTRAZENECA

100MG

N013077 001

**DISCONTINUED DRUG PRODUCT LIST**

6-225(of 375)

\*\* See List Footnote

LIDOCAINE HYDROCHLORIDE

## INJECTABLE; INJECTION

## ALPHACAINE HYDROCHLORIDE

CARLISLE	2%	A084721 001
LIDOCAINE HYDROCHLORIDE		
ABBOTT	10%	A087980 001 Feb 02, 1983
	20%	A089362 001 May 25, 1988
ABRAXIS PHARM	1%	A080420 001
	1%	A086761 001
	1.5%	A080420 005
	2%	A080420 002
	2%	A080420 004
	2%	A086761 002
	2%	N017508 001
	4%	N017508 002
	20%	N017508 004
AKORN	1%	A085037 001
	2%	A085037 002
BEL MAR	1%	A080710 001
	2%	A080760 001
BELMORA LLC	2%	A080504 001
DELL LABS	1%	A083387 001
	2%	A083388 001
ELKINS SINK	0.5%	A085131 001
	4%	A084626 001
GD SEARLE LLC	1%	A083135 001
	2%	A083135 002
HOSPIRA	1%	A040013 001 Jun 23, 1995
	1.5%	A088330 001 May 17, 1984
	2%	A088331 001 May 17, 1984
INTL MEDICATION	1%	N017701 002
	2%	N017701 001
	1GM/VIAL	N018543 001
	2GM/VIAL	N018543 002
LUITPOLD	2%	A083198 001
LYPHOMED	1%	A080390 001
	2%	A080390 002
MILES	1%	A080414 001
	2%	A080414 002
MYLAN LABS LTD	0.5%	A091056 001 Dec 08, 2010
	0.5%	A091058 001 Sep 30, 2010
	1%	A091056 002 Dec 08, 2010
	1%	A091058 002 Sep 30, 2010
	2%	A202242 001 Apr 11, 2014
WATSON LABS	1%	A080377 001
	1%	A083627 001
	2%	A080377 002
	2%	A083627 002
WEST-WARD PHARMS INT	1%	A080407 001
	2%	A080407 002
WYETH AYERST	1%	A083083 001
	2%	A083083 002
LIDOCAINE HYDROCHLORIDE 0.1% AND DEXTROSE 5% IN PLASTIC CONTAINER		
BAXTER HLTHCARE	100MG/100ML	N018461 001
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER		
B BRAUN	200MG/100ML	N018967 001 Mar 30, 1984
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5%		
HOSPIRA	200MG/100ML	A083158 005
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER		
ABBOTT	200MG/100ML	N018954 001 Jul 09, 1985
HOSPIRA	200MG/100ML	N018388 001
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER		
B BRAUN	400MG/100ML	N018967 002 Mar 30, 1984
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5%		
HOSPIRA	400MG/100ML	A083158 006
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER		
HOSPIRA	400MG/100ML	N018388 002
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER		
B BRAUN	800MG/100ML	N018967 003 Mar 30, 1984

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-226(of 375)

\*\* See List Footnote

LIDOCAINE HYDROCHLORIDE

## INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER HOSPIRA	800MG/100ML	N018388 003 Nov 05, 1982
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER HOSPIRA	1.5%	A088326 001 Jul 31, 1984
	10%	A088367 001 Jul 31, 1984
	20%	A088368 001 Jul 31, 1984
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE INT'L MEDICATION	4%	N017702 002
MYLAN LABS LTD	2%	A090665 001 Sep 27, 2010
WEST-WARD PHARMS INT	1%	A084625 001
	2%	A084625 002
LIDOCATON PHARMATON	2%	A084727 001 Aug 17, 1983
LIDOPEN MERIDIAN MEDCL TECHN	10%	N017549 001
XYLOCAINE ASTRAZENECA	1%	N010418 005
	1.5%	N010418 009
	2%	N010418 007
XYLOCAINE DENTAL DENTSPLY PHARM	2%	N021380 001
XYLOCAINE PRESERVATIVE FREE FRESENIUS KABI USA	10%	N016801 003
INJECTABLE; SPINAL XYLOCAINE 1.5% W/ DEXTROSE 7.5% FRESENIUS KABI USA	1.5%	N016297 001
XYLOCAINE 5% W/ GLUCOSE 7.5% ASTRAZENECA	5%	N010496 002 Jul 07, 1982
JELLY; TOPICAL ANESTACON	2%	A080429 001
LIDOCAINE HYDROCHLORIDE G AND W LABS INC	2%	A081318 001 Apr 29, 1993
SOLUTION; ORAL LIDOCAINE HYDROCHLORIDE VISCOSUS ACTAVIS MID ATLANTIC	2%	A086578 001
INT'L MEDICATION	2%	A086389 001 Feb 02, 1982
XYLOCAINE VISCOSUS FRESENIUS KABI USA	2% **	N009470 001
SOLUTION; TOPICAL LARYNGOTRACHEAL ANESTHESIA KIT KENDALL IL	4%	A087931 001 Jun 10, 1983
LIDOCAINE HYDROCHLORIDE PACO	4%	A089688 001 Jun 30, 1989
LTA II KIT HOSPIRA	4%	A088542 001 Jul 31, 1984
PEDIATRIC LTA KIT ABBOTT	2%	A088572 001 Jul 31, 1984
HOSPIRA	2%	A085995 001

LIDOCAINE HYDROCHLORIDE; OXYTETRACYCLINE

## INJECTABLE; INJECTION

TERRAMYCIN PFIZER	2%;50MG/ML	A060567 001
	2%;125MG/ML	A060567 002

LIDOCAINE; PRilocaine

## DISC; TOPICAL

EMLA ASTRAZENECA	2.5%;2.5%	N020962 001 Feb 04, 1998
---------------------	-----------	--------------------------

LINCOMYCIN HYDROCHLORIDE

## CAPSULE; ORAL

LINCOCIN PHARMACIA AND UPJOHN	EQ 250MG BASE	N050316 001
	EQ 500MG BASE	N050316 002

**DISCONTINUED DRUG PRODUCT LIST**

6-227(of 375)

\*\* See List Footnote

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE

WATSON LABS

EQ 300MG BASE/ML

A063180 001 Apr 16, 1991

LINDANE

CREAM; TOPICAL

K WELL

REED AND CARNICK

1%

A084218 001

1%

N006309 001

LOTION; TOPICAL

GAMENE

SOLA BARNES HIND

1%

A084989 001

K WELL

REED AND CARNICK

1%

A084218 002

1%

N006309 003

SCABENE

STIEFEL

1%

A086769 001

SHAMPOO; TOPICAL

GAMENE

SOLA BARNES HIND

1%

A084988 001

K WELL

REED AND CARNICK

1%

A084219 001

1%

N010718 001

SCABENE

STIEFEL

1%

A087940 001 Apr 08, 1983

LINEZOLID

TABLET; ORAL

ZYVOX

+ PHARMACIA AND UPJOHN 400MG \*\*

N021130 001 Apr 18, 2000

LIOTHYRONINE SODIUM

TABLET; ORAL

LIOTHYRONINE SODIUM

WATSON LABS

EQ 0.025MG BASE

A085755 001 Jan 25, 1982

EQ 0.05MG BASE

A085753 001 Feb 03, 1982

LIOTRIX (T4;T3)

TABLET; ORAL

EUTHROID-0.5

PARKE DAVIS

0.03MG; 0.0075MG

N016680 001

EUTHROID-1

PARKE DAVIS

0.06MG; 0.015MG

N016680 002

EUTHROID-2

PARKE DAVIS

0.12MG; 0.03MG

N016680 003

EUTHROID-3

PARKE DAVIS

0.18MG; 0.045MG

N016680 004

THYROLAR-5

FOREST LABS

0.25MG; 0.0625MG

N016807 006

LISINOPRIL

TABLET; ORAL

LISINOPRIL

SANDOZ

2.5MG

A075903 001 Jul 01, 2002

2.5MG

A075999 001 Jul 01, 2002

5MG

A075903 002 Jul 01, 2002

5MG

A075999 002 Jul 01, 2002

10MG

A075903 003 Jul 01, 2002

10MG

A075999 003 Jul 01, 2002

20MG

A075903 004 Jul 01, 2002

20MG

A075999 004 Jul 01, 2002

30MG

A075903 005 Jul 01, 2002

30MG

A075999 005 Jul 01, 2002

40MG

A075903 006 Jul 01, 2002

40MG

A075999 006 Jul 01, 2002

TEVA

2.5MG

A075783 001 Jul 01, 2002

5MG

A075783 002 Jul 01, 2002

10MG

A075783 003 Jul 01, 2002

20MG

A075783 004 Jul 01, 2002

30MG

A075783 005 Jul 01, 2002

40MG

A075783 006 Jul 01, 2002

**DISCONTINUED DRUG PRODUCT LIST**

6-228(of 375)

\*\* See List Footnote

LISINOPRIL

TABLET;ORAL PRINIVIL MERCK	2.5MG	N019558 006 Jan 28, 1994
----------------------------------	-------	--------------------------

LITHIUM CARBONATE

CAPSULE;ORAL ESKALITH NOVEN THERAP LITHIUM CARBONATE ABLE	300MG 150MG 300MG 300MG 600MG	N016860 001 A076823 001 Jun 29, 2004 A076121 001 Sep 27, 2001 A076823 002 Jun 29, 2004 A076823 003 Jun 29, 2004
APOTEX INC USL PHARMA WATSON LABS	300MG 300MG 300MG	A076795 001 Nov 22, 2004 A072542 001 Feb 01, 1989 A070407 001 Mar 19, 1987
LITHONATE SOLVAY	300MG	N016782 001
TABLET;ORAL ESKALITH JDS PHARMS LITHANE BAYER PHARMS	300MG	N017971 001 N018833 001 Jul 18, 1985
LITHIUM CARBONATE HIKMA INTL PHARMS PFIZER	300MG 300MG	A078715 001 Dec 28, 2010 N016834 001
LITHOTABS SOLVAY	300MG	N016980 001
TABLET, EXTENDED RELEASE;ORAL ESKALITH CR JDS PHARMS LITHIUM CARBONATE ABLE BARR	450MG ** 300MG 300MG 450MG	N018152 001 Mar 29, 1982 A076382 001 Apr 21, 2003 A076170 001 Jun 10, 2002 A076366 001 Aug 21, 2003
HIKMA INTL PHARMS	450MG	A076490 001 Jun 17, 2003

LITHIUM CITRATE

SYRUP;ORAL LITHONATE SOLVAY	EQ 300MG CARBONATE/5ML	N017672 001
-----------------------------------	------------------------	-------------

LOMEFLOXACIN HYDROCHLORIDE

TABLET;ORAL MAXAQWIN PHARMACIA	EQ 400MG BASE	N020013 001 Feb 21, 1992
--------------------------------------	---------------	--------------------------

LOPERAMIDE HYDROCHLORIDE

CAPSULE;ORAL IMODIUM J AND J CONSUMER INC	2MG ** 2MG **	N017690 001 N017694 001
LOPERAMIDE HYDROCHLORIDE FOSUN PHARMA ROXANE TEVA	2MG 2MG 2MG	A072993 001 Aug 28, 1992 A073080 001 Nov 27, 1991 A073122 001 Aug 30, 1991
SOLUTION;ORAL IMODIUM JANSSEN PHARMS	1MG/5ML	N019037 001 Jul 31, 1984
LOPERAMIDE HYDROCHLORIDE ALPHARMA US PHARMS DURAMED PHARMS BARR TEVA WATSON LABS	1MG/5ML 1MG/5ML 1MG/5ML 1MG/5ML	A073187 001 Sep 15, 1992 A074991 001 Dec 29, 1997 A073478 001 Jun 23, 1995 A073062 001 May 28, 1993
TABLET;ORAL LOPERAMIDE HYDROCHLORIDE ABLE CONTRACT PHARMACAL PERRIGO	2MG 2MG 2MG	A073528 001 Nov 30, 1993 A073254 001 Jul 30, 1993 A074194 001 Oct 30, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-229(of 375)

\*\* See List Footnote

LOPERAMIDE HYDROCHLORIDE

TABLET, CHEWABLE;ORAL  
 IMODIUM A-D EZ CHEWS  
 + J AND J CONSUMER INC 2MG  
 N020448 001 Jul 24, 1997

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET, CHEWABLE;ORAL  
 IMODIUM MULTI-SYMPOTM RELIEF  
 + J AND J CONSUMER INC 2MG;125MG  
 N020606 001 Jun 26, 1996

LOPINAVIR; RITONAVIR

CAPSULE;ORAL  
 KALETRA  
 ABBVIE 133.3MG;33.3MG  
 N021226 001 Sep 15, 2000

LORACARBEF

CAPSULE;ORAL  
 LORABID  
 KING PHARMS 200MG  
 400MG  
 N050668 001 Dec 31, 1991  
 N050668 002 Apr 05, 1996  
 FOR SUSPENSION;ORAL  
 LORABID  
 KING PHARMS 100MG/5ML  
 200MG/5ML  
 N050667 001 Dec 31, 1991  
 N050667 002 Dec 31, 1991

LORATADINE

SYRUP;ORAL  
 CLARITIN HIVES RELIEF  
 + BAYER HEALTHCARE LLC 1MG/ML \*\*  
 N020641 003 Nov 19, 2003  
 LORATADINE  
 APOTEX INC 1MG/ML  
 RANBAXY LABS LTD 1MG/ML  
 A075565 001 Oct 05, 2004  
 A076529 001 Aug 20, 2004  
 TABLET;ORAL  
 LORATADINE  
 PERRIGO 10MG  
 N021512 001 Jun 24, 2004

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE;ORAL  
 LORATADINE AND PSEUDOEPHEDRINE SULFATE  
 ACTAVIS LABS FL INC 5MG;120MG  
 A076208 001 Jan 28, 2004

LORAZEPAM

INJECTABLE;INJECTION  
 LORAZEPAM  
 AKORN 2MG/ML  
 BEDFORD 2MG/ML  
 4MG/ML  
 DAVA PHARMS INC 2MG/ML  
 4MG/ML  
 HOSPIRA 2MG/ML  
 2MG/ML  
 4MG/ML  
 4MG/ML  
 SAGENT AGILA LLC 2MG/ML  
 2MG/ML  
 4MG/ML  
 4MG/ML  
 WATSON LABS 2MG/ML  
 4MG/ML  
 WATSON LABS INC 1MG/0.5ML  
 2MG/ML  
 2MG/ML  
 4MG/ML  
 4MG/ML  
 WEST-WARD PHARMS INT 2MG/ML  
 4MG/ML  
 A074974 001 Jul 23, 1998  
 A077076 001 Jul 13, 2005  
 A077076 002 Jul 13, 2005  
 A074793 001 Mar 16, 2000  
 A074793 002 Mar 16, 2000  
 A074280 001 May 27, 1994  
 A074300 001 Apr 12, 1994  
 A074280 002 May 27, 1994  
 A074300 003 Mar 19, 1997  
 A200217 001 Apr 04, 2017  
 A200542 001 Apr 28, 2017  
 A200217 002 Apr 04, 2017  
 A200542 002 Apr 28, 2017  
 A074276 001 Apr 15, 1994  
 A074276 002 Apr 15, 1994  
 A074551 003 Sep 12, 1996  
 A074535 001 Sep 12, 1996  
 A074551 001 Sep 12, 1996  
 A074535 002 Sep 12, 1996  
 A074551 002 Sep 12, 1996  
 A074496 001 Sep 28, 1998  
 A074496 002 Sep 28, 1998

SOLUTION;ORAL  
 LORAZEPAM  
 ROXANE 0.5MG/5ML  
 A074648 001 Mar 18, 1997

TABLET;ORAL  
 LORAZ  
 QUANTUM PHARMICS 0.5MG  
 A070200 001 Aug 09, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-230(of 375)

\*\* See List Footnote

LORAZEPAMTABLET;ORAL  
LORAZ

	1MG	A070201 001 Aug 09, 1985
	2MG	A070202 001 Aug 09, 1985
LORAZEPAM		
AM THERAP	0.5MG	A070727 001 Mar 07, 1986
	1MG	A070728 001 Mar 07, 1986
	2MG	A070729 001 Mar 07, 1986
ANDA REPOSITORY	0.5MG	A072553 001 Mar 29, 1991
	1MG	A072554 001 Mar 29, 1991
	2MG	A072555 001 Mar 29, 1991
HALSEY	0.5MG	A071434 001 Sep 01, 1987
	1MG	A071435 001 Sep 01, 1987
	2MG	A071436 001 Sep 01, 1987
MUTUAL PHARM	0.5MG	A070472 001 Dec 10, 1985
	1MG	A070473 001 Dec 10, 1985
	2MG	A070474 001 Dec 10, 1985
MYLAN	2MG	A071591 001 Oct 13, 1987
PAR PHARM	0.5MG	A070675 001 Dec 01, 1986
	1MG	A070676 001 Dec 01, 1986
	2MG	A070677 001 Dec 01, 1986
SANDOZ	0.5MG	A071193 001 Apr 15, 1988
	1MG	A071194 001 Apr 15, 1988
	2MG	A071195 001 Apr 15, 1988
SUPERPHARM	0.5MG	A071245 001 Feb 09, 1987
	1MG	A071246 001 Feb 09, 1987
	2MG	A071247 001 Feb 09, 1987
USL PHARMA	1MG	A070539 001 Dec 22, 1986
	2MG	A070540 001 Dec 22, 1986
WARNER CHILCOTT	1MG	A071038 001 Jan 12, 1988
	2MG	A071039 001 Jan 12, 1988
WATSON LABS	0.5MG	A071086 001 Mar 23, 1987
	0.5MG	A071117 001 Jul 24, 1986
	1MG	A071087 001 Mar 23, 1987
	1MG	A071118 001 Jul 24, 1986
	2MG	A071088 001 Mar 23, 1987
	2MG	A071110 001 Jul 24, 1986

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS;OPHTHALMIC

LOTEMAX

PHARMOS	0.5%	N020841 001 Mar 09, 1998
---------	------	--------------------------

LOVASTATIN

TABLET;ORAL

LOVASTATIN

MYLAN	10MG	A075935 001 Dec 17, 2001
	20MG	A075935 002 Dec 17, 2001
	40MG	A075935 003 Dec 17, 2001

MEVACOR

+ MERCK	10MG **	N019643 002 Mar 28, 1991
+	20MG **	N019643 003 Aug 31, 1987
+	40MG **	N019643 004 Dec 14, 1988

TABLET, EXTENDED RELEASE;ORAL

ALTOPREV

COVIS PHARMA BV	10MG	N021316 001 Jun 26, 2002
-----------------	------	--------------------------

LOXAPINE HYDROCHLORIDE

CONCENTRATE;ORAL

LOXITANE C

ACTAVIS LABS UT INC	EQ 25MG BASE/ML	N017658 001
---------------------	-----------------	-------------

INJECTABLE;INJECTION

LOXITANE IM

ACTAVIS LABS UT INC	EQ 50MG BASE/ML	N018039 001
---------------------	-----------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-231(of 375)

\*\* See List Footnote

LOXAPINE SUCCINATE

CAPSULE;ORAL

LOXITANE

+ ACTAVIS LABS UT INC	EQ 5MG BASE **	N017525 001
+	EQ 10MG BASE **	N017525 002
+	EQ 25MG BASE **	N017525 003
+	EQ 50MG BASE **	N017525 004

TABLET;ORAL

LOXITANE

+ ACTAVIS LABS UT INC	EQ 10MG BASE **	N017525 006
+	EQ 25MG BASE **	N017525 007
+	EQ 50MG BASE **	N017525 008

LUCINACTANT

SUSPENSION;INTRATRACHEAL

SURFAXIN

WINDTREE THERAP 8.5ML N021746 001 Mar 06, 2012

LUTROPIN ALFA

INJECTABLE;SUBCUTANEOUS

LUVERIS

EMD SERONO 75 IU/VIAL N021322 001 Oct 08, 2004

LYPRESSIN

SOLUTION;NASAL

DIAPID

NOVARTIS 0.185MG/ML N016755 001

MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE;INJECTION

PLASMA-LYTE 56 IN PLASTIC CONTAINER

BAXTER HLTHCARE 32MG/100ML;128MG/100ML;234MG/100ML N019047 001 Jun 15, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE;INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

B BRAUN 30MG/100ML;37MG/100ML;0.82MG/100ML;370M G/100ML;530MG/100ML;500MG/100ML;12MG/10 OML N019006 001 Apr 04, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE;INJECTION

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN 30MG/100ML;37MG/100ML;370MG/100ML;530MG /100ML;500MG/100ML N018252 001

SOLUTION;IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA INC 14MG/100ML;37MG/100ML;222MG/100ML;526MG /100ML;502MG/100ML N018406 001

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA INC 30MG/100ML;37MG/100ML;222MG/100ML;526MG /100ML;502MG/100ML N018406 002 Jul 08, 1982

SYNOVALYTE IN PLASTIC CONTAINER

BAXTER HLTHCARE 30MG/100ML;37MG/100ML;368MG/100ML;526MG /100ML;502MG/100ML N019326 001 Jan 25, 1985

MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET;ORAL

MAGNESIUM HYDROXIDE AND OMEPRAZOLE AND SODIUM BICARBONATE

SANTARUS 343MG;20MG;750MG N022456 001 Dec 04, 2009  
343MG;40MG;750MG N022456 002 Dec 04, 2009

TABLET, CHEWABLE;ORAL

ZEGERID

SANTARUS 700MG;20MG;600MG N021850 001 Mar 24, 2006  
700MG;40MG;600MG N021850 002 Mar 24, 2006MAGNESIUM SULFATE; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; POTASSIUM SULFATE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

SOLUTION;ORAL

SUCLEAR

+ BRAINTREE LABS 1.6GM/BOT,3.13GM/BOT,17.5GM/BOT,N/A,N/A ,N/A,N/A;N/A,N/A,N/A,N/A,210GM,0.74GM,2.86G M,5.6GM \*\* N203595 001 Jan 18, 2013

**DISCONTINUED DRUG PRODUCT LIST**

6-232(of 375)

\*\* See List Footnote

MALATHION

LOTION;TOPICAL

MALATHION

MYLAN PHARMS INC 0.5%

A078743 001 Mar 06, 2009

MANGAFODIPIR TRISODIUM

INJECTABLE;INJECTION

TESLASCAN

IC TARGETS 37.9MG/ML

N020652 001 Nov 26, 1997

MANGANESE CHLORIDE TETRAHYDRATE

FOR SOLUTION;ORAL

LUMENHANCE

BRACCO 3.49MG/GM

N020686 001 Dec 19, 1997

MANGANESE SULFATE

INJECTABLE;INJECTION

MANGANESE SULFATE

ABRAXIS PHARM EQ 0.1MG MANGANESE/ML

N019228 001 May 05, 1987

MANNITOL

INJECTABLE;INJECTION

MANNITOL 10%

B BRAUN 10GM/100ML

N016080 002

HOSPIRA 10GM/100ML

N016269 002

MILES 10GM/100ML

N016472 002

MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER

B BRAUN 10GM/100ML

N016080 006

MANNITOL 15%

B BRAUN 15GM/100ML

N016080 003

HOSPIRA 15GM/100ML

N016269 003

MILES 15GM/100ML

N016472 005

MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%

B BRAUN 15GM/100ML

N016080 005

MANNITOL 20%

B BRAUN 20GM/100ML

N014738 001

20GM/100ML

N016080 004

HOSPIRA 20GM/100ML

N016269 004

MILES 20GM/100ML

N016472 004

MANNITOL 25%

ABRAXIS PHARM 12.5GM/50ML

A086754 001

HOSPIRA 12.5GM/50ML

N016269 005

IGI LABS INC 12.5GM/50ML

A089239 001 May 06, 1987

12.5GM/50ML

A089240 001 May 06, 1987

MERCK 12.5GM/50ML

N005620 001

WATSON LABS 12.5GM/50ML

A087460 001 Jun 27, 1983

MANNITOL 5%

B BRAUN 5GM/100ML

N016080 001

HOSPIRA 5GM/100ML

N016269 001

MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%

B BRAUN 5GM/100ML

N016080 007

POWDER;INHALATION

ARIDOL KIT

PHARMAXIS LTD N/A,5MG,10MG,20MG,40MG

N022368 001 Oct 05, 2010

SOLUTION;IRRIGATION

RESECTISOL

B BRAUN 5GM/100ML

N016704 002

MANNITOL; SORBITOL

SOLUTION;IRRIGATION

SORBITOL-MANNITOL

HOSPIRA 540MG/100ML;2.7GM/100ML

A080224 001

SORBITOL-MANNITOL IN PLASTIC CONTAINER

HOSPIRA 540MG/100ML;2.7GM/100ML

N017636 001

MAPROTILINE HYDROCHLORIDE

TABLET;ORAL

LUDIOMIL

NOVARTIS 25MG

N017543 001

50MG

N017543 002

75MG

N017543 003 Sep 30, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-233(of 375)

\*\* See List Footnote

MAPROTILINE HYDROCHLORIDE

TABLET;ORAL

MAPROTILINE HYDROCHLORIDE

AM THERAP	25MG	A072129 001	Jan 14, 1988
	50MG	A072130 001	Jan 14, 1988
	75MG	A072131 001	Jan 14, 1988
WATSON LABS	25MG	A071943 001	Dec 30, 1987
	50MG	A071944 001	Dec 30, 1987
	75MG	A071945 001	Dec 30, 1987
WATSON LABS TEVA	25MG	A072164 001	Jun 01, 1988
	50MG	A072162 001	Jun 01, 1988
		A072163 001	Jun 01, 1988

MASOPROCOL

CREAM;TOPICAL

ACTINEX

UNIV AZ CANCER CTR	10%	N019940 001	Sep 04, 1992
--------------------	-----	-------------	--------------

MAZINDOL

TABLET;ORAL

MAZANOR

WYETH AYERST	1MG	N017980 002	
	2MG	N017980 001	
SANOREX			
+ NOVARTIS	1MG **	N017247 001	
+	2MG **	N017247 002	

MEBENDAZOLE

TABLET, CHEWABLE;ORAL

VERMOX

+ JANSSEN PHARMS	100MG **	N017481 001
------------------	----------	-------------

MEBUTAMATE

TABLET;ORAL

DORMATE

MEDPOINTE PHARM HLC	600MG	N017374 001
---------------------	-------	-------------

MECAMYLAMINE HYDROCHLORIDE

TABLET;ORAL

INVERSINE

+ TARGACEPT	2.5MG **	N010251 001
-------------	----------	-------------

MECASERMIN RINFABATE RECOMBINANT

INJECTABLE;SUBCUTANEOUS

IPLEX

INSMED	36MG/0.6ML	N021884 001	Dec 12, 2005
--------	------------	-------------	--------------

MECLIZINE HYDROCHLORIDE

TABLET;ORAL

ANTIVERT

CASPER PHARMA LLC	12.5MG	N010721 006	
	25MG	N010721 004	
	50MG	N010721 001	Jan 20, 1982

MECLIZINE HYDROCHLORIDE

ABC HOLDING	12.5MG	A085253 001	
	25MG	A085252 001	
AMNEAL PHARMS	50MG	A201451 003	Feb 23, 2011
ANABOLIC	25MG	A085891 001	
ANI PHARMS INC	12.5MG	A084975 001	
	25MG	A084657 001	
BUNDY	12.5MG	A084382 001	
	25MG	A084872 001	
IVAX SUB TEVA PHARMS	12.5MG	A083784 001	
KV PHARM	12.5MG	A085524 001	
	25MG	A085523 001	
MYLAN PHARMS INC	50MG	A202640 003	Sep 17, 2012
PAR PHARM	50MG	A089674 001	Mar 31, 1988
PLIVA	12.5MG	A088732 001	Dec 11, 1985
	25MG	A088734 001	Dec 11, 1985
RISING PHARMS INC	12.5MG	A040179 001	Jan 30, 1997
	25MG	A040179 002	Jan 30, 1997
SUPERPHARM	12.5MG	A089113 001	Aug 20, 1985
	25MG	A089114 001	Aug 20, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-234(of 375)

\*\* See List Footnote

MECLIZINE HYDROCHLORIDE

TABLET;ORAL

MECLIZINE HYDROCHLORIDE

UDL	12.5MG 25MG	A088256 001 Jun 13, 1983 A088257 001 Jun 13, 1983
VANGARD	12.5MG 25MG	A087877 001 Apr 20, 1982 A087620 001 Jan 04, 1982
WATSON LABS	12.5MG 12.5MG 25MG	A085195 001 A085269 001 A085740 001

TABLET, CHEWABLE;ORAL

ANTIVERT

CASPER PHARMA LLC	25MG	N010721 005
MECLIZINE HYDROCHLORIDE		
IVAX SUB TEVA PHARMS	25MG	A084976 001
NEXGEN PHARMA INC	25MG	A086392 001

PLIVA 25MG A088733 001 Dec 11, 1985

MECLOCYCLINE SULFOSALICYLATE

CREAM;TOPICAL

MECLAN

JOHNSON AND JOHNSON 1% N050518 001

MECLOFENAMATE SODIUM

CAPSULE;ORAL

MECLODUM

QUANTUM PHARMICS EQ 50MG BASE A071380 001 Jul 14, 1987  
EQ 100MG BASE A071381 001 Jul 14, 1987

MECLOFENAMATE SODIUM		
AM THERAP	EQ 50MG BASE	A071362 001 Feb 10, 1987
	EQ 100MG BASE	A071363 001 Feb 10, 1987
BARR	EQ 50MG BASE	A072848 001 Mar 20, 1989
	EQ 100MG BASE	A072809 001 Mar 20, 1989
PAR PHARM	EQ 50MG BASE	A072077 001 Mar 10, 1988
	EQ 100MG BASE	A072078 001 Mar 10, 1988
SANDOZ	EQ 50MG BASE	A072262 001 Nov 29, 1988
	EQ 100MG BASE	A072263 001 Nov 29, 1988
USL PHARMA	EQ 50MG BASE	A071007 001 Mar 25, 1988
	EQ 100MG BASE	A071008 001 Mar 25, 1988
VITARINE	EQ 50MG BASE	A071710 001 Jun 15, 1988
	EQ 100MG BASE	A071684 001 Jun 15, 1988
WATSON LABS	EQ 50MG BASE	A070400 001 Nov 25, 1986
	EQ 50MG BASE	A071468 001 Apr 15, 1987
	EQ 50MG BASE	A071640 001 Aug 11, 1987
	EQ 100MG BASE	A070401 001 Nov 25, 1986
	EQ 100MG BASE	A071641 001 Aug 11, 1987
WATSON LABS TEVA	EQ 100MG BASE	A071469 001 Apr 15, 1987
MECLOMEN		
PARKE DAVIS	EQ 50MG BASE	N018006 001
	EQ 100MG BASE	N018006 002

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

+ PHARMACIA AND UPJOHN 100MG/ML \*\* N012541 002

MEDROXYPROGESTERONE ACETATE		
SANDOZ INC	150MG/ML	A078711 001 May 20, 2009
TEVA PHARMS USA	150MG/ML	A076552 001 Oct 27, 2004

TABLET;ORAL

AMEN

AMARIN PHARMS 10MG A083242 001

CURRETAB	10MG	A085686 001
SOLVAY	10MG	
CYCRIN		
ESI	2.5MG	A081239 001 Oct 30, 1992
	5MG	A081240 001 Oct 30, 1992
	10MG	A089386 001 Sep 09, 1987

MEDROXYPROGESTERONE ACETATE		
DURAMED PHARMS BARR	2.5MG	A040311 001 Dec 01, 1999
	5MG	A040311 002 Dec 01, 1999
	10MG	A040311 003 Dec 01, 1999

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-235(of 375)

\*\* See List Footnote

MEDROXYPROGESTERONE ACETATE

TABLET;ORAL

MEDROXYPROGESTERONE ACETATE

USL PHARMA 10MG

A088484 001 Jul 26, 1984

MEDRYSONE

SUSPENSION;OPHTHALMIC

HMS

ALLERGAN 1%

N016624 003

MEFLOQUINE HYDROCHLORIDE

TABLET;ORAL

LARIAM

+ ROCHE 250MG \*\*

N019591 001 May 02, 1989

MEFLOQUINE HYDROCHLORIDE

HIKMA INTL PHARMS 250MG

A077699 001 Apr 21, 2010

SANDOZ 250MG

A076175 001 Feb 20, 2002

US ARMY WALTER REED 250MG \*\*

N019578 001 May 02, 1989

MEGESTROL ACETATE

SUSPENSION;ORAL

MEGACE

+ BRISTOL MYERS SQUIBB 40MG/ML

N020264 001 Sep 10, 1993

MEGESTROL ACETATE

APOTEX INC 40MG/ML

A077404 001 Feb 16, 2006

TABLET;ORAL

MEGACE

+ BRISTOL MYERS SQUIBB 20MG \*\*

N016979 001

+ 40MG \*\*

N016979 002

MEGESTROL ACETATE

TEVA 40MG

A074745 001 Feb 27, 1998

USL PHARMA 20MG

A070646 001 Oct 02, 1987

40MG

A070647 001 Oct 02, 1987

MELOXICAM

SUSPENSION;ORAL

MOBIC

+ BOEHRINGER INGELHEIM 7.5MG/5ML

N021530 001 Jun 01, 2004

TABLET;ORAL

MELOXICAM

ANDA REPOSITORY 7.5MG

A077935 001 Jul 19, 2006

15MG

A077935 002 Jul 19, 2006

CR DOUBLE CRANE 7.5MG

A078039 001 Dec 14, 2006

15MG

A078039 002 Dec 14, 2006

IMPAX LABS INC 7.5MG

A077930 001 Jul 19, 2006

15MG

A077930 002 Jul 19, 2006

MYLAN 7.5MG

A077934 001 Jul 20, 2006

15MG

A077934 002 Jul 20, 2006

ROXANE 7.5MG

A077925 001 Jul 19, 2006

15MG

A077925 002 Jul 19, 2006

SUN PHARM INDs INC 7.5MG

A077937 001 Jul 19, 2006

15MG

A077937 002 Jul 19, 2006

YABAO PHARM 7.5MG

A077933 001 Jul 19, 2006

15MG

A077933 002 Jul 19, 2006

MELPHALAN HYDROCHLORIDE

INJECTABLE;INJECTION

ALKERAN

+ APOTEX INC EQ 50MG BASE/VIAL

N020207 001 Nov 18, 1992

MELPHALAN HYDROCHLORIDE

MYLAN INSTITUTIONAL EQ 50MG BASE/VIAL

A090299 001 Oct 27, 2009

MEMANTINE HYDROCHLORIDE

TABLET;ORAL

MEMANTINE HYDROCHLORIDE

ORCHID HLTHCARE 5MG

A090044 001 Mar 12, 2012

10MG

A090044 002 Mar 12, 2012

**DISCONTINUED DRUG PRODUCT LIST**

6-236(of 375)

\*\* See List Footnote

MENADIOL SODIUM DIPHOSPHATE

INJECTABLE; INJECTION

KAPPADIONE

LILLY	10MG/ML	N005725 001
SYNKAYVITE		
ROCHE	5MG/ML	N003718 004
	10MG/ML	N003718 006
	37.5MG/ML	N003718 008

TABLET; ORAL

SYNKAYVITE

ROCHE	5MG	N003718 010
-------	-----	-------------

MENADIONE

TABLET; ORAL

MENADIONE

LILLY	5MG	N002139 003
-------	-----	-------------

MENOTROPINS (FSH; LH)

INJECTABLE; INJECTION

HUMEGON

ORGANON USA INC	75 IU/VIAL; 75 IU/VIAL	N020328 001 Sep 01, 1994
	150 IU/VIAL; 150 IU/VIAL	N020328 002 Sep 01, 1994

MENOTROPINS

FERRING	75 IU/VIAL; 75 IU/VIAL	A073598 001 Jan 30, 1997
	150 IU/VIAL; 150 IU/VIAL	A073599 001 Jan 30, 1997

PERGONAL

SERONO	75 IU/AMP; 75 IU/AMP	N017646 001
	150 IU/AMP; 150 IU/AMP	N017646 002 May 20, 1985

REPRONEX

FERRING	150 IU/VIAL; 150 IU/VIAL	N021047 002 Aug 27, 1999
---------	--------------------------	--------------------------

INJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS

REPRONEX

FERRING	75 IU/VIAL; 75 IU/VIAL	N021047 001 Aug 27, 1999
---------	------------------------	--------------------------

MEPENZOLATE BROMIDE

SOLUTION; ORAL

CANTIL

SANOFI AVENTIS US	25MG/5ML	N010679 004
-------------------	----------	-------------

TABLET; ORAL

CANTIL

+ SANOFI AVENTIS US	25MG	N010679 003
---------------------	------	-------------

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

US PHARM HOLDINGS	25MG/ML	N005010 007
	50MG/ML	N005010 002
	75MG/ML	N005010 009
	100MG/ML	N005010 003

MEPERIDINE HYDROCHLORIDE

ABBOTT	25MG/ML	A080388 001
	50MG/ML	A080385 001
	50MG/ML	A080387 001
	75MG/ML	A080389 001
	100MG/ML	A080386 001

BAXTER HLTHCARE	25MG/ML	A088279 001 Jun 15, 1984
	50MG/ML	A088280 001 Jun 15, 1984
	75MG/ML	A088281 001 Jun 15, 1984
	100MG/ML	A088282 001 Jun 15, 1984

IGI LABS INC	25MG/ML	A089781 001 Mar 31, 1989
	50MG/ML	A089782 001 Mar 31, 1989
	50MG/ML	A089783 001 Mar 31, 1989
	75MG/ML	A089784 001 Mar 31, 1989

	100MG/ML	A089785 001 Mar 31, 1989
	100MG/ML	A089786 001 Mar 31, 1989
	100MG/ML	A089787 001 Mar 31, 1989
	100MG/ML	A089788 001 Mar 31, 1989

INTL MEDICATION	10MG/ML	A086332 001
PARKE DAVIS	50MG/ML	A080364 002
	75MG/ML	A080364 003
	100MG/ML	A080364 001

**DISCONTINUED DRUG PRODUCT LIST**

6-237(of 375)

\*\* See List Footnote

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HYDROCHLORIDE

WATSON LABS	50MG/ML 100MG/ML	A073444 001 Mar 17, 1992 A073445 001 Mar 17, 1992
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE		
HOSPIRA	10MG/ML	A040305 001 Mar 10, 1999
INTL MEDICATION	10MG/ML	A081309 001 Aug 30, 1993
SPECGX LLC	10MG/ML	A040163 001 May 12, 1997
WATSON LABS	10MG/ML	A073443 001 Mar 17, 1992

SYRUP; ORAL

DEMEROL

US PHARM HOLDINGS 50MG/5ML \*\*

N005010 005

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

DURAMED PHARMS BARR	50MG 100MG	A040318 001 Oct 05, 1999 A040318 002 Oct 05, 1999
SUN PHARM INDUSTRIES	50MG 100MG	A080448 001 A080448 002
WATSON LABS	50MG 100MG	A040186 001 Jun 30, 1997 A040186 002 Jun 30, 1997
WYETH AYERST	50MG	A080454 001

MEPERIDINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERGAN

WEST-WARD PHARMS INT 25MG/ML; 25MG/ML

N011730 001

MEPHENTERMINE SULFATE

INJECTABLE; INJECTION

WYAMINE SULFATE

BAXTER HLTHCARE CORP EQ 15MG BASE/ML  
EQ 30MG BASE/MLN008248 002  
N008248 001MEPHENYTOIN

TABLET; ORAL

MESANTOIN

+ NOVARTIS 100MG \*\* N006008 001

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE

SOLVAY	3%	A084777 002 Apr 18, 1982
CARBOCAINE		
+ EASTMAN KODAK	3% **	N012125 003
MEPIVACAINE HYDROCHLORIDE		
BELMORA LLC	3%	A083559 001
INTL MEDICATION SYS	1%	A087509 001 Oct 05, 1982
WATSON LABS	1%	A088769 001 Nov 20, 1984
	2%	A088770 001 Nov 20, 1984
POLOCAINE		
DENTSPLY PHARM	3%	A088653 001 Aug 21, 1984

MEPREDNISONE

TABLET; ORAL

BETAPAR

SCHERING 4MG N016053 002

MEPROBAMATE

CAPSULE; ORAL

EQUANIL

WYETH AYERST	400MG	N012455 002
CAPSULE, EXTENDED RELEASE; ORAL		
MEPROSPAN		
MEDPOINTE PHARM HLC	200MG 400MG	N011284 001 N011284 002

TABLET; ORAL

AMOSENE

FERNDALE LABS 400MG A084030 001

BAMATE

ALRA	200MG 400MG	A080380 001 A080380 002
------	----------------	----------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-238(of 375)

\*\* See List Footnote

**MEPROBAMATE**

TABLET;ORAL

EQUANIL

WYETH AYERST	200MG	N010028 005
	400MG	N010028 004
MEPRIAM		
TEVA	400MG	N016069 001
MEPROBAMATE		
AUROLIFE PHARMA LLC	400MG	A080655 001
BARR	600MG	A084230 001
ELKINS SINK	200MG	N015426 002
	400MG	N015426 001
HEATHER	400MG	N016928 003
	600MG	A084329 001
IMPAX LABS	200MG	N014322 002
	400MG	N014322 001
IVAX SUB TEVA PHARMS	200MG	N015438 001
	400MG	N015438 002
	600MG	A084181 001
IVC INDS	400MG	A084153 001
LANNETT	200MG	N014882 002
	400MG	N014882 001
LEDERLE	400MG	A086299 001
LEE KM	400MG	A089538 001 Nov 25, 1987
MALLARD	400MG	N015072 002
MK LABS	200MG	N014368 004
	400MG	N014368 002
MYLAN	400MG	A083618 001
NEXGEN PHARMA INC	200MG	A084220 001
	400MG	A084589 001
PARKE DAVIS	200MG	A084744 001
	400MG	A084744 002
PERRIGO	200MG	A084546 001
	400MG	A084547 001
PHARMAVITE	400MG	A084438 001
PUREPAC PHARM	200MG	A084804 001
	400MG	A084804 002
PVT FORM	400MG	N014601 001
ROXANE	600MG	A084332 001
SANDOZ	200MG	N014547 002
	400MG	N014547 001
SCHERER LABS	400MG	A083343 001
SOLVAY	200MG	A084435 001
STANLABS PHARM	200MG	N014474 002
	400MG	N014474 004
SUN PHARM INDUSTRIES	200MG	A080699 001
	400MG	A080699 002
TABLICAPS	400MG	A083494 001
TARO	200MG	A200998 001 May 23, 2011
	400MG	A200998 002 May 23, 2011
USL PHARMA	200MG	A087825 001 Mar 18, 1982
	400MG	A087826 001 Mar 18, 1982
VALEANT PHARM INTL	200MG	N015139 006
	400MG	N015139 005
VANGARD	400MG	A088011 001 Jul 14, 1982
WATSON LABS	200MG	A085720 001
	400MG	A085721 001
	600MG	A084274 001
	600MG	A085719 001
WEST WARD	200MG	N015417 003
	400MG	N015417 002
WHITEWORTH TOWN PLSN	200MG	A083830 001
	400MG	A083442 001
MILTOWN		
+ MEDPOINTE PHARM HLC	200MG **	N009698 004
+	400MG **	N009698 002
	600MG	A083919 001
NEURAMATE		
HALSEY	200MG	N014359 002

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-239(of 375)

\*\* See List Footnote

**MEPROBAMATE**

TABLET;ORAL NEURAMATE	400MG	N014359 001
TRANMEP SOLVAY	400MG 400MG	A084369 001 N016249 001

**MEQUINOL; TRETINOIN**

SOLUTION;TOPICAL SOLAGE	2%;0.01%	N020922 001 Dec 10, 1999
AQUA PHARMS		

**MEROPENEM**

INJECTABLE;INJECTION MEROPENEM		
SANDOZ	500MG/VIAL 1GM/VIAL	A091201 001 Mar 29, 2011 A091201 002 Mar 29, 2011

**MERSALYL SODIUM; THEOPHYLLINE**

INJECTABLE;INJECTION MERSALYL-THEOPHYLLINE		
WATSON LABS	100MG/ML;50MG/ML	A084875 001

**MESALAMINE**

SUPPOSITORY;RECTAL CANASA		
FOREST LABS LLC	500MG	N021252 001 Jan 05, 2001
ROWASA + MEDA PHARMS	500MG **	N019919 001 Dec 18, 1990
TABLET, DELAYED RELEASE;ORAL ASACOL		
APIL	400MG	N019651 001 Jan 31, 1992

**MESNA**

INJECTABLE;INTRAVENOUS MESNA		
MYLAN LABS LTD	100MG/ML	A203364 001 Jul 18, 2014

**MESORIDAZINE BESYLATE**

CONCENTRATE;ORAL SERENTIL		
NOVARTIS	EQ 25MG BASE/ML	N016997 001
INJECTABLE;INJECTION SERENTIL		
NOVARTIS	EQ 25MG BASE/ML	N016775 001
TABLET;ORAL SERENTIL		
NOVARTIS	EQ 10MG BASE EQ 25MG BASE EQ 50MG BASE EQ 100MG BASE	N016774 001 N016774 002 N016774 003 N016774 004

**MESTRANOL; NORETHINDRONE**

TABLET;ORAL-20 NORINYL		
ACTAVIS LABS UT INC	0.1MG;2MG	N013625 004
TABLET;ORAL-21 NORETHIN 1/50M-21		
WATSON LABS	0.05MG;1MG	A071539 001 Apr 12, 1988
NORETHINDRONE AND MESTRANOL WATSON LABS	0.05MG;1MG	A070758 001 Jul 01, 1988
NORINYL 1+50 21-DAY ACTAVIS LABS UT INC	0.05MG;1MG	N013625 002
NORINYL 1+80 21-DAY GD SEARLE LLC	0.08MG;1MG	N016724 001
ORTHO-NOVUM 1/50 21 ORTHO MCNEIL PHARM	0.05MG;1MG	N012728 004
ORTHO-NOVUM 1/80 21 ORTHO MCNEIL PHARM	0.08MG;1MG	N016715 001
ORTHO-NOVUM 10-21 ORTHO MCNEIL PHARM	0.06MG;10MG	N012728 001

**DISCONTINUED DRUG PRODUCT LIST**

6-240(of 375)

\*\* See List Footnote

MESTRANOL; NORETHINDRONE

TABLET;ORAL-21			
ORTHO-NOVUM 2-21			
ORTHO MCNEIL PHARM	0.1MG;2MG		N012728 005
TABLET;ORAL-28			
NORETHIN 1/50M-28			
WATSON LABS	0.05MG;1MG		A071540 001 Apr 12, 1988
NORETHINDRONE AND MESTRANOL			
WATSON LABS	0.05MG;1MG		A070759 001 Jul 01, 1988
NORINYL 1+80 28-DAY			
GD SEARLE LLC	0.08MG;1MG		N016725 001
ORTHO-NOVUM 1/50 28			
ORTHO MCNEIL JANSSEN	0.05MG;1MG		N016709 001
ORTHO-NOVUM 1/80 28			
ORTHO MCNEIL PHARM	0.08MG;1MG		N016715 002

MESTRANOL; NORETHYNODREL

TABLET;ORAL			
ENOVID			
GD SEARLE LLC	0.075MG;5MG		N010976 008
	0.15MG;9.85MG		N010976 005
TABLET;ORAL-20			
ENOVID			
GD SEARLE LLC	0.075MG;5MG		N010976 004
ENOVID-E			
GD SEARLE LLC	0.1MG;2.5MG		N010976 006
TABLET;ORAL-21			
ENOVID-E 21			
GD SEARLE LLC	0.1MG;2.5MG		N010976 007

METAPROTERENOL SULFATE

AEROSOL, METERED;INHALATION			
ALUPENT			
BOEHRINGER INGELHEIM	0.65MG/INH		N016402 001
SOLUTION;INHALATION			
ALUPENT			
BOEHRINGER INGELHEIM	0.4%		N018761 002 Oct 10, 1986
	0.6%		N018761 001 Jun 30, 1983
	5%		N017659 001
METAPROTERENOL SULFATE			
APOTEX INC	0.4%		A075402 001 Feb 28, 2001
	0.6%		A075403 001 Feb 28, 2001
ASTRAZENECA	0.4%		A071275 001 Jul 27, 1988
	0.6%		A071018 001 Jul 27, 1988
DEY	0.33%		A071806 001 Aug 05, 1988
	0.5%		A071805 001 Aug 05, 1988
	5%		A070805 001 Aug 17, 1987
MYLAN SPECIALITY LP	0.4%		A071786 001 Aug 05, 1988
	0.6%		A070804 001 Aug 17, 1987
NEPHRON	0.4%		A071855 001 Jul 14, 1988
	0.6%		A071726 001 Jul 14, 1988
WOCKHARDT	0.4%		A075586 001 May 30, 2002
	0.6%		A075586 002 May 30, 2002
	5%		A072190 001 Jun 07, 1988
PROMETA			
MURO	5%		A073340 001 Mar 30, 1992
SYRUP;ORAL			
ALUPENT			
BOEHRINGER INGELHEIM	10MG/5ML		N017571 001
METAPROTERENOL SULFATE			
APOTEX INC	10MG/5ML		A075235 001 Jan 27, 2000
G AND W LABS INC	10MG/5ML		A072761 001 Feb 27, 1992
	10MG/5ML		A073034 001 Aug 30, 1991
MORTON GROVE	10MG/5ML		A071656 001 Oct 13, 1987
WOCKHARDT	10MG/5ML		A074702 001 Mar 24, 1997
PROMETA			
MURO	10MG/5ML		A072023 001 Sep 15, 1988
TABLET;ORAL			
ALUPENT			
BOEHRINGER INGELHEIM	10MG		N015874 002

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-241(of 375)

\*\* See List Footnote

METAPROTERENOL SULFATETABLET;ORAL  
ALUPENT

	20MG	N015874 001
METAPROTERENOL SULFATE		
AM THERAP	10MG	A072054 001 Jun 23, 1988
	20MG	A072055 001 Jun 23, 1988
TEVA	10MG	A072519 001 Mar 30, 1990
	20MG	A072520 001 Mar 30, 1990
USL PHARMA	10MG	A071013 001 Jan 25, 1988
	20MG	A071014 001 Jan 25, 1988
WATSON LABS	10MG	A073013 001 Jan 31, 1991
	20MG	A072795 001 Jan 31, 1991

METARAMINOL BITARTRATEINJECTABLE;INJECTION  
ARAMINE

+ MERCK	EQ 10MG BASE/ML **	N009509 002 Dec 22, 1987
METARAMINOL BITARTRATE		
ABRAXIS PHARM	EQ 10MG BASE/ML	A080431 001
ELKINS SINK	EQ 10MG BASE/ML	A083363 001
GD SEARLE LLC	EQ 10MG BASE/ML	A086418 001
	EQ 20MG BASE/ML	A086418 002

METAXALONETABLET;ORAL  
METAXALONE

EPIC PHARMA LLC	640MG	N022503 001 Jun 01, 2015
SKELAXIN		
+ KING PHARMS	400MG **	N013217 001

METFORMIN HYDROCHLORIDETABLET;ORAL  
GLUCOPHAGE

+ BRISTOL MYERS SQUIBB	625MG **	N020357 003 Nov 05, 1998
+	750MG **	N020357 004 Nov 05, 1998

METFORMIN HYDROCHLORIDE		
BARR	500MG	A075971 001 Jan 25, 2002
	850MG	A075971 002 Jan 25, 2002
	1GM	A075971 003 Jan 25, 2002
IPCA LABS LTD	500MG	A078422 001 Aug 06, 2007
	850MG	A078422 002 Aug 06, 2007
	1GM	A078422 003 Aug 06, 2007
IVAX SUB TEVA PHARMS	500MG	A075975 001 Jan 24, 2002
	625MG	A075975 004 Jan 24, 2002
	750MG	A075975 005 Jan 24, 2002
	850MG	A075975 002 Jan 24, 2002
	1GM	A075975 003 Jan 24, 2002
MYLAN PHARMS INC	500MG	A075969 001 Jan 29, 2002
	850MG	A075969 002 Jan 29, 2002
	1GM	A075969 003 Jan 29, 2002
TEVA	500MG	A076328 001 Dec 16, 2002
	850MG	A076328 002 Dec 16, 2002
	1GM	A076328 003 Dec 16, 2002
WATSON LABS	500MG	A075979 001 Jan 24, 2002
	850MG	A075979 002 Jan 24, 2002
	1GM	A075979 003 Jan 24, 2002
WATSON LABS FLORIDA	500MG	A075961 001 Jan 25, 2002
	850MG	A075961 002 Jan 25, 2002
	1GM	A075961 003 Jan 25, 2002

TABLET, EXTENDED RELEASE;ORAL

METFORMIN HYDROCHLORIDE		
ACTAVIS ELIZABETH	500MG	A076450 001 Oct 01, 2004
	750MG	A076878 001 Apr 13, 2005
BARR	500MG	A076496 001 Nov 25, 2005
IMPAK LABS	500MG	A076249 001 Jul 30, 2004
	750MG	A076985 001 Sep 13, 2005
IVAX SUB TEVA PHARMS	500MG	A076545 001 Dec 01, 2003
MYLAN	500MG	A076650 001 Sep 13, 2005
	750MG	A077113 001 Sep 08, 2005
RANBAXY LABS LTD	500MG	A076413 001 Jun 18, 2004

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-242(of 375)

\*\* See List Footnote

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL  
METFORMIN HYDROCHLORIDE

SANDOZ	750MG	A077211 001 Jun 29, 2005
SUN PHARM INDUSTRIES	500MG	A076223 001 Dec 14, 2004
TORRENT PHARMS LTD	500MG	A077124 001 Dec 21, 2005
WATSON LABS INC	750MG	A079226 001 Feb 18, 2010
	500MG	A076818 001 Dec 14, 2004

METFORMIN HYDROCHLORIDE; REPAGLINIDE

TABLET; ORAL  
PRANDIMET  
+ NOVO NORDISK INC 500MG; 1MG  
+ 500MG; 2MG

N022386 001	Jun 23, 2008
N022386 002	Jun 23, 2008

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL  
AVANDAMET  
+ SB PHARMCO 500MG; EQ 1MG BASE \*\*  
+ 500MG; EQ 2MG BASE \*\*  
+ 500MG; EQ 4MG BASE \*\*  
+ 1GM; EQ 2MG BASE \*\*  
+ 1GM; EQ 4MG BASE \*\*

N021410 001	Oct 10, 2002
N021410 002	Oct 10, 2002
N021410 003	Oct 10, 2002
N021410 004	Aug 25, 2003
N021410 005	Aug 25, 2003

METHACHOLINE CHLORIDE

FOR SOLUTION; INHALATION  
PROVOCHOLINE  
+ METHAPHARM 1600MG/VIAL

N019193 002 Aug 29, 2016

METHACYCLINE HYDROCHLORIDE

CAPSULE; ORAL  
RONDOMYCIN  
MEDPOINTE PHARM HLC EQ 140MG BASE  
EQ 280MG BASE

A060641 001
A060641 002

SYRUP; ORAL  
RONDOMYCIN  
MEDPOINTE PHARM HLC EQ 70MG BASE/5ML

A060641 003

METHADONE HYDROCHLORIDE

POWDER; FOR RX COMPOUNDING  
METHADONE HYDROCHLORIDE  
MALLINCKRODT INC 50GM/BOT  
100GM/BOT  
500GM/BOT

N006383 002
N006383 003
N006383 004

SYRUP; ORAL  
DOLOPHINE HYDROCHLORIDE  
WEST-WARD PHARMS INT 10MG/30ML

N006134 004

TABLET; ORAL  
METHADONE HYDROCHLORIDE

ROXANE 5MG	A088108 001 Mar 08, 1983
10MG	A088109 001 Mar 08, 1983
40MG	A074081 001 Apr 28, 1995
SANDOZ 5MG	A040241 001 May 29, 1998

TABLET, DISPERSIBLE; ORAL  
WESTADONE

SANDOZ 2.5MG	N017108 001
--------------	-------------

TABLET, EFFERVESCENT; ORAL  
WESTADONE

SANDOZ 5MG	N017108 002
10MG	N017108 003
40MG	N017108 004

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL  
METHAMPEX  
TEVA 10MG

A083889 001

METHAMPHETAMINE HYDROCHLORIDE 5MG	A040529 001 Feb 25, 2004
REXAR 5MG	A084931 001
ABLE 10MG	A084931 002
TEVA 5MG	A086359 001

**DISCONTINUED DRUG PRODUCT LIST**

6-243(of 375)

\*\* See List Footnote

METHAMPHETAMINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL

DESOXYN

RECORDATI RARE	5MG	N005378 004
	10MG	N005378 003
	15MG	N005378 005

METHANTHELINE BROMIDE

TABLET;ORAL

BANTHINE

SHIRE	50MG	N007390 001
-------	------	-------------

METHARBITAL

TABLET;ORAL

GEMONIL

ABBVIE	100MG	N008322 001
--------	-------	-------------

METHAZOLAMIDE

TABLET;ORAL

METHAZOLAMIDE

APPLIED ANAL	25MG	A040011 001 Jul 17, 1997
	50MG	A040011 002 Jul 17, 1997
SANDOZ	25MG	A040102 001 Aug 28, 1996
	50MG	A040102 002 Aug 28, 1996
NEPTAZANE		
+ LEDERLE	25MG **	N011721 002 Nov 25, 1991
+	50MG **	N011721 001

METHDILAZINE

TABLET, CHEWABLE;ORAL

TACARYL

WESTWOOD SQUIBB	3.6MG	N011950 009
-----------------	-------	-------------

METHDILAZINE HYDROCHLORIDE

SYRUP;ORAL

METHDILAZINE HYDROCHLORIDE

ALPHARMA US PHARMS	4MG/5ML	A087122 001
TACARYL		
WESTWOOD SQUIBB	4MG/5ML	N011950 007
TABLET;ORAL		
TACARYL		
WESTWOOD SQUIBB	8MG	N011950 006

METHICILLIN SODIUM

INJECTABLE; INJECTION

STAPHCILLIN

APOTHECON	EQ 900MG BASE/VIAL	A061449 001
	EQ 900MG BASE/VIAL	N050117 001
	EQ 3.6GM BASE/VIAL	A061449 002
	EQ 3.6GM BASE/VIAL	N050117 002
	EQ 5.4GM BASE/VIAL	A061449 003
	EQ 5.4GM BASE/VIAL	N050117 003

METHIMAZOLE

TABLET;ORAL

METHIMAZOLE

ECI PHARMS LLC	15MG	A040619 003 Jul 12, 2005
	20MG	A040547 004 Feb 18, 2005
MYLAN	20MG	A040350 003 Jun 07, 2001
TAPAZOLE		
+ KING PHARMS	5MG **	N007517 002
+	10MG **	N007517 004

METHIXENE HYDROCHLORIDE

TABLET;ORAL

TREST

NOVARTIS	1MG	N013420 001
----------	-----	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-244(of 375)

\*\* See List Footnote

METHOCARBAMOL

INJECTABLE; INJECTION

METHOCARBAMOL

MARSAM PHARMS LLC	100MG/ML	A089849 001 Dec 27, 1991
WATSON LABS	100MG/ML	A086459 001
<b>TABLET; ORAL</b>		
<b>DELAXIN</b>		
FERNDALE LABS	500MG	A085454 001
<b>FORBAXIN</b>		
FOREST LABS	750MG	A085136 001
<b>METHOCARBAMOL</b>		
ABLE	500MG	A040413 001 Mar 17, 2003
	750MG	A040413 002 Mar 17, 2003
AM THERAP	500MG	A089417 001 Feb 11, 1987
	750MG	A089418 001 Feb 11, 1987
ASCOT	500MG	A087660 001 Oct 27, 1982
	750MG	A087661 001 Oct 27, 1982
CLONMEL HLTHCARE	500MG	A085961 001
	750MG	A085963 001
HEATHER	500MG	A084675 001
	750MG	A084924 001
IMPAX LABS	500MG	A084927 001
	750MG	A084928 001
INWOOD LABS	500MG	A085137 001
IVAX SUB TEVA PHARMS	500MG	A084648 001
	750MG	A084649 001
KV PHARM	500MG	A085660 001
	750MG	A085658 001
LANNETT HOLDINGS INC	500MG	A084756 002 Mar 31, 2003
	750MG	A084756 001
MYLAN	500MG	A084259 001
	750MG	A084323 001
NYLOS	750MG	A085033 001
PIONEER PHARMS	500MG	A088731 001 Dec 13, 1985
	750MG	A089082 001 Dec 13, 1985
PURACAP PHARM	500MG	A084231 002
	750MG	A084471 001
PUREPAC PHARM	500MG	A085718 001
	750MG	A085718 002
ROXANE	500MG	A088646 001 Feb 29, 1984
	750MG	A088647 001 Feb 29, 1984
SANDOZ	500MG	A084616 001
	500MG	A087283 001
	750MG	A084615 001
	750MG	A087282 001
SOLVAY	500MG	A084448 001
	750MG	A084449 001
SUN PHARM INDUSTRIES	500MG	A084488 001
	750MG	A084486 001
SUPERPHARM	500MG	A087589 001 Jan 22, 1982
	750MG	A087590 001 Jan 22, 1982
TABLICAPS	500MG	A084846 001
UPSHER SMITH	500MG	A087453 001
	750MG	A087454 001
WATSON LABS	500MG	A083605 001
	500MG	A085180 001
	750MG	A083605 002
	750MG	A085192 001

METHOHEXITAL SODIUM

INJECTABLE; INJECTION

BREVITAL SODIUM

PAR STERILE PRODUCTS	200MG/VIAL	N011559 004 Dec 21, 2012
	5GM/VIAL	N011559 003

**DISCONTINUED DRUG PRODUCT LIST**

6-245(of 375)

\*\* See List Footnote

**METHOTREXATE**

SOLUTION;SUBCUTANEOUS

OTREXUP

+ ANTARES PHARMA INC 7.5MG/0.4ML (7.5MG/0.4ML) N204824 005 Nov 07, 2014

OTREXUP PFS

+ ANTARES PHARMA INC	10MG/0.4ML (10MG/0.4ML)	N204824 009 May 31, 2017
+	15MG/0.6ML (15MG/0.6ML)	N204824 010 May 31, 2017
+	17.5MG/0.7ML (17.5MG/0.7ML)	N204824 011 May 31, 2017
+	20MG/0.8ML (20MG/0.8ML)	N204824 012 May 31, 2017
+	22.5MG/0.9ML (22.5MG/0.9ML)	N204824 013 May 31, 2017
+	25MG/ML (25MG/ML)	N204824 014 May 31, 2017

RASUVO

+ MEDAC PHARMA INC 27.5MG/0.55ML (27.5MG/0.55ML) N205776 009 Jul 10, 2014

**METHOTREXATE SODIUM**

INJECTABLE;INJECTION

ABITREXATE

ABIC	EQ 25MG BASE/ML	A089161 001 Mar 10, 1987
	EQ 50MG BASE/VIAL	A089354 001 Jul 17, 1987
	EQ 100MG BASE/VIAL	A089355 001 Jul 17, 1987
	EQ 250MG BASE/VIAL	A089356 001 Jul 17, 1987

FOLEX

PHARMACIA AND UPJOHN	EQ 25MG BASE/VIAL	A087695 001 Apr 08, 1983
	EQ 50MG BASE/VIAL	A087695 002 Apr 08, 1983
	EQ 100MG BASE/VIAL	A087695 003 Apr 08, 1983
	EQ 250MG BASE/VIAL	A088954 001 Oct 24, 1985

FOLEX PFS

PHARMACIA AND UPJOHN	EQ 25MG BASE/ML	A081242 001 Aug 23, 1991
	EQ 25MG BASE/ML	A089180 001 Jan 03, 1986

METHOTREXATE LPF

HOSPIRA	EQ 25MG BASE/ML	N011719 007 Mar 31, 1982
---------	-----------------	--------------------------

METHOTREXATE PRESERVATIVE FREE

HOSPIRA	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N011719 014 Apr 13, 2005
+	EQ 500MG BASE/20ML (EQ 25MG BASE/ML) **	N011719 013 Apr 13, 2005
	EQ 2.5GM BASE/100ML (EQ 25MG BASE/ML)	N011719 011 Apr 13, 2005

METHOTREXATE SODIUM

ABRAXIS PHARM	EQ 2.5MG BASE/ML	A089323 001 Jun 13, 1986
	EQ 20MG BASE/VIAL	A088935 001 Oct 11, 1985
	EQ 25MG BASE/ML	A089263 001 Jun 13, 1986
	EQ 25MG BASE/ML	A089322 001 Jun 13, 1986
	EQ 50MG BASE/VIAL	A088936 001 Oct 11, 1985
	EQ 100MG BASE/VIAL	A088937 001 Oct 11, 1985

HOSPIRA	EQ 2.5MG BASE/ML	N011719 004
	EQ 20MG BASE/VIAL	N011719 001
	EQ 25MG BASE/ML	N011719 005
	EQ 50MG BASE/VIAL	N011719 003
	EQ 100MG BASE/VIAL	N011719 006

NORBROOK	EQ 25MG BASE/ML	A088648 001 May 09, 1986
PHARMACHEMIE USA	EQ 25MG BASE/ML	A089158 001 Jul 08, 1988

METHOTREXATE SODIUM PRESERVATIVE FREE

HOSPIRA	EQ 1GM BASE/VIAL	N011719 009 Apr 07, 1988
---------	------------------	--------------------------

MEXATE

BRISTOL	EQ 20MG BASE/VIAL	A086358 001
	EQ 50MG BASE/VIAL	A086358 002
	EQ 100MG BASE/VIAL	A086358 003
	EQ 250MG BASE/VIAL	A086358 004

MEXATE-AQ

BRISTOL MYERS	EQ 25MG BASE/ML	A088760 001 Feb 14, 1985
---------------	-----------------	--------------------------

MEXATE-AQ PRESERVED

BRISTOL MYERS SQUIBB	EQ 25MG BASE/ML	A089887 001 Apr 14, 1989
----------------------	-----------------	--------------------------

TABLET;ORAL

METHOTREXATE SODIUM		
DURAMED PHARMS BARR	EQ 2.5MG BASE	A040233 001 Jun 17, 1999

**DISCONTINUED DRUG PRODUCT LIST**

6-246(of 375)

\*\* See List Footnote

METHOXAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

VASOXYL

GLAXOSMITHKLINE

10MG/ML

N006772 002

20MG/ML

N006772 001

METHOXALEN

CAPSULE; ORAL

8-MOP

+ VALEANT PHARM INTL 10MG

N009048 001

METHOXALEN

IDT AUSTRALIA LTD

10MG

A087781 001 Jun 08, 1982

LOTION; TOPICAL

OXSORALEN

+ VALEANT PHARM INTL 1%

N009048 002

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

PVT FORM

2.5MG

A080970 001

PAMINE

FOUGERA PHARMS

2.5MG \*\*

N008848 001

PAMINE FORTE

FOUGERA PHARMS

5MG \*\*

N008848 002 Mar 25, 2003

METHYCLOTHIAZIDE

TABLET; ORAL

AQUATENSEN

MEDPOINTE PHARM HLC

5MG

N017364 001

ENDURON

+ ABBVIE

2.5MG \*\*

N012524 001

+

5MG \*\*

N012524 004

METHYCLOTHIAZIDE

FOSUN PHARMA

2.5MG

A089835 001 Aug 18, 1988

5MG

A089837 001 Aug 18, 1988

IVAX PHARMS

2.5MG

A087913 001 Jun 03, 1982

5MG

A087786 001 May 18, 1982

MYLAN

2.5MG

A087671 001 Aug 17, 1982

PAR PHARM

2.5MG

A089135 001 Feb 12, 1986

5MG

A089136 001 Feb 12, 1986

USL PHARMA

5MG

A088745 001 Mar 21, 1985

WATSON LABS

2.5MG

A085487 001 Mar 11, 1982

2.5MG

A088750 001 Sep 06, 1984

5MG

A085476 001 Mar 11, 1982

5MG

A088724 001 Sep 06, 1984

METHYCLOTHIAZIDE; PARGYLINE HYDROCHLORIDE

TABLET; ORAL

EUTRON

ABBOTT

5MG; 25MG

N016047 001

METHYCLOTHIAZIDE; RESERPINE

TABLET; ORAL

DIUTENSEN-R

MEDPOINTE PHARM HLC 2.5MG; 0.1MG

N012708 005

METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL

METVIXIA

GALDERMA LABS LP

EQ 16.8% BASE

N021415 001 Jul 27, 2004

METHYLDOPA

SUSPENSION; ORAL

ALDOMET

MERCK

250MG/5ML

N018389 001

TABLET; ORAL

ALDOMET

+ MERCK

125MG \*\*

N013400 003

+

250MG \*\*

N013400 001

+

500MG \*\*

N013400 002

METHYLDOPA

ACCORD HLTHCARE

125MG

A070070 003 Oct 15, 1985

DURAMED PHARMS BARR

250MG

A071006 001 Dec 16, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-247(of 375)

\*\* See List Footnote

METHYLDOPATABLET;ORAL  
METHYLDOPA

FOSUN PHARMA	500MG	A071009 001	Dec 16, 1986
	250MG	N018934 001	Jun 29, 1984
	500MG	N018934 002	Jun 29, 1984
HALSEY	125MG	A071751 001	Mar 28, 1988
	250MG	A071752 001	Mar 28, 1988
	500MG	A071753 001	Mar 28, 1988
PAR PHARM	125MG	A070535 001	Jan 02, 1987
	250MG	A070536 001	Jan 02, 1987
	500MG	A070537 001	Jan 02, 1987
PARKE DAVIS	125MG	A070331 001	Apr 15, 1986
	250MG	A070332 001	Apr 15, 1986
	500MG	A070333 001	Apr 15, 1986
PLIVA	125MG	A072126 001	Jul 07, 1988
	250MG	A072127 001	Jul 07, 1988
	500MG	A072128 001	Jul 07, 1988
PUREPAC PHARM	125MG	A070749 001	Feb 07, 1986
	250MG	A070750 001	Feb 07, 1986
	500MG	A070452 001	Feb 07, 1986
ROXANE	125MG	A070192 001	Apr 25, 1986
	250MG	A070193 001	Apr 25, 1986
	500MG	A070194 001	Apr 25, 1986
SANDOZ	125MG	A071700 001	Mar 02, 1988
SUN PHARM INDUSTRIES	125MG	A070073 001	Oct 09, 1986
	250MG	A070060 001	Oct 09, 1986
	500MG	A070074 001	Oct 09, 1986
SUPERPHARM	250MG	A070669 001	Jun 23, 1989
	500MG	A070670 001	Jun 23, 1989
TEVA	125MG	A071105 001	Dec 05, 1986
	250MG	A071106 001	Dec 05, 1986
	500MG	A071067 001	Dec 05, 1986
WATSON LABS	125MG	A070245 001	Feb 25, 1986
	125MG	A070260 001	Jun 24, 1985
	250MG	A070246 001	Feb 25, 1986
	250MG	A070261 001	Jun 24, 1985
	250MG	A070703 001	Jun 06, 1986
	500MG	A070247 001	Feb 25, 1986
	500MG	A070262 001	Jun 24, 1985

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

ALDOMET

+ MERCK	50MG/ML **	N013401 001
METHYLDOPATE HYDROCHLORIDE		
ABRAXIS PHARM	50MG/ML	A070652 001
BAXTER HLTHCARE	50MG/ML	A070291 001
HOSPIRA	50MG/ML	A070691 001
	50MG/ML	A070698 001
	50MG/ML	A070699 001
	50MG/ML	A070849 001
MARSAM PHARMS LLC	50MG/ML	A071812 001
SMITH AND NEPHEW	50MG/ML	A070841 001
TEVA PARENTERAL	50MG/ML	A072974 001

METHYLERGONOVINE MALEATE

TABLET;ORAL

METHERGINE

+ EDISON THERAPS LLC	0.2MG **	N006035 003
----------------------	----------	-------------

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

RITALIN LA

+ NOVARTIS	60MG **	N021284 005	Oct 27, 2014
------------	---------	-------------	--------------

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

ABLE	5MG	A040404 001	Mar 29, 2001
	10MG	A040404 002	Mar 29, 2001
	20MG	A040404 003	Mar 29, 2001
ACTAVIS ELIZABETH	5MG	A040321 001	Feb 05, 2002

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-248(of 375)

\*\* See List Footnote

METHYLPHENIDATE HYDROCHLORIDE

TABLET;ORAL

METHYLPHENIDATE HYDROCHLORIDE

10MG	A040321	002	Feb 05,	2002
20MG	A040321	003	Feb 05,	2002

TABLET, CHEWABLE;ORAL

METHYLINE

+ SPECGX LLC	2.5MG **	N021475	001	Apr 15,	2003
+	5MG **	N021475	002	Apr 15,	2003
+	10MG **	N021475	003	Apr 15,	2003

TABLET, EXTENDED RELEASE;ORAL

METADATE ER

UCB INC	10MG	A040306	001	Oct 20,	1999
---------	------	---------	-----	---------	------

METHYLPHENIDATE HYDROCHLORIDE

ABLE	20MG	A076032	001	May 09,	2001
ACTAVIS ELIZABETH	20MG	A075450	001	Dec 21,	2001
WATSON LABS	20MG	A040410	001	Feb 09,	2001

METHYLPREDNISOLONE

TABLET;ORAL

MEDROL

PHARMACIA AND UPJOHN	24MG	N011153	005
----------------------	------	---------	-----

METHYLPREDNISOLONE

HEATHER	4MG	A085650	001
PAR PHARM	16MG	A089207	001
	24MG	A089208	001
	32MG	A089209	001
SANDOZ	4MG	A087341	001
WATSON LABS	4MG	A086161	001
	16MG	A086159	001

METHYLPREDNISOLONE ACETATE

ENEMA;RECTAL

MEDROL

PHARMACIA AND UPJOHN	40MG/BOT	N018102	001
----------------------	----------	---------	-----

INJECTABLE;INJECTION

M-PREDROL

BEL MAR	40MG/ML	A086666	001
	80MG/ML	A087135	001

METHYLPREDNISOLONE ACETATE

AKORN	40MG/ML	A086903	001	Oct 20,	1982
	80MG/ML	A086903	002	Oct 20,	1982
WATSON LABS	20MG/ML	A085597	001		
	20MG/ML	A087248	001		
	40MG/ML	A085374	001		
	40MG/ML	A085600	001		
	80MG/ML	A085595	001		
	80MG/ML	A086507	001		

OINTMENT;TOPICAL

MEDROL ACETATE

PHARMACIA AND UPJOHN	0.25%	N012421	001
	1%	N012421	002

METHYLPREDNISOLONE ACETATE; NEOMYCIN SULFATE

CREAM;TOPICAL

NEO-MEDROL ACETATE

PHARMACIA AND UPJOHN	0.25%;EQ 3.5MG BASE/GM	A060611	002
	1%;EQ 3.5MG BASE/GM	A060611	001

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE;INJECTION

A-METHAPRED

ABBOTT	EQ 40MG BASE/VIAL	A089573	001	Feb 22,	1991
	EQ 125MG BASE/VIAL	A089574	001	Feb 22,	1991
	EQ 500MG BASE/VIAL	A089575	001	Feb 22,	1991
	EQ 1GM BASE/VIAL	A089576	001	Feb 22,	1991
HOSPIRA	EQ 40MG BASE/VIAL	A085853	001		
	EQ 125MG BASE/VIAL	A085855	001		
	EQ 500MG BASE/VIAL	A085854	001		
	EQ 500MG BASE/VIAL	A089173	001	Aug 18,	1987
	EQ 1GM BASE/VIAL	A085852	001		

**DISCONTINUED DRUG PRODUCT LIST**

6-249(of 375)

\*\* See List Footnote

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-METHAPRED

HOSPIRA INC	EQ 1GM BASE/VIAL EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL	A089174 001 Aug 18, 1987 A040793 001 Nov 25, 2008 A040827 001 Nov 25, 2008
METHYLPREDNISOLONE ELKINS SINK	EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A086906 002 A086906 003 A086906 004
ORGANON USA INC	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A087535 001 Jun 25, 1982 A087535 002 Jun 25, 1982
METHYLPREDNISOLONE SODIUM SUCCINATE ABRAXIS PHARM	EQ 40MG BASE/VIAL EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 500MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL	A088676 001 Jun 08, 1984 A089143 001 Mar 28, 1986 A088677 001 Jun 08, 1984 A089144 001 Mar 28, 1986 A088678 001 Jun 08, 1984 A089186 001 Mar 28, 1986 A089187 001 Mar 28, 1986 A088679 001 Jun 08, 1984 A089188 001 Mar 28, 1986 A089189 001 Mar 28, 1986
BEDFORD LABS	EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 1GM BASE/VIAL	A040662 001 Feb 21, 2007 A040641 002 Feb 21, 2007 A040641 003 Feb 21, 2007 A040709 001 Feb 21, 2007 A040641 004 Feb 21, 2007 A040709 002 Feb 21, 2007
ELKINS SINK INTL MEDICATION	EQ 40MG BASE/VIAL EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A086906 001 A087812 001 Feb 09, 1983 A087813 001 Feb 09, 1983 A087851 001 Feb 09, 1983 A087852 001 Feb 09, 1983
TEVA PARENTERAL	EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A081266 001 Nov 30, 1992 A081267 001 Nov 30, 1992 A081268 001 Nov 30, 1992
WATSON LABS	EQ 40MG BASE/VIAL EQ 125MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	A086953 001 Jul 22, 1982 A087030 001 Jul 22, 1982 A088523 001 Jul 24, 1984 A088524 001 Jul 24, 1984

METHYLPREDNISOLONE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEO-MEDROL

PHARMACIA AND UPJOHN 0.1%; EQ 3.5MG BASE/GM

A060645 001

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

HEATHER 10MG

A084967 001

VIRILON

STAR PHARMS FL 10MG

A087750 001 Nov 24, 1982

TABLET; Buccal

ANDROID 5

VALEANT PHARM INTL 5MG

A087222 001

ORETON

SCHERING 10MG

A080281 001

TABLET; Buccal, Sublingual

METANDREN

NOVARTIS 5MG  
10MG  
10MG  
25MGN003240 004  
N003240 001  
N003240 005  
N003240 003

METHYLTESTOSTERONE

IMPAK LABS 10MG  
LILLY 10MG  
25MG  
PUREPAC PHARM 10MG  
10MGA084287 001  
A080256 001  
A080256 002  
A080308 001  
A080475 001

**DISCONTINUED DRUG PRODUCT LIST**

6-250(of 375)

\*\* See List Footnote

METHYLTESTOSTERONE

TABLET;BUCCAL, SUBLINGUAL  
METHYLTESTOSTERONE

PVT FORM	10MG	A080475 002
TABLICAPS	25MG	A080475 003
USL PHARMA	5MG	A083836 001
	10MG	A085125 001
	10MG	A080271 001
TABLET;ORAL		
ANDROID 10		
VALEANT PHARMS NORTH	10MG	A086450 001
METHYLTESTOSTERONE		
IMPAX LABS	25MG	A084310 001
INWOOD LABS	10MG	A080839 001
	25MG	A080973 001
KV PHARM	10MG	A084312 001
LANNETT	10MG	A087092 001 Nov 05, 1982
	25MG	A087111 001 Jan 27, 1983
PARKE DAVIS	10MG	A084244 001
	25MG	A084241 001
PUREPAC PHARM	10MG	A080309 001
	25MG	A080310 001
PVT FORM	5MG	A080214 001
	10MG	A080214 002
	25MG	A080214 003
TABLICAPS	10MG	A080313 001
	25MG	A085270 001
WATSON LABS	10MG	A080933 001
	25MG	A080931 001
WEST WARD	10MG	A084331 001
	25MG	A084331 002
	25MG	A084642 001
ORETON METHYL		
SCHERING	10MG	N003158 001
	25MG	N003158 002

METHYPRYLON

CAPSULE;ORAL		
NOLUDAR		
ROCHE	300MG	N009660 008
ELIXIR;ORAL		
NOLUDAR		
ROCHE	50MG/5ML	N009660 007
TABLET;ORAL		
NOLUDAR		
ROCHE	50MG	N009660 002
	200MG	N009660 004

METHYSERGIDE MALEATE

TABLET;ORAL		
SANSERT		
NOVARTIS	2MG	N012516 001

METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE;ORAL		
METOCLOPRAMIDE INTENSOL		
ROXANE	EQ 10MG BASE/ML	A072995 001 Jan 30, 1992
INJECTABLE;INJECTION		
METOCLOPRAMIDE HYDROCHLORIDE		
BEDFORD	EQ 5MG BASE/ML	A072155 001 Mar 30, 1992
	EQ 5MG BASE/ML	A072244 001 Mar 30, 1992
	EQ 5MG BASE/ML	A072247 001 May 18, 1992
HOSPIRA	EQ 5MG BASE/ML	A070505 001 Jun 23, 1989
	EQ 5MG BASE/ML	A070506 001 Jun 22, 1989
	EQ 5MG BASE/ML	A070847 001 Nov 07, 1988
	EQ 5MG BASE/ML	A071291 001 Mar 03, 1989
	EQ 5MG BASE/ML	A071990 001 Jan 18, 1989
	EQ 5MG BASE/ML	A073117 001 Jan 17, 1991
	EQ 5MG BASE/ML	A074147 001 Aug 02, 1996
LYPHOMED	EQ 10MG BASE/2ML	A070293 001 Jan 24, 1986
NORBROOK	EQ 10MG BASE/2ML	A070892 001 Aug 26, 1988

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-251(of 375)

\*\* See List Footnote

METOCLOPRAMIDE HYDROCHLORIDE

## INJECTABLE; INJECTION

## METOCLOPRAMIDE HYDROCHLORIDE

SMITH AND NEPHEW	EQ 5MG BASE/ML EQ 10MG BASE/2ML	A070623 001 Mar 02, 1987 A070622 001 Mar 02, 1987
------------------	------------------------------------	--

## REGLAN

WEST-WARD PHARMS INT	EQ 5MG BASE/ML EQ 10MG BASE/ML	N017862 001 N017862 004 May 28, 1987
----------------------	-----------------------------------	---

## SOLUTION; ORAL

## METOCLOPRAMIDE HYDROCHLORIDE

ACTAVIS MID ATLANTIC	EQ 5MG BASE/5ML	A071340 001 Aug 18, 1988
MORTON GROVE	EQ 5MG BASE/5ML	A070949 001 Mar 06, 1987
PACO	EQ 5MG BASE/5ML	A071665 001 Dec 05, 1988
ROXANE	EQ 5MG BASE/5ML	A072038 001 Dec 05, 1988
SILARX	EQ 5MG BASE/5ML	A073680 001 Oct 27, 1992
TEVA	EQ 5MG BASE/5ML	A070819 001 Jul 10, 1987
	EQ 5MG BASE/5ML	A071315 001 Jun 30, 1993

## REGLAN

+ ROBINS AH	EQ 5MG BASE/5ML **	N018821 001 Mar 25, 1983
-------------	--------------------	--------------------------

## TABLET; ORAL

## CLOPRA

QUANTUM PHARMICS	EQ 5MG BASE EQ 10MG BASE	A072384 001 Jun 02, 1988 A070294 001 Jul 29, 1985
------------------	-----------------------------	--

## CLOPRA-"YELLOW"

QUANTUM PHARMICS	EQ 10MG BASE	A070632 001 Oct 28, 1985
------------------	--------------	--------------------------

## MAXOLON

KING PHARMS	EQ 10MG BASE	A070106 001 Mar 04, 1986
-------------	--------------	--------------------------

## METOCLOPRAMIDE HYDROCHLORIDE

CLONMEL	EQ 10MG BASE	A072639 001 May 09, 1991
FOSUN PHARMA	EQ 5MG BASE	A074478 001 Oct 05, 1995
	EQ 10MG BASE	A072215 001 Jan 30, 1990
	EQ 10MG BASE	A074478 002 Oct 05, 1995
HALSEY	EQ 10MG BASE	A070906 001 Oct 28, 1986
INTERPHARM	EQ 10MG BASE	A071213 001 Sep 24, 1986
MUTUAL PHARM	EQ 10MG BASE	A070660 001 Feb 10, 1987
NORTHSTAR HLTHCARE	EQ 5MG BASE	A078374 001 Nov 30, 2007
	EQ 10MG BASE	A078374 002 Nov 30, 2007
PAR PHARM	EQ 10MG BASE	A070342 001 Mar 25, 1986
SANDOZ	EQ 5MG BASE	A072436 001 Jun 22, 1989
	EQ 10MG BASE	A070850 001 Feb 03, 1987
SCHERING	EQ 10MG BASE	A070598 001 Feb 02, 1987
SUN PHARM INDUSTRIES	EQ 5MG BASE	A071536 002 Jan 16, 1997
	EQ 10MG BASE	A071536 001 Apr 28, 1993
SUPERPHARM	EQ 10MG BASE	A070926 001 Jun 26, 1987
USL PHARMA	EQ 10MG BASE	A070339 001 Jul 29, 1985
WATSON LABS	EQ 10MG BASE	A070363 001 Mar 02, 1987
	EQ 10MG BASE	A070453 001 Jun 06, 1986
	EQ 10MG BASE	A070511 001 Jan 22, 1986
	EQ 10MG BASE	A070645 001 May 11, 1987

## TABLET, ORALLY DISINTEGRATING; ORAL

## METOZOLV ODT

+ SALIX PHARMS	EQ 10MG BASE **	N022246 002 Sep 04, 2009
----------------	-----------------	--------------------------

## REGLAN ODT

MEDA PHARMS	EQ 5MG BASE	N021793 001 Jun 10, 2005
	EQ 10MG BASE	N021793 002 Jun 10, 2005

METOCURINE IODIDE

## INJECTABLE; INJECTION

## METUBINE IODIDE

LILLY	2MG/ML	N006632 003
-------	--------	-------------

METOLAZONE

## TABLET; ORAL

## DIULO

GD SEARLE LLC	2.5MG 5MG 10MG	N018535 001 N018535 002 N018535 003
---------------	----------------------	---

## METOLAZONE

ROXANE	10MG	A076482 002 Apr 29, 2004
TEVA	2.5MG	A076600 001 Jan 06, 2004

**DISCONTINUED DRUG PRODUCT LIST**

6-252(of 375)

\*\* See List Footnote

**METOLAZONE**TABLET;ORAL  
METOLAZONE

WATSON LABS	5MG 10MG 10MG	A076833 001 Mar 01, 2004 A075543 003 Dec 24, 2003 A076891 001 Jul 21, 2004
MYKROX	0.5MG	N019532 001 Oct 30, 1987
UCB INC		

**METOPROLOL FUMARATE**TABLET, EXTENDED RELEASE;ORAL  
LOPRESSOR

NOVARTIS	EQ 100MG TARTRATE EQ 200MG TARTRATE EQ 300MG TARTRATE EQ 400MG TARTRATE	N019786 001 Dec 27, 1989 N019786 002 Dec 27, 1989 N019786 003 Dec 27, 1989 N019786 004 Dec 27, 1989
----------	--	--

**METOPROLOL SUCCINATE**TABLET, EXTENDED RELEASE;ORAL  
METOPROLOL SUCCINATE

NESHER PHARMS	EQ 25MG TARTRATE EQ 50MG TARTRATE EQ 100MG TARTRATE EQ 200MG TARTRATE	A077779 001 Mar 20, 2008 A077176 001 May 14, 2008 A076640 002 May 18, 2007 A076640 001 May 18, 2007
SANDOZ	EQ 25MG TARTRATE EQ 50MG TARTRATE EQ 100MG TARTRATE EQ 200MG TARTRATE	A076969 001 Jul 31, 2006 A076969 002 May 18, 2007 A076969 003 Mar 20, 2008 A076969 004 Mar 20, 2008

**METOPROLOL TARTRATE**INJECTABLE;INJECTION  
METOPROLOL TARTRATE

WATSON LABS	1MGM/ML	A074032 001 Dec 21, 1993
TABLET;ORAL		
METOPROLOL TARTRATE		
APOTHECON	50MG 100MG	A074258 001 Jan 27, 1994 A074258 002 Jan 27, 1994
FOSUN PHARMA	50MG 100MG	A073288 001 Mar 25, 1994 A073289 001 Mar 25, 1994
MYLAN	50MG 100MG	A073666 001 Dec 21, 1993 A073666 002 Dec 21, 1993
PRINSTON INC	50MG 100MG	A074453 001 Apr 27, 1995 A074453 002 Apr 27, 1995
PUREPAC PHARM	50MG 100MG	A074380 001 Jul 29, 1994 A074380 002 Jul 29, 1994
TEVA	50MG 100MG	A074143 001 Sep 30, 1994 A074143 002 Sep 30, 1994
TEVA PHARMS	50MG 100MG	A074333 001 Jan 27, 1994 A074333 002 Jan 27, 1994

**METRIZAMIDE**

INJECTABLE;INJECTION

AMIPAQUE		
GE HEALTHCARE	2.5GM/VIAL 3.75GM/VIAL 6.75GM/VIAL 13.5GM/VIAL	N017982 003 Sep 12, 1983 N017982 001 N017982 002 N017982 004 Sep 12, 1983

**METRONIDAZOLE**

CAPSULE;ORAL

METRONIDAZOLE		
ABLE	375MG	A076505 001 Nov 13, 2003
INJECTABLE;INJECTION		
FLAGYL I.V. RTU IN PLASTIC CONTAINER		
PFIZER	500MG/100ML	N018353 002
METRO I.V.		
B BRAUN	500MG/100ML	N018674 001 Aug 31, 1982
METRONIDAZOLE		
ABBOTT	500MG/100ML	N018889 001 Nov 18, 1983
ABRAXIS PHARM	500MG/100ML	A070071 001 Dec 03, 1984
INTL MEDICATION	500MG/100ML	A070004 001 May 08, 1985
WATSON LABS	500MG/100ML	A070042 001 Dec 20, 1984

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-253(of 375)

\*\* See List Footnote

METRONIDAZOLEINJECTABLE; INJECTION  
METRONIDAZOLE

WEST-WARD PHARMS INT	500MG/100ML 500MG/100ML	A070170 001 Apr 01, 1986 N018907 001 Mar 30, 1984
<b>TABLET; ORAL</b>		
METROMIDOL LABS AF	250MG 500MG	A074523 001 Oct 24, 1996 A074523 002 Oct 24, 1996
METRONIDAZOLE ABLE	250MG 500MG	A076519 001 Jun 27, 2003 A076519 002 Jun 27, 2003
CHARTWELL MOLECULES	250MG 500MG	N018845 001 Aug 18, 1983 N018930 001 Aug 18, 1983
HALSEY	250MG 500MG	A070021 001 Apr 02, 1985 A070593 001 Feb 27, 1986
IVAX SUB TEVA PHARMS	250MG 500MG	N018517 001 N018517 002 May 05, 1982
LNK	250MG	N019029 001 Apr 10, 1984
MUTUAL PHARM	250MG 500MG	N018818 001 Feb 16, 1983 N018818 002 Feb 16, 1983
SANDOZ	250MG 500MG	N018620 001 Mar 04, 1982 N018740 001 Oct 22, 1982 N018620 002 Jun 02, 1983 N018740 002 Oct 22, 1982
SUPERPHARM	250MG 500MG	A070008 001 Dec 11, 1984 A070009 001 Dec 11, 1984
WATSON LABS	250MG 500MG	N018599 001 Sep 17, 1982 N018764 001 Sep 17, 1982 N018599 002 Feb 13, 1984 N018764 002 Dec 20, 1982
PROTOSTAT ORTHO MCNEIL PHARM	250MG 500MG	N018871 001 Mar 02, 1983 N018871 002 Mar 02, 1983
SATRIC SAVAGE LABS	250MG 500MG	A070029 001 Mar 19, 1985 A070731 001 Jun 08, 1987
<b>TABLET, EXTENDED RELEASE; ORAL</b>		
METRONIDAZOLE ABLE	750MG	A076462 001 Jun 25, 2003

METRONIDAZOLE HYDROCHLORIDEINJECTABLE; INJECTION  
FLAGYL I.V.  
PFIZER  
METRONIDAZOLE HYDROCHLORIDE  
ABRAXIS PHARMEQ 500MG BASE/VIAL \*\*  
N018353 001  
EQ 500MG BASE/VIAL  
A070295 001 Oct 15, 1985METYRAPONETABLET; ORAL  
METOPIRONE  
HRA PHARMA250MG  
N012911 001MEXILETINE HYDROCHLORIDECAPSULE; ORAL  
MEXILETINE HYDROCHLORIDE  
IDT AUSTRALIA LTD  
WATSON LABS150MG  
200MG  
250MG  
150MG  
150MG  
200MG  
200MG  
250MG  
250MG  
A074450 001 May 16, 1996  
A074450 002 May 16, 1996  
A074450 003 May 16, 1996  
A074711 001 Feb 26, 1997  
A074865 001 Apr 13, 1998  
A074711 002 Feb 26, 1997  
A074865 002 Apr 13, 1998  
A074711 003 Feb 26, 1997  
A074865 003 Apr 13, 1998MEXITIL  
BOEHRINGER INGELHEIM150MG  
200MG  
250MG  
N018873 002 Dec 30, 1985  
N018873 003 Dec 30, 1985  
N018873 004 Dec 30, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-254(of 375)

\*\* See List Footnote

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

BAYER PHARMS	EQ 1GM BASE/VIAL	A062333 001
	EQ 1GM BASE/VIAL	A062372 005 Jan 13, 1983
	EQ 1GM BASE/VIAL	N050549 001
	EQ 2GM BASE/VIAL	A062333 002
	EQ 2GM BASE/VIAL	A062372 001 May 13, 1982
	EQ 2GM BASE/VIAL	N050549 002
	EQ 3GM BASE/VIAL	A062333 003
	EQ 3GM BASE/VIAL	A062372 002 May 13, 1982
	EQ 3GM BASE/VIAL	A062697 001 Jan 22, 1987
	EQ 3GM BASE/VIAL	N050549 003
	EQ 4GM BASE/VIAL	A062333 004
	EQ 4GM BASE/VIAL	A062372 003 May 13, 1982
	EQ 4GM BASE/VIAL	A062697 002 Jan 22, 1987
	EQ 4GM BASE/VIAL	N050549 004
	EQ 20GM BASE/VIAL	A062372 004 Mar 02, 1988
	EQ 20GM BASE/VIAL	N050549 005 Mar 02, 1988

MICONAZOLE

INJECTABLE; INJECTION

MONISTAT

JANSSEN PHARMA	10MG/ML	N018040 001
----------------	---------	-------------

MICONAZOLE NITRATE

CREAM;TOPICAL

MONISTAT-DERM

INSIGHT PHARMS	2%	N017494 001
----------------	----	-------------

CREAM;VAGINAL

MICONAZOLE NITRATE

TEVA	2%	A074136 001 Jan 04, 1995
TEVA PHARMS	2%	A074030 001 Oct 30, 1992

CREAM, SUPPOSITORY;TOPICAL, VAGINAL

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC	2%,100MG	A074586 001 Jul 17, 1997
----------------------	----------	--------------------------

LOTION;TOPICAL

MONISTAT-DERM

INSIGHT PHARMS	2%	N017739 001
----------------	----	-------------

TAMPON;VAGINAL

MONISTAT 5

PERSONAL PRODS	100MG	N018592 001 Oct 27, 1989
----------------	-------	--------------------------

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

APOTHECON	EQ 1MG BASE/ML	A075620 001 Nov 01, 2000
	EQ 5MG BASE/ML	A075620 002 Nov 01, 2000
	EQ 5MG BASE/ML	A075641 001 Oct 19, 2000
BAXTER HLTHCARE CORP	EQ 1MG BASE/ML	A075637 001 Oct 31, 2000
	EQ 5MG BASE/ML	A075637 002 Oct 31, 2000
BEDFORD	EQ 5MG BASE/ML	A075249 001 Jun 23, 2000
BEN VENUE	EQ 5MG BASE/ML	A075455 001 Jun 20, 2000
HOSPIRA	EQ 1MG BASE/ML	A075396 001 Jun 20, 2000
	EQ 5MG BASE/ML	A075396 002 Jun 20, 2000
	EQ 5MG BASE/ML	A075484 001 Jun 20, 2000
HOSPIRA INC	EQ 1MG BASE/ML	A075409 002 Jun 20, 2000
	EQ 5MG BASE/ML	A075409 001 Jun 20, 2000
IGI LABS INC	EQ 5MG BASE/ML	A075263 001 Jun 26, 2000
INTL MEDICATED	EQ 1MG BASE/ML	A076144 001 Jan 26, 2005
	EQ 5MG BASE/ML	A076144 002 Jan 26, 2005
INTL MEDICATION	EQ 1MG BASE/ML	A076020 001 Jul 16, 2004
	EQ 5MG BASE/ML	A076020 002 Jul 16, 2004
WOCKHARDT	EQ 1MG BASE/ML	A078141 001 May 30, 2008
	EQ 5MG BASE/ML	A078511 001 Nov 10, 2008
	EQ 5MG BASE/ML	A078511 002 May 30, 2008
	EQ 5MG BASE/ML	A078511 002 Nov 10, 2008

VERSED

+	HLR	EQ 1MG BASE/ML **	N018654 002 May 26, 1987
+		EQ 5MG BASE/ML **	N018654 001 Dec 20, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-255(of 375)

\*\* See List Footnote

MIDAZOLAM HYDROCHLORIDE

SYRUP;ORAL

MIDAZOLAM HYDROCHLORIDE

APOTEX INC	EQ 2MG BASE/ML	A077115 001 Sep 09, 2005
SUN PHARM INDS LTD	EQ 2MG BASE/ML	A076058 001 Mar 15, 2002
VERSED		
+ ROCHE	EQ 2MG BASE/ML **	N020942 001 Oct 15, 1998

MIDODRINE HYDROCHLORIDE

TABLET;ORAL

PROAMATINE

+ SHIRE LLC	2.5MG	N019815 001 Sep 06, 1996
+	5MG	N019815 002 Sep 06, 1996
+	10MG	N019815 003 Mar 20, 2002

MILNACIPRAN HYDROCHLORIDE

TABLET;ORAL

MILNACIPRAN HYDROCHLORIDE

WINDLAS HLTHCARE	12.5MG	A205071 001 Jan 27, 2016
	25MG	A205071 002 Jan 27, 2016
	50MG	A205071 003 Jan 27, 2016
	100MG	A205071 004 Jan 27, 2016

MILRINONE LACTATE

INJECTABLE;INJECTION

MILRINONE LACTATE

BAXTER HLTHCARE CORP	EQ 1MG BASE/ML	A076427 001 Sep 21, 2004
HOSPIRA	EQ 1MG BASE/ML	A075830 001 May 28, 2002
	EQ 1MG BASE/ML	A075884 001 May 28, 2002
MYLAN INSTITUTIONAL	EQ 1MG BASE/ML	A076428 001 Jun 16, 2003
WEST-WARD PHARMS INT	EQ 1MG BASE/ML	A075852 001 May 28, 2002

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A076414 001 Aug 18, 2004
BAXTER HLTHCARE	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A076259 001 Aug 08, 2002
RENAISSANCE SSA LLC	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A077151 001 Jul 20, 2005
WEST-WARD PHARMS INT	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075510 001 May 28, 2002

PRIMACOR

+ SANOFI AVENTIS US	EQ 1MG BASE/ML **	N019436 001 Dec 31, 1987
PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER		
+ SANOFI AVENTIS US	EQ 10MG BASE/100ML **	N020343 001 Aug 09, 1994
+	EQ 15MG BASE/100ML **	N020343 002 Aug 09, 1994
+	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	N020343 003 Aug 09, 1994
+	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	N020343 004 Aug 09, 1994
	**	

MINOCYCLINE HYDROCHLORIDE

CAPSULE;ORAL

MINOCIN

+ PRECISION DERMAT	EQ 75MG BASE **	N050649 003 Feb 12, 2001
TRIAZ PHARMS	EQ 50MG BASE	N050315 002
	EQ 100MG BASE	N050315 001

CAPSULE, EXTENDED RELEASE;ORAL

XIMINO

SUN PHARM INDS LTD	EQ 67.5MG BASE	N201922 002 Jul 11, 2012
	EQ 112.5MG BASE	N201922 004 Jul 11, 2012

INJECTABLE;INJECTION

MINOCIN

LEDERLE	EQ 100MG BASE/VIAL	A062139 001
---------	--------------------	-------------

SUSPENSION;ORAL

MINOCIN

PRECISION DERMAT	EQ 50MG BASE/5ML	N050445 001
------------------	------------------	-------------

TABLET;ORAL

MINOCYCLINE HYDROCHLORIDE

+ TRIAX PHARMS	EQ 50MG BASE **	N050451 003 Aug 10, 1982
+	EQ 100MG BASE **	N050451 002 Aug 10, 1982

TABLET, EXTENDED RELEASE;ORAL

MINOCYCLINE HYDROCHLORIDE

BARR LABS INC	EQ 45MG BASE	A065485 001 Mar 17, 2009
	EQ 65MG BASE	A065485 004 May 18, 2012
	EQ 80MG BASE	A065485 007 Apr 26, 2017
	EQ 90MG BASE	A065485 002 Mar 17, 2009

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-256(of 375)

\*\* See List Footnote

MINOCYCLINE HYDROCHLORIDETABLET, EXTENDED RELEASE;ORAL  
MINOCYCLINE HYDROCHLORIDE

	EQ 105MG BASE	A065485 008	Apr 26, 2017
	EQ 115MG BASE	A065485 005	May 18, 2012
	EQ 135MG BASE	A065485 003	Mar 17, 2009
IMPAX LABS INC	EQ 45MG BASE	A090024 001	Feb 03, 2009
	EQ 90MG BASE	A090024 002	Feb 03, 2009
	EQ 135MG BASE	A090024 003	Feb 03, 2009
LUPIN LTD	EQ 55MG BASE	A091424 002	Nov 30, 2011
MYLAN PHARMS INC	EQ 45MG BASE	A090911 001	Jul 20, 2010
	EQ 80MG BASE	A203443 002	Aug 22, 2014
	EQ 90MG BASE	A090911 002	Jul 20, 2010
	EQ 105MG BASE	A203443 003	Aug 22, 2014
	EQ 135MG BASE	A090911 003	Jul 20, 2010
SOLODYNE			
+ MEDICIS	EQ 45MG BASE **	N050808 001	May 08, 2006
+	EQ 90MG BASE **	N050808 002	May 08, 2006
+	EQ 135MG BASE **	N050808 003	May 08, 2006

MINOXIDIL

SOLUTION;TOPICAL

## MINOXIDIL (FOR MEN)

APOTEX INC	2%	A074924 001	Apr 29, 1998
BAUSCH AND LOMB	2%	A074643 001	Apr 09, 1996
COPLEY PHARM	2%	A074500 001	May 23, 1996
SIGHT PHARMS	2%	A074743 002	Oct 18, 1996
TEVA	2%	A074589 001	Apr 05, 1996

## MINOXIDIL (FOR WOMEN)

APOTEX INC	2%	A074924 002	Apr 29, 1998
SIGHT PHARMS	2%	A074743 001	Oct 18, 1996

## MINOXIDIL EXTRA STRENGTH (FOR MEN)

APOTEX INC	5%	A075839 001	Oct 01, 2001
------------	----	-------------	--------------

TABLET;ORAL

## LONITEN

+ PHARMACIA AND UPJOHN	2.5MG **	N018154 001
+	10MG **	N018154 003

## MINODYL

QUANTUM PHARMICS	2.5MG	A072153 001	Jul 13, 1988
	10MG	A071534 001	Mar 19, 1987

## MINOXIDIL

ROYCE LABS	2.5MG	A071799 001	Nov 10, 1987
	10MG	A071796 001	Nov 10, 1987
USL PHARMA	2.5MG	A071537 001	Dec 16, 1988

MIRTAZAPINE

TABLET;ORAL

## MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076241 001	Jun 25, 2003
	15MG	A076308 001	Jun 20, 2003
	30MG	A076241 002	Jun 25, 2003
	30MG	A076308 002	Jun 20, 2003
	45MG	A076241 003	Jun 25, 2003
	45MG	A076308 003	Jun 20, 2003
ACTAVIS LABS FL INC	15MG	A076336 001	Jun 20, 2003
	30MG	A076336 002	Jun 20, 2003
	45MG	A076336 003	Jun 20, 2003
IVAX SUB TEVA PHARMS	15MG	A076244 001	Dec 22, 2003
	30MG	A076244 002	Dec 22, 2003
	45MG	A076244 003	Dec 22, 2003
MYLAN PHARMS INC	15MG	A076176 001	Jun 19, 2003
	30MG	A076176 002	Jun 19, 2003
	45MG	A076176 003	Jun 19, 2003
ROXANE	15MG	A076270 001	Jun 19, 2003
	30MG	A076270 002	Jun 19, 2003
	45MG	A076270 003	Jun 19, 2003
UPSHER-SMITH LABS	15MG	A076189 001	Jun 19, 2003
	30MG	A076189 002	Jun 19, 2003
	45MG	A076189 003	Jun 19, 2003

**DISCONTINUED DRUG PRODUCT LIST**

6-257(of 375)

\*\* See List Footnote

MIRTAZAPINETABLET, ORALLY DISINTEGRATING;ORAL  
MIRTAZAPINE

ACTAVIS ELIZABETH	15MG	A076689 001 Aug 31, 2005
	15MG	A077959 001 Feb 14, 2011
	30MG	A076689 002 Aug 31, 2005
	30MG	A077959 002 Feb 14, 2011
	45MG	A076689 003 Aug 31, 2005
	45MG	A077959 003 Feb 14, 2011

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN		
HOSPIRA	20MG/VIAL	A064106 001 Nov 29, 1995
MITOZYTREX		
+ SUPERGEN	5MG/VIAL **	N050763 001 Nov 14, 2002
MUTAMYCIN		
+ BRISTOL	5MG/VIAL	N050450 001
+ BRISTOL MYERS	20MG/VIAL	N050450 002
	5MG/VIAL	A062336 001
	20MG/VIAL	A062336 002
	40MG/VIAL	A062336 003 Mar 10, 1988

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

MITOXANTRONE HYDROCHLORIDE		
FRESENIUS KABI ONCOL	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	A078606 001 May 14, 2008
	EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML)	A078606 002 May 14, 2008
	EQ 30MG BASE/15ML (EQ 2MG BASE/ML)	A078606 003 May 14, 2008
NOVANTRONE		
EMD SERONO	EQ 20MG BASE/10ML (EQ 2MG BASE/ML)	N019297 001 Dec 23, 1987
+	EQ 25MG BASE/12.5ML (EQ 2MG BASE/ML) **	N019297 002 Dec 23, 1987
+	EQ 30MG BASE/15ML (EQ 2MG BASE/ML) **	N019297 003 Dec 23, 1987

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER		
ABBVIE	EQ 0.5MG BASE/ML	N020098 002 Jan 22, 1992
	EQ 50MG BASE/100ML	N020098 003 Jan 22, 1992
MIVACURIUM CHLORIDE		
MYLAN LABS LTD	EQ 2MG BASE/ML	A078562 001 Apr 30, 2009
SOLUTION; INTRAVENOUS		
MIVACRON		
+ ABBVIE	EQ 2MG BASE/ML (EQ 2MG BASE/ML) **	N020098 001 Jan 22, 1992

MOEXIPRIL HYDROCHLORIDE

TABLET;ORAL

UNIVASC		
UCB INC	7.5MG **	N020312 001 Apr 19, 1995
	15MG **	N020312 002 Apr 19, 1995

MOLINDONE HYDROCHLORIDE

CAPSULE;ORAL

MOBAN		
+ ENDO PHARMS	5MG **	N017111 001
+	10MG **	N017111 002

CONCENTRATE;ORAL

MOBAN		
ENDO PHARMS	20MG/ML	N017938 001

TABLET;ORAL

MOBAN		
+ ENDO PHARMS	5MG **	N017111 004
+	10MG **	N017111 005
+	25MG **	N017111 006
+	50MG **	N017111 007
+	100MG **	N017111 008

**DISCONTINUED DRUG PRODUCT LIST**

6-258(of 375)

\*\* See List Footnote

MOMETASONE FUROATE

CREAM;TOPICAL ELOCON	MERCK SHARP DOHME	0.1%	N019625 001 May 06, 1987
OINTMENT;TOPICAL MOMETASONE FUROATE	TARO	0.1%	A076624 001 Dec 03, 2004

MONOBENZONE

CREAM;TOPICAL BENOQUIN	VALEANT PHARM INTL	20%	N008173 003
---------------------------	--------------------	-----	-------------

MONOCTANOIN

LIQUID;PERFUSION, BILIARY MOCTANIN	ETHITEK	100%	N019368 001 Oct 29, 1985
---------------------------------------	---------	------	--------------------------

MORICIZINE HYDROCHLORIDE

TABLET;ORAL ETHMOZINE	SHIRE	200MG 250MG 300MG	N019753 001 Jun 19, 1990 N019753 002 Jun 19, 1990 N019753 003 Jun 19, 1990
--------------------------	-------	-------------------------	--

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE;ORAL AVINZA	KING PHARMS LLC	30MG 45MG 60MG 75MG 90MG 120MG	N021260 001 Mar 20, 2002 N021260 005 Dec 18, 2008 N021260 002 Mar 20, 2002 N021260 006 Dec 18, 2008 N021260 003 Mar 20, 2002 N021260 004 Mar 20, 2002
--	-----------------	---	--

## INJECTABLE;INJECTION

MORPHINE SULFATE	+ HOSPIRA INC ICU MEDICAL INC SPECGX LLC	15MG/ML 0.5MG/ML 1MG/ML 2MG/ML	N202515 005 Nov 14, 2011 N019917 001 Oct 30, 1992 N020631 001 Jul 03, 1996 N020631 002 Jul 03, 1996
	WATSON LABS	0.5MG/ML 0.5MG/ML 1MG/ML 1MG/ML	A073373 001 Sep 30, 1991 A073375 001 Sep 30, 1991 A073374 001 Sep 30, 1991 A073376 001 Sep 30, 1991

## INJECTABLE, LIPOSOMAL;EPIDURAL

DEPODUR	PACIRA PHARMS INC	10MG/ML (10MG/ML) 15MG/1.5ML (10MG/ML) 20MG/2ML (10MG/ML)	N021671 001 May 18, 2004 N021671 002 May 18, 2004 N021671 003 May 18, 2004
---------	-------------------	---	--

## TABLET, EXTENDED RELEASE;ORAL

MORPHINE SULFATE	WATSON LABS	100MG	A075656 001 Jan 30, 2001
ORAMORPH SR	XANODYNE PHARMS INC	15MG 30MG 60MG 100MG	N019977 004 Nov 23, 1994 N019977 001 Aug 15, 1991 N019977 002 Aug 15, 1991 N019977 003 Aug 15, 1991

MOXALACTAM DISODIUM

INJECTABLE;INJECTION MOXAM	LILLY	EQ 250MG BASE/VIAL EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL EQ 2GM BASE/VIAL EQ 10GM BASE/VIAL	N050550 001 N050550 002 N050550 003 N050550 004 N050550 008
-------------------------------	-------	---	---

**DISCONTINUED DRUG PRODUCT LIST**

6-259(of 375)

\*\* See List Footnote

MOXIFLOXACIN HYDROCHLORIDE

SOLUTION;IV (INFUSION)

AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER  
+ BAYER HLTHCARE 400MG/250ML (1.6MG/ML)

N021277 001 Nov 30, 2001

MUPIROCIN

OINTMENT;TOPICAL

BACTROBAN

+ GLAXOSMITHKLINE 2% \*\*

N050591 001 Dec 31, 1987

MYCOPHENOLATE MOFETIL

CAPSULE;ORAL

MYCOPHENOLATE MOFETIL

DR REDDYS LABS LTD 250MG  
JUBILANT CADISTA 250MG  
ZYDUS PHARMS USA INC 250MGA091315 001 Oct 27, 2011  
A090762 001 Dec 15, 2014  
A065433 001 May 04, 2009

TABLET;ORAL

MYCOPHENOLATE MOFETIL

DR REDDYS LABS LTD 500MG  
JUBILANT CADISTA 500MG  
ZYDUS PHARMS USA INC 500MGA090464 001 Sep 13, 2010  
A090661 001 Dec 15, 2014  
A065477 001 May 04, 2009NABUMETONE

TABLET;ORAL

NABUMETONE

COPLEY PHARM 750MG  
OXFORD PHARMS 500MG  
750MG  
SANDOZ 500MG  
750MG  
SCIEGEN PHARMS INC 500MG  
750MGA075179 001 Jun 06, 2000  
A079093 001 Feb 27, 2009  
A079093 002 Feb 27, 2009  
A075590 001 Feb 25, 2002  
A075590 002 Feb 25, 2002  
A078420 001 Sep 24, 2008  
A078420 002 Sep 24, 2008

RELAFEN

+ SMITHKLINE BEECHAM 500MG \*\*  
+ 750MG \*\*N019583 001 Dec 24, 1991  
N019583 002 Dec 24, 1991NADOLOL

TABLET;ORAL

CORGARD

US WORLDMEDS LLC 120MG  
160MGN018063 003  
N018063 004

NADOLOL

IVAX SUB TEVA PHARMS 120MG  
160MG  
TEVA PHARMS 80MG  
120MG  
160MGA074255 002 Jan 24, 1996  
A074255 003 Jan 24, 1996  
A074368 001 Aug 31, 1994  
A074368 002 Aug 31, 1994  
A074368 003 Aug 31, 1994NAFCILLIN SODIUM

CAPSULE;ORAL

UNIPEN

WYETH AYERST EQ 250MG BASE

N050111 001

FOR SOLUTION;ORAL

UNIPEN

WYETH AYERST EQ 250MG BASE/5ML

N050199 001

INJECTABLE;INJECTION

NAFCILLIN SODIUM

APOTHECON EQ 500MG BASE/VIAL  
EQ 1GM BASE/VIAL  
EQ 2GM BASE/VIAL  
EQ 4GM BASE/VIAL  
SANDOZ EQ 500MG BASE/VIAL  
WATSON LABS INC EQ 500MG BASE/VIAL  
EQ 1GM BASE/VIAL  
EQ 1.5GM BASE/VIAL  
EQ 2GM BASE/VIAL  
EQ 4GM BASE/VIAL  
EQ 10GM BASE/VIALA061984 001  
A061984 002  
A061984 003  
A061984 005  
A062527 001 Aug 02, 1984  
A062844 001 Oct 26, 1988  
A062844 002 Oct 26, 1988  
A062844 003 Oct 26, 1988  
A062844 004 Oct 26, 1988  
A062844 005 Oct 26, 1988  
A063008 001 Sep 29, 1988

NALLPEN

GLAXOSMITHKLINE EQ 500MG BASE/VIAL  
EQ 1GM BASE/VIAL  
EQ 1GM BASE/VIALA061999 001  
A061999 002  
A062755 001 Dec 19, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-260(of 375)

\*\* See List Footnote

NAFCILLIN SODIUMINJECTABLE; INJECTION  
NALLPEN

UNIPEN		EQ 2GM BASE/VIAL	A061999 003
WYETH AYERST		EQ 2GM BASE/VIAL	A062755 002 Dec 19, 1986
+		EQ 10GM BASE/VIAL	A061999 004
UNIPEN			
WYETH AYERST		EQ 500MG BASE/VIAL **	A062717 001 Dec 16, 1986
+		EQ 500MG BASE/VIAL **	N050320 001
		EQ 1GM BASE/VIAL **	A062717 002 Dec 16, 1986
		EQ 2GM BASE/VIAL **	A062717 004 Dec 16, 1986
+		EQ 2GM BASE/VIAL **	N050320 003
+		EQ 4GM BASE/VIAL **	N050320 004
+		EQ 10GM BASE/VIAL **	N050320 005
+		EQ 20GM BASE/VIAL **	N050320 006
UNIPEN IN PLASTIC CONTAINER			
WYETH AYERST		EQ 1GM BASE/VIAL **	N050320 002
TABLET; ORAL			
UNIPEN			
WYETH AYERST		EQ 500MG BASE	N050462 001

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE			
ABRAXIS PHARM		10MG/ML	A070751 001 Jul 02, 1986
		20MG/ML	A070752 001 Sep 24, 1986
NALBUPHINE HYDROCHLORIDE			
ABBOTT		20MG/ML	A070917 001 Feb 03, 1989
ABBVIE		1.5MG/ML	N020200 001 Mar 12, 1993
BARR		10MG/ML	A074471 001 Mar 19, 1998
		20MG/ML	A074471 002 Mar 19, 1998
IGI LABS INC		10MG/ML	A072070 001 Apr 10, 1989
		10MG/ML	A072071 001 Apr 10, 1989
		20MG/ML	A072072 001 Apr 10, 1989
		20MG/ML	A072073 001 Apr 10, 1989
		20MG/ML	A072074 001 Apr 10, 1989
		20MG/ML	A072075 001 Apr 10, 1989
NUBAIN			
+	PAR PHARM INC	10MG/ML **	N018024 001
+		20MG/ML **	N018024 002 May 27, 1982

NALIDIXIC ACID

SUSPENSION; ORAL

NEGGRAM			
SANOFI AVENTIS US		250MG/5ML	N017430 001
TABLET; ORAL			
NALIDIXIC ACID			
SUN PHARM INDUSTRIES		250MG	A070270 001 Jun 29, 1988
		500MG	A070271 001 Jun 29, 1988
		1GM	A070272 001 Jun 29, 1988
WATSON LABS		250MG	A071936 001 Jun 29, 1988
		500MG	A072061 001 Jun 29, 1988
		1GM	A071919 001 Jun 29, 1988
NEGGRAM			
SANOFI AVENTIS US		250MG	N014214 002
		500MG	N014214 004
		1GM	N014214 005

NALMEFENE HYDROCHLORIDE

INJECTABLE; INJECTION

REVEX			
+	EUROHLTH INTL SARL	EQ 0.1MG BASE/ML **	N020459 001 Apr 17, 1995
+		EQ 1MG BASE/ML **	N020459 002 Apr 17, 1995

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE			
WEST-WARD PHARMS INT		0.4MG/ML	A070298 001 Sep 24, 1986
		0.4MG/ML	A070496 001 Sep 24, 1986
WYETH AYERST		0.02MG/ML	A070188 001 Sep 24, 1986
		0.02MG/ML	A070189 001 Sep 24, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-261(of 375)

\*\* See List Footnote

NALOXONE HYDROCHLORIDEINJECTABLE; INJECTION  
NALOXONE

	0.4MG/ML	A070190 001 Sep 24, 1986
	0.4MG/ML	A070191 001 Sep 24, 1986
NALOXONE HYDROCHLORIDE		
ABRAXIS PHARM	0.02MG/ML	A070648 001 Nov 17, 1986
	0.02MG/ML	A070661 001 Nov 17, 1986
	0.4MG/ML	A070649 001 Nov 17, 1986
	1MG/ML	A071604 001 Dec 16, 1988
ASTRAZENECA	0.02MG/ML	A072081 001 Apr 11, 1989
EUROHLTH INTL SARL	0.02MG/ML	A071272 001 May 24, 1988
	1MG/ML	A071273 001 May 24, 1988
	1MG/ML	A071274 001 May 24, 1988
HOSPIRA	0.02MG/ML	A071287 001 May 24, 1988
	0.02MG/ML	A070171 001 Sep 24, 1986
	0.02MG/ML	A070252 001 Jan 16, 1987
	0.02MG/ML	A070253 001 Jan 16, 1987
	0.4MG/ML	A070255 001 Jan 07, 1987
IGI LABS INC	0.02MG/ML	A072082 001 Apr 11, 1989
	0.02MG/ML	A072083 001 Apr 11, 1989
	0.02MG/ML	A072084 001 Apr 11, 1989
	0.02MG/ML	A072085 001 Apr 11, 1989
	0.4MG/ML	A072086 001 Apr 11, 1989
	0.4MG/ML	A072087 001 Apr 11, 1989
	0.4MG/ML	A072088 001 Apr 11, 1989
	0.4MG/ML	A072089 001 Apr 11, 1989
	0.4MG/ML	A072090 001 Apr 11, 1989
	1MG/ML	A072091 001 Apr 11, 1989
	1MG/ML	A072092 001 Apr 11, 1989
	1MG/ML	A072093 001 Apr 11, 1989
INTL MEDICATION	0.4MG/ML	A070417 001 Sep 24, 1986
	1MG/ML	A072115 001 Apr 27, 1988
MARSAM PHARMS LLC	0.4MG/ML	A071811 001 Jul 19, 1988
SMITH AND NEPHEW	0.02MG/ML	A071671 001 Nov 17, 1987
	0.4MG/ML	A071681 001 Nov 17, 1987
	0.4MG/ML	A071682 001 Nov 17, 1987
SOLOPAK	0.02MG/ML	A071672 001 Nov 17, 1987
	0.4MG/ML	A071683 001 Nov 17, 1987
WATSON LABS	0.4MG/ML	A071339 001 Nov 18, 1987
NARCAN		
+ ADAPT	0.02MG/ML **	N016636 002
+	0.4MG/ML **	N016636 001
+	1MG/ML **	N016636 003 Jun 14, 1982
BRISTOL MYERS SQUIBB	0.4MG/ML	A071083 001 Jul 28, 1988
	1MG/ML	A071084 001 Jul 28, 1988
	1MG/ML	A071311 001 Jul 28, 1988
SOLUTION;INTRAMUSCULAR, SUBCUTANEOUS		
EVZIO		
+ KALEO INC	0.4MG/0.4ML (0.4MG/0.4ML)	N205787 001 Apr 03, 2014

NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDETABLET, EXTENDED RELEASE; ORAL  
TARGINITQ

+ PURDUE PHARMA LP	5MG;10MG	N205777 001 Jul 23, 2014
+	10MG;20MG	N205777 002 Jul 23, 2014
+	20MG;40MG	N205777 003 Jul 23, 2014

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDETABLET;ORAL  
TALWIN NX  
SANOFI AVENTIS US EQ 0.5MG BASE;EQ 50MG BASE \*\* N018733 001 Dec 16, 1982NALTREXONE HYDROCHLORIDE

TABLET;ORAL		
NALTREXONE HYDROCHLORIDE		
SANDOZ	50MG	A075434 001 Mar 08, 2000
REVIA		
TEVA WOMENS	50MG	N018932 001 Nov 20, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-262(of 375)

\*\* See List Footnote

NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDECAPSULE, EXTENDED RELEASE;ORAL  
TROXYCA ER

PFIZER INC	1.2MG;10MG	N207621 001	Aug 19, 2016
	2.4MG;20MG	N207621 002	Aug 19, 2016
	3.6MG;30MG	N207621 003	Aug 19, 2016
	4.8MG;40MG	N207621 004	Aug 19, 2016
	7.2MG;60MG	N207621 005	Aug 19, 2016
	9.6MG;80MG	N207621 006	Aug 19, 2016

NANDROLONE DECANOATEINJECTABLE;INJECTION  
DECA-DURABOLIN

ASPEN GLOBAL INC	50MG/ML	N013132 001	Jun 12, 1986
+	100MG/ML	N013132 002	Jun 12, 1986
	200MG/ML **	N013132 003	Jun 12, 1986
NANDROLONE DECANOATE			
ABRAXIS PHARM	100MG/ML	A088290 001	Oct 03, 1983
	200MG/ML	A088317 001	Oct 14, 1983
AKORN	100MG/ML	A087519 001	Sep 28, 1983
WATSON LABS	50MG/ML	A086385 001	Jan 13, 1984
	50MG/ML	A087598 001	Oct 06, 1983
	50MG/ML	A088554 001	Feb 10, 1986
	100MG/ML	A086598 001	Jan 13, 1984
	100MG/ML	A087599 001	Oct 06, 1983
	200MG/ML	A088128 001	Dec 05, 1983

NANDROLONE PHENPROPIONATEINJECTABLE;INJECTION  
DURABOLIN

ORGANON USA INC	25MG/ML	N011891 001	
	50MG/ML	N011891 002	
NANDROLONE PHENPROPIONATE			
WATSON LABS	25MG/ML	A086386 001	Jun 17, 1983
	50MG/ML	A087488 001	Jun 17, 1983

NAPHAZOLINE HYDROCHLORIDESOLUTION/DROPS;OPHTHALMIC  
ALBALON

ALLERGAN	0.1% **	A080248 001	
NAFAZAIR			
BAUSCH AND LOMB	0.1%	A040073 001	May 25, 1994
PHARMAFAIR	0.1%	A088101 001	Apr 15, 1983
NAPHCON FORTE			
ALCON	0.1%	A080229 001	
OPCON			
BAUSCH AND LOMB	0.1%	A087506 001	
VASOCON			
NOVARTIS	0.1%	A080235 002	Mar 24, 1983

NAPROXENTABLET;ORAL  
NAPROXEN

CHARTWELL MOLECULES	250MG	A074410 001	Apr 28, 1995
	375MG	A074410 002	Apr 28, 1995
	500MG	A074410 003	Apr 28, 1995
DAVA PHARMS INC	250MG	A074105 001	Dec 21, 1993
	375MG	A074105 002	Dec 21, 1993
	500MG	A074105 003	Dec 21, 1993
HAMILTON PHARMS	250MG	A074110 001	Oct 30, 1992
	375MG	A074110 002	Oct 30, 1992
	500MG	A074110 003	Oct 30, 1992
HIKMA INTL PHARMS	250MG	A076494 001	Jan 14, 2004
	375MG	A076494 002	Jan 14, 2004
	500MG	A076494 003	Jan 14, 2004
IVAX SUB TEVA PHARMS	250MG	A074111 001	Feb 28, 1995
	375MG	A074111 002	Feb 28, 1995
	500MG	A074111 003	Feb 28, 1995
PLIVA	250MG	A074182 001	Jun 27, 1996
	375MG	A074182 002	Jun 27, 1996
	500MG	A074182 003	Jun 27, 1996

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-263(of 375)

\*\* See List Footnote

NAPROXEN

TABLET;ORAL

NAPROXEN

PUREPAC PHARM	250MG	A074263 001	Dec 21, 1993
	375MG	A074263 002	Dec 21, 1993
	500MG	A074263 003	Dec 21, 1993
ROXANE	250MG	A074211 001	Feb 28, 1994
	375MG	A074211 002	Feb 28, 1994
	500MG	A074211 003	Feb 28, 1994
SANDOZ	250MG	A074140 001	Dec 21, 1993
	375MG	A074140 002	Dec 21, 1993
	500MG	A074140 003	Dec 21, 1993
TEVA	250MG	A074129 001	Dec 21, 1993
	250MG	A074216 001	Apr 11, 1996
	375MG	A074129 002	Dec 21, 1993
	375MG	A074216 002	Apr 11, 1996
	500MG	A074129 003	Dec 21, 1993
	500MG	A074216 003	Apr 11, 1996
TEVA PHARMS	250MG	A074207 001	Dec 21, 1993
	375MG	A074207 002	Dec 21, 1993
	500MG	A074207 003	Dec 21, 1993
WATSON LABS	250MG	A074457 001	May 31, 1995
	375MG	A074457 002	May 31, 1995
	500MG	A074457 003	May 31, 1995
WATSON LABS TEVA	250MG	A074163 001	Feb 10, 1995
	375MG	A074163 002	Feb 10, 1995
	500MG	A074163 003	Feb 10, 1995

TABLET, DELAYED RELEASE;ORAL

NAPROXEN

ACTAVIS ELIZABETH	375MG	A074936 001	Feb 24, 1998
	500MG	A074936 002	Feb 24, 1998
MYLAN PHARMS INC	375MG	A075390 001	Apr 19, 2001
	500MG	A075390 002	Apr 19, 2001
SANDOZ	375MG	A075061 001	Feb 18, 1998
	500MG	A075061 002	Feb 18, 1998

NAPROXEN SODIUM

TABLET;ORAL

NAPROXEN SODIUM

ABLE	EQ 250MG BASE	A076544 001	Aug 22, 2003
	EQ 500MG BASE	A076544 002	Aug 22, 2003
CONTRACT PHARMACAL	EQ 200MG BASE	A074789 001	Feb 27, 1997
HAMILTON PHARMS	EQ 250MG BASE	A074106 001	Aug 31, 1993
	EQ 500MG BASE	A074106 002	Aug 31, 1993
HIKMA	EQ 250MG BASE	A074480 002	Feb 18, 1998
	EQ 500MG BASE	A074480 001	May 14, 1996
IVAX SUB TEVA PHARMS	EQ 250MG BASE	A074230 001	Mar 14, 1995
	EQ 500MG BASE	A074230 002	Mar 14, 1995
MYLAN	EQ 250MG BASE	A074367 001	Aug 31, 1994
	EQ 500MG BASE	A074367 002	Aug 31, 1994
PLIVA	EQ 250MG BASE	A074242 001	Jun 20, 1996
	EQ 500MG BASE	A074242 002	Jun 20, 1996
PUREPAC PHARM	EQ 250MG BASE	A074319 001	Mar 20, 1995
	EQ 500MG BASE	A074319 002	Mar 20, 1995
ROXANE	EQ 250MG BASE	A074257 001	Dec 21, 1993
	EQ 500MG BASE	A074257 002	Dec 21, 1993
SANDOZ	EQ 200MG BASE	A074646 001	Jan 13, 1997
	EQ 250MG BASE	A074162 001	Dec 21, 1993
	EQ 250MG BASE	A074495 001	Dec 05, 1994
	EQ 500MG BASE	A074162 002	Dec 21, 1993
	EQ 500MG BASE	A074495 002	Dec 05, 1994
TEVA	EQ 250MG BASE	A074142 001	Dec 21, 1993
	EQ 500MG BASE	A074142 002	Dec 21, 1993
TEVA PHARMS	EQ 250MG BASE	A074289 001	Jan 27, 1994
	EQ 500MG BASE	A074289 002	Jan 27, 1994
WATSON LABS	EQ 250MG BASE	A074195 001	Dec 21, 1993
	EQ 250MG BASE	A074455 001	May 31, 1995
	EQ 500MG BASE	A074195 002	Dec 21, 1993
	EQ 500MG BASE	A074455 002	May 31, 1995

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-264(of 375)

\*\* See List Footnote

NATEGLINIDE

TABLET;ORAL

NATEGLINIDE

TEVA PHARMS

60MG

120MG

A077467 001 Sep 09, 2009

A077467 002 Sep 09, 2009

NEBIVOLOL HYDROCHLORIDE

TABLET;ORAL

NEBIVOLOL HYDROCHLORIDE

ALKEM LABS LTD

EQ 2.5MG BASE

A203741 001 Jun 24, 2015

EQ 5MG BASE

A203741 002 Jun 24, 2015

EQ 10MG BASE

A203741 003 Jun 24, 2015

EQ 20MG BASE

A203741 004 Jun 24, 2015

AMERIGEN PHARMS LTD

EQ 2.5MG BASE

A203659 001 Apr 16, 2015

EQ 5MG BASE

A203659 002 Apr 16, 2015

EQ 10MG BASE

A203659 003 Apr 16, 2015

EQ 20MG BASE

A203659 004 Apr 16, 2015

GLENMARK PHARMS LTD

EQ 2.5MG BASE

A203821 001 May 25, 2017

EQ 5MG BASE

A203821 002 May 25, 2017

EQ 10MG BASE

A203821 003 May 25, 2017

EQ 20MG BASE

A203821 004 May 25, 2017

INDCHEMIE HEALTH

EQ 2.5MG BASE

A203828 001 Jul 29, 2015

EQ 5MG BASE

A203828 002 Jul 29, 2015

EQ 10MG BASE

A203828 003 Jul 29, 2015

EQ 20MG BASE

A203828 004 Jul 29, 2015

WATSON LABS INC

EQ 2.5MG BASE

A203683 001 Nov 27, 2015

EQ 5MG BASE

A203683 002 Nov 27, 2015

EQ 10MG BASE

A203683 003 Nov 27, 2015

EQ 20MG BASE

A203683 004 Nov 27, 2015

NEDOCROMIL SODIUM

AEROSOL, METERED;INHALATION

TILADE

KING PHARMS LLC

1.75MG/INH

N019660 001 Dec 30, 1992

SOLUTION;INHALATION

TILADE

SANOFI AVENTIS US

0.5%

N020750 001 Oct 01, 1997

NEFAZODONE HYDROCHLORIDE

TABLET;ORAL

NEFAZODONE HYDROCHLORIDE

DR REDDYS LABS INC

50MG

A076309 001 Sep 16, 2003

100MG

A076309 002 Sep 16, 2003

150MG

A076309 003 Sep 16, 2003

200MG

A076309 004 Sep 16, 2003

250MG

A076309 005 Sep 16, 2003

IDT AUSTRALIA LTD

50MG

A076072 001 Sep 16, 2003

100MG

A076072 002 Sep 16, 2003

150MG

A076072 003 Sep 16, 2003

200MG

A076072 004 Sep 16, 2003

250MG

A076072 005 Sep 16, 2003

IVAX SUB TEVA PHARMS

50MG

A075763 001 Sep 16, 2003

100MG

A075763 002 Sep 16, 2003

150MG

A075763 003 Sep 16, 2003

200MG

A075763 004 Sep 16, 2003

250MG

A075763 005 Sep 16, 2003

MYLAN

100MG

A076129 002 Sep 16, 2003

150MG

A076129 003 Sep 16, 2003

200MG

A076129 004 Sep 16, 2003

250MG

A076129 005 Sep 16, 2003

ROXANE

50MG

A076196 001 Sep 16, 2003

100MG

A076196 002 Sep 16, 2003

150MG

A076196 003 Sep 16, 2003

200MG

A076196 004 Sep 16, 2003

250MG

A076196 005 Sep 16, 2003

SANDOZ

50MG

A076302 001 Sep 16, 2003

100MG

A076302 002 Sep 16, 2003

150MG

A076302 003 Sep 16, 2003

200MG

A076302 004 Sep 16, 2003

250MG

A076302 005 Sep 16, 2003

SUN PHARM INDs LTD

50MG

A076409 001 Sep 16, 2003

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-265(of 375)

\*\* See List Footnote

NEFAZODONE HYDROCHLORIDE

TABLET;ORAL

NEFAZODONE HYDROCHLORIDE

	100MG	A076409 002	Sep 16, 2003
	150MG	A076409 003	Sep 16, 2003
	200MG	A076409 004	Sep 16, 2003
	250MG	A076409 005	Sep 16, 2003
WATSON LABS	100MG	A076073 002	Sep 16, 2003
	150MG	A076073 003	Sep 16, 2003
	200MG	A076073 004	Sep 16, 2003
	250MG	A076073 005	Sep 16, 2003
SERZONE			
+ BRISTOL MYERS SQUIBB	50MG **	N020152 001	Dec 22, 1994
+	100MG **	N020152 002	Dec 22, 1994
+	150MG **	N020152 003	Dec 22, 1994
+	200MG **	N020152 004	Dec 22, 1994
+	250MG **	N020152 005	Dec 22, 1994
+	300MG **	N020152 006	Dec 22, 1994

NELFINAVIR MESYLATE

POWDER;ORAL

VIRACEPT

AGOURON PHARMS EQ 50MG BASE/SCOOPFUL

N020778 001 Mar 14, 1997

NEOMYCIN SULFATE

POWDER;FOR RX COMPOUNDING

NEO-RX

X GEN PHARMS 100%

A061579 001

SOLUTION;ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 87.5MG BASE/5ML

N050285 001

NEO-FRADIN

X GEN PHARMS EQ 87.5MG BASE/5ML

A065010 001 May 23, 2002

TABLET;ORAL

MYCIFRADIN

PHARMACIA AND UPJOHN EQ 350MG BASE

A060520 001

NEOBIOTIC

PFIZER EQ 350MG BASE

A060475 001

NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB 500MG

A060365 001

LANNETT 500MG

A060607 001

LILLY 500MG

A060385 001

ROXANE 500MG

A062173 001

SANDOZ 500MG

A061586 001

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM;TOPICAL

NEOSPORIN

GLAXOSMITHKLINE EQ 3.5MG BASE/GM;10,000 UNITS/GM

N050176 002 Jan 14, 1985

OINTMENT;OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/GM;10,000 UNITS/GM

N050344 002

SOLUTION/DROPS;OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/ML;16,250 UNITS/ML

A062339 001 Nov 30, 1984

EQ 3.5MG BASE/ML;16,250 UNITS/ML

N050456 001

NEOMYCIN SULFATE; POLYMYXIN B SULFATE; PREDNISOLONE ACETATE

SUSPENSION/DROPS;OPHTHALMIC

POLY-PRED

ALLERGAN EQ 0.35% BASE;10,000 UNITS/ML;0.5%

N050081 002

NEOMYCIN SULFATE; PREDNISOLONE ACETATE

OINTMENT;OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/GM;0.25%

A061039 002

EQ 3.5MG BASE/GM;0.5%

A061039 001

SUSPENSION/DROPS;OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/ML;0.25%

A061037 001

**DISCONTINUED DRUG PRODUCT LIST**

6-266(of 375)

\*\* See List Footnote

NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATE

OINTMENT;OPHTHALMIC

NEO-HYDELTRASOL

MERCK

EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE

N050378 001

NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

MYTREX A

SAVAGE LABS

EQ 3.5MG BASE/GM;0.1%

A062598 001 Jul 21, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA

EQ 3.5MG BASE/GM;0.1%

A062600 001 Jul 21, 1986

PHARMADERM

EQ 3.5MG BASE/GM;0.1%

A062595 001 Jul 21, 1986

OINTMENT;TOPICAL

MYTREX A

SAVAGE LABS

EQ 3.5MG BASE/GM;0.1%

A062609 001 May 23, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA

EQ 3.5MG BASE/GM;0.1%

A062608 001 May 23, 1986

PHARMADERM

EQ 3.5MG BASE/GM;0.1%

A062607 001 May 23, 1986

NETILMICIN SULFATE

INJECTABLE;INJECTION

NETROMYCIN

SCHERING

EQ 10MG BASE/ML

N050544 001 Feb 28, 1983

EQ 25MG BASE/ML

N050544 002 Feb 28, 1983

EQ 100MG BASE/ML

N050544 003 Feb 28, 1983

NIACIN

CAPSULE;ORAL

WAMPOCAP

MEDPOINTE PHARM HLC 500MG

N011073 003

TABLET;ORAL

NIACIN

EVERYLIFE

500MG

A083203 001

HALSEY

500MG

A083453 001

HIKMA PHARMS

500MG

A083718 001

IMPAK LABS

500MG

A083115 001

IVAX SUB TEVA PHARMS

500MG

A083180 001

MK LABS

500MG

A083525 001

PUREPAC PHARM

500MG

A083271 001

SANDOZ

500MG

A083306 001

TABLICAPS

500MG

A084237 001

WATSON LABS

500MG

A083136 001

500MG

A083305 001

500MG

A085172 001

NICOLAR

SANOFI AVENTIS US 500MG

A083823 001

TABLET, EXTENDED RELEASE;ORAL

NIASPAN

ABBVIE

375MG

N020381 001 Jul 28, 1997

NIASPAN TITRATION STARTER PACK

ABBVIE

375MG;500MG;750MG

N020381 005 Jul 28, 1997

NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; TYROSINE

SUSPENSION;ORAL

TPN

INTL MINERALS

15MG/5ML;3.75MG/5ML;600MG/5ML

N008378 003

NICARDIPINE HYDROCHLORIDE

CAPSULE;ORAL

CARDENE

CHIESI USA INC

20MG \*\*

N019488 001 Dec 21, 1988

30MG \*\*

N019488 002 Dec 21, 1988

NICARDIPINE HYDROCHLORIDE

WATSON LABS

20MG

A074670 001 Oct 28, 1996

30MG

A074670 002 Oct 28, 1996

CAPSULE, EXTENDED RELEASE;ORAL

CARDENE SR

+ CHIESI USA INC

30MG \*\*

N020005 001 Feb 21, 1992

+

45MG \*\*

N020005 002 Feb 21, 1992

+

60MG \*\*

N020005 003 Feb 21, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-267(of 375)

\*\* See List Footnote

NICLOSAMIDE

TABLET, CHEWABLE;ORAL

NICLOCIDE

BAYER PHARMS

500MG

N018669 001 May 14, 1982

NICOTINE

FILM, EXTENDED RELEASE;TRANSDERMAL

NICOTROL

MCNEIL CONS

15MG/16HR

N020536 001 Jul 03, 1996

PROSTEP

AVEVA

11MG/24HR

N019983 003 Dec 23, 1998

22MG/24HR

N019983 004 Dec 23, 1998

NICOTINE POLACRILEX

GUM, CHEWING;BUCCAL

NICOTINE POLACRILEX

IVAX SUB TEVA PHARMS EQ 2MG BASE

A076880 001 Feb 18, 2009

EQ 4MG BASE

A077850 001 Feb 18, 2009

NIFEDIPINE

CAPSULE;ORAL

ADALAT

BAYER PHARMS

10MG

N019478 001 Nov 27, 1985

20MG

N019478 002 Sep 17, 1986

NIFEDIPINE

CHASE LABS NJ

10MG

A072409 001 Jul 04, 1990

20MG

A073421 001 Jun 19, 1991

TEVA

10MG

A072651 001 Feb 19, 1992

PROCARDIA

+ PFIZER

20MG \*\*

N018482 002 Jul 24, 1986

TABLET, EXTENDED RELEASE;ORAL

NIFEDIPINE

MARTEC USA LLC

90MG

A075414 003 Mar 23, 2004

MYLAN

30MG

A075108 001 Dec 17, 1999

MYLAN LABS LTD

30MG

A090602 001 Sep 13, 2010

60MG

A090602 002 Sep 13, 2010

90MG

A090602 003 Sep 13, 2010

NILUTAMIDE

TABLET;ORAL

NILANDRON

CONCORDIA PHARMS INC 50MG

N020169 001 Sep 19, 1996

NIMODIPINE

CAPSULE;ORAL

NIMOTOP

+ BAYER PHARMS

30MG \*\*

N018869 001 Dec 28, 1988

NISOLDIPINE

TABLET, EXTENDED RELEASE;ORAL

SULAR

+ COVIS PHARMA BV

10MG \*\*

N020356 001 Feb 02, 1995

+

20MG \*\*

N020356 002 Feb 02, 1995

+

25.5MG \*\*

N020356 006 Jan 02, 2008

+

30MG \*\*

N020356 003 Feb 02, 1995

+

40MG \*\*

N020356 004 Feb 02, 1995

NITRIC OXIDE

GAS;INHALATION

INOMAX

+ MALLINCKRODT HOSP

100PPM \*\*

N020845 002 Dec 23, 1999

NITROFURANTOIN

CAPSULE;ORAL

NITROFURANTOIN

WATSON LABS

50MG

A084326 001

100MG

A084326 002

TABLET;ORAL

FURADANTIN

PROCTER AND GAMBLE

50MG

N008693 001

100MG

N008693 002

FURALAN

LANNETT

50MG

A080017 001

**DISCONTINUED DRUG PRODUCT LIST**

6-268(of 375)

\*\* See List Footnote

NITROFURANTOINTABLET;ORAL  
FURALAN

	100MG	A080017 002
NITROFURANTOIN		
ELKINS SINK	50MG	A080003 001
	100MG	A080003 002
IVAX SUB TEVA PHARMS	50MG	A080078 002
	100MG	A080078 001
SANDOZ	50MG	A080043 001
	100MG	A080043 002
WATSON LABS	50MG	A080447 001
	50MG	A085797 001
	100MG	A080447 002
	100MG	A085796 001
WHITEWORTH TOWN PLSN	100MG	A084085 002

NITROFURANTOIN\_SODIUMINJECTABLE;INJECTION  
IVADANTIN  
PROCTER AND GAMBLE EQ 180MG BASE/VIAL

N012402 001

NITROFURANTOIN\_MACROCRYSTALLINECAPSULE;ORAL  
NITROFURANTOIN

MYLAN	100MG	A074967 002 Jul 09, 1997
SANDOZ	25MG	A074336 001 Jan 25, 1995
	50MG	A074336 002 Jan 25, 1995
	100MG	A074336 003 Jan 25, 1995
WATSON LABS	25MG	A073696 001 Dec 31, 1992
	50MG	A073696 002 Dec 31, 1992
	100MG	A073696 003 Dec 31, 1992
NITROFURANTOIN MACROCRYSTALLINE		
WATSON LABS	50MG	A070248 001 Jun 24, 1988
	100MG	A070249 001 Jun 24, 1988

NITROFURANTOIN; NITROFURANTOIN\_MACROCRYSTALLINECAPSULE;ORAL  
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)  
RANBAXY LABS LTD 75MG;25MG

A076951 001 Mar 30, 2005

NITROFURAZONECREAM;TOPICAL  
FURACIN

SHIRE	0.2%	A083789 001
-------	------	-------------

DRESSING;TOPICAL

ACTIN-N		
---------	--	--

SHERWOOD MEDCL 0.2%

N017343 001

OINTMENT;TOPICAL

FURACIN		
---------	--	--

SHIRE 0.2%

N005795 001

## NITROFURAZONE

AMBIX	0.2%	A086077 001
LANNETT	0.2%	A084393 001
PERRIGO NEW YORK	0.2%	A084968 001
TARO	0.2%	A086156 001
WENDT	0.2%	A086766 001

POWDER;TOPICAL

FURACIN		
---------	--	--

SHIRE 0.2%

A083791 001

SOLUTION;TOPICAL

NITROFURAZONE		
---------------	--	--

PERRIGO NEW YORK 0.2%

A085130 001

WENDT 0.2%

A087081 001

NITROGLYCERIN

AEROSOL;SUBLINGUAL

NITROLINGUAL

POHL BOSKAMP 0.4MG/SPRAY

N018705 001 Oct 31, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-269(of 375)

\*\* See List Footnote

**NITROGLYCERIN**

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

KREMERS URBAN PHARMS	0.2MG/HR 0.4MG/HR	A075115 001 Aug 10, 2004 A075115 002 Aug 10, 2004
MYLAN TECHNOLOGIES	0.1MG/HR 0.2MG/HR 0.4MG/HR 0.6MG/HR	A074992 004 Nov 12, 1999 A074992 003 Nov 12, 1999 A074992 002 Nov 12, 1999 A074992 001 Nov 12, 1999
TRANSDERM-NITRO		
+ NOVARTIS	0.1MG/HR **	N020144 001 Feb 27, 1996
+	0.2MG/HR **	N020144 002 Feb 27, 1996
+	0.4MG/HR **	N020144 003 Feb 27, 1996
+	0.6MG/HR **	N020144 004 Feb 27, 1996
+	0.8MG/HR **	N020144 005 Feb 27, 1996
INJECTABLE; INJECTION		
NITRO IV		
POHL BOSKAMP	5MG/ML	N018672 002 Aug 30, 1983
NITRO-BID		
SANOFI AVENTIS US	5MG/ML 10MG/ML	N018621 001 Jan 05, 1982 A071159 001 Feb 28, 1990
NITROGLYCERIN		
ABRAXIS PHARM	5MG/ML 5MG/ML	A070077 001 Dec 13, 1985 A071203 001 May 08, 1987
+ HOSPIRA	5MG/ML **	N018531 001
INTL MEDICATION	5MG/ML	A070026 001 Sep 10, 1985
LUITPOLD	5MG/ML	A071492 001 May 24, 1988
SMITH AND NEPHEW	5MG/ML 5MG/ML	A070633 001 Jun 19, 1986 A070634 001 Jun 19, 1986
NITROGLYCERIN IN DEXTROSE 5%		
HOSPIRA	0.1MG/ML	A074083 001 Oct 26, 1994
NITROL		
RORER	0.8MG/ML	N018774 001 Jan 19, 1983
NITRONAL		
POHL BOSKAMP	1MG/ML	N018672 001 Aug 30, 1983
NITROSTAT		
PARKE DAVIS	0.8MG/ML 5MG/ML 5MG/ML 10MG/ML 10MG/ML	N018588 001 A070863 001 Jan 08, 1987 N018588 002 Dec 23, 1983 A070871 001 Jan 08, 1987 A070872 001 Jan 08, 1987
TRIDIL		
HOSPIRA	0.5MG/ML 5MG/ML	N018537 002 Jun 16, 1983 N018537 001

**NIZATIDINE**

CAPSULE; ORAL

AXID

SMITHKLINE BEECHAM	150MG 300MG	N019508 001 Apr 12, 1988 N019508 002 Apr 12, 1988
NIZATIDINE		
ANI PHARMS INC	150MG 300MG	A075461 001 Jul 08, 2002 A075461 002 Jul 08, 2002
APOTEX INC	150MG 300MG	A076383 001 Jan 23, 2003 A076383 002 Jan 23, 2003
MYLAN PHARMS INC	150MG 300MG	A075934 001 Jul 09, 2002 A075934 002 Jul 09, 2002

SOLUTION; ORAL

AXID

+ BRAINTREE	15MG/ML **	N021494 001 May 25, 2004
-------------	------------	--------------------------

**NONOXYNOL-9**

AEROSOL; VAGINAL

DELFEN

PERSONAL PRODS	12.5%	N014349 002
----------------	-------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-270(of 375)

\*\* See List Footnote

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

METRICS PHARM

EQ 1MG BASE/ML

A040522 001 Sep 30, 2004

NOREPINEPHRINE BITARTRATE; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAIN AND NOVOCAIN W/ LEVOPHED

EASTMAN KODAK

EQ 0.033MG BASE/ML; 2%; 0.4%

N008592 003

NORETHINDRONE

TABLET; ORAL

NORLUTIN

PARKE DAVIS

5MG

N010895 002

NORETHINDRONE ACETATE

TABLET; ORAL

NORLUTATE

PARKE DAVIS

5MG

N012184 002

NORFLOXACIN

SOLUTION/DROPS; OPHTHALMIC

CHIBROXIN

MERCK

0.3%

N019757 001 Jun 17, 1991

TABLET; ORAL

NOROXIN

+ MERCK

400MG \*\*

N019384 002 Oct 31, 1986

NORGESTREL

TABLET; ORAL

OPILL

+ LABORATOIRE HRA

0.075MG

N017031 001

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

AVENTYL HYDROCHLORIDE

LILLY

EQ 10MG BASE

N014684 001

EQ 25MG BASE

N014684 002

NORTRIPTYLINE HYDROCHLORIDE

AUROLIFE PHARMA LLC

EQ 10MG BASE

A074835 001 Jun 30, 1997

EQ 25MG BASE

A074835 002 Jun 30, 1997

EQ 50MG BASE

A074835 003 Jun 30, 1997

EQ 75MG BASE

A074835 004 Jun 30, 1997

IDT AUSTRALIA LTD

EQ 10MG BASE

A074054 001 Dec 31, 1992

EQ 25MG BASE

A074054 002 Dec 31, 1992

EQ 50MG BASE

A074054 003 Dec 31, 1992

EQ 75MG BASE

A074054 004 Dec 31, 1992

MYLAN

EQ 10MG BASE

A074234 001 Jul 26, 1993

EQ 25MG BASE

A074234 002 Jul 26, 1993

EQ 50MG BASE

A074234 003 Jul 26, 1993

EQ 75MG BASE

A074234 004 Jul 26, 1993

TEVA

EQ 10MG BASE

A073667 001 Apr 11, 1996

EQ 25MG BASE

A073667 002 Apr 11, 1996

EQ 50MG BASE

A073667 003 Apr 11, 1996

EQ 75MG BASE

A073667 004 Apr 11, 1996

SOLUTION; ORAL

AVENTYL

+ RANBAXY

EQ 10MG BASE/5ML \*\*

N014685 001

PAMELOR

SPECGX LLC

EQ 10MG BASE/5ML

N018012 001

NYSTATIN

CREAM; TOPICAL

CANDEX

BAYER PHARMS

100,000 UNITS/GM

A061810 001

MYCOSTATIN

DELCOR ASSET CORP

100,000 UNITS/GM \*\*

A060575 001

MYKINAC

ALPHARMA US PHARMS

100,000 UNITS/GM

A062387 001 Jul 29, 1982

NILSTAT

LEDERLE

100,000 UNITS/GM

A061445 001

**DISCONTINUED DRUG PRODUCT LIST**

6-271(of 375)

\*\* See List Footnote

NYSTATIN

CREAM;TOPICAL		
NYSTATIN		
TARO	100,000 UNITS/GM	A062457 001 Jul 28, 1983
LOTION;TOPICAL		
CANDEX		
BAYER PHARMS	100,000 UNITS/ML	N050233 001
OINTMENT;TOPICAL		
MYCOSTATIN		
DELCOR ASSET CORP	100,000 UNITS/GM **	A060571 001
MYKINAC		
ALPHARMA US PHARMS	100,000 UNITS/GM	A062731 001 Sep 22, 1986
NILSTAT		
LEDERLE	100,000 UNITS/GM	A061444 001
PASTILLE;ORAL		
MYCOSTATIN		
DELCOR ASSET CORP	200,000 UNITS	N050619 001 Apr 09, 1987
POWDER;ORAL		
BARSTATIN 100		
BARLAN	100%	A062489 001 Apr 27, 1988
NILSTAT		
+ DAVA PHARMS INC	100% **	N050576 001 Dec 22, 1983
NYSTATIN		
PADDOCK LLC	100%	A062613 001 Nov 26, 1985
POWDER;TOPICAL		
MYCOSTATIN		
DELCOR ASSET CORP	100,000 UNITS/GM **	A060578 001
NYSTATIN		
NESHER PHARMS	100,000 UNITS/GM	A065321 001 Aug 18, 2006
SUPPOSITORY;VAGINAL		
NYSSERT		
WARNER CHILCOTT	100,000 UNITS	N050478 001
SUSPENSION;ORAL		
MYCOSTATIN		
DELCOR ASSET CORP	100,000 UNITS/ML	A061533 001
NILSTAT		
+ GLENMARK GENERICS	100,000 UNITS/ML	N050299 001
NYSTATIN		
ALLIED PHARMA INC	100,000 UNITS/ML	A062832 001 Dec 27, 1991
ALPHARMA US PHARMS	100,000 UNITS/ML	A062571 001 Oct 29, 1985
G AND W LABS INC	100,000 UNITS/ML	A062776 001 Dec 17, 1987
MORTON GROVE	100,000 UNITS/ML	A062835 001 Nov 19, 1987
PHARMADERM	100,000 UNITS/ML	A062518 001 Jul 06, 1984
PHARMAFAIR	100,000 UNITS/ML	A062541 001 Jan 16, 1985
TEVA	100,000 UNITS/ML	A062670 001 Jun 18, 1987
NYSTEX		
SAVAGE LABS	100,000 UNITS/ML	A062519 001 Jul 06, 1984
TABLET;ORAL		
MYCOSTATIN		
DELCOR ASSET CORP	500,000 UNITS	A060574 001
NILSTAT		
LEDERLE	500,000 UNITS	A061151 001
NYSTATIN		
QUANTUM PHARMS	500,000 UNITS	A062525 001 Oct 29, 1984
SANDOZ	500,000 UNITS	A062065 001
USL PHARMA	500,000 UNITS	A062524 001 Nov 26, 1985
WATSON LABS	500,000 UNITS	A062402 001 Dec 16, 1982
TABLET;VAGINAL		
KOROSTATIN		
HOLLAND RANTOS	100,000 UNITS	A061718 001
MYCOSTATIN		
DELCOR ASSET CORP	100,000 UNITS	A060577 001
NILSTAT		
LEDERLE	100,000 UNITS	A061325 001
NYSTATIN		
FOUGERA	100,000 UNITS	A062459 001 Nov 09, 1983
PHARMADERM	100,000 UNITS	A062460 001 Nov 09, 1983
QUANTUM PHARMS	100,000 UNITS	A062509 001 Apr 03, 1984
SANDOZ	100,000 UNITS	A061965 001

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-272(of 375)

\*\* See List Footnote

NYSTATIN

TABLET;VAGINAL

NYSTATIN

TEVA	100,000 UNITS	A062502 001 Dec 23, 1983
WATSON LABS	100,000 UNITS	A062176 001

NYSTATIN; TRIAMCINOLOLONE ACETONIDE

CREAM;TOPICAL

MYCO-TRIACET II

TEVA	100,000 UNITS/GM; 0.1%	A061954 002 Sep 20, 1985
------	------------------------	--------------------------

MYCOLOG-II

DELCOR ASSET CORP	100,000 UNITS/GM; 0.1% **	A060576 002 May 01, 1985
MYLAN PHARMS INC	100,000 UNITS/GM; 0.1% **	A062606 001 May 15, 1985

MYTREX F

SAVAGE LABS	100,000 UNITS/GM; 0.1%	A062597 001 Oct 08, 1985
-------------	------------------------	--------------------------

NYSTATIN AND TRIAMCINOLOLONE ACETONIDE

ALPHARMA US PHARMS	100,000 UNITS/GM; 0.1%	A063010 001 Dec 20, 1988
PERRIGO NEW YORK	100,000 UNITS/GM; 0.1%	A062186 002 Jun 06, 1985
PHARMAFAIR	100,000 UNITS/GM; 0.1%	A062657 001 Jul 30, 1986
TARO	100,000 UNITS/GM; 0.1%	A062347 001 Mar 30, 1987

NYSTATIN TRIAMCINOLOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM; 0.1%	A062596 001 Oct 08, 1985
------------	------------------------	--------------------------

OINTMENT;TOPICAL

MYCO-TRIACET II

TEVA	100,000 UNITS/GM; 0.1%	A062045 002 Nov 26, 1985
------	------------------------	--------------------------

MYCOLOG-II

MYLAN PHARMS INC	100,000 UNITS/GM; 0.1% **	A060572 001 Jun 28, 1985
------------------	---------------------------	--------------------------

MYTREX F

SAVAGE LABS	100,000 UNITS/GM; 0.1%	A062601 001 Oct 09, 1985
-------------	------------------------	--------------------------

NYSTATIN AND TRIAMCINOLOLONE ACETONIDE

PERRIGO NEW YORK	100,000 UNITS/GM; 0.1%	A062280 002 Oct 10, 1985
PHARMAFAIR	100,000 UNITS/GM; 0.1%	A062656 001 Jul 30, 1986

NYSTATIN-TRIAMCINOLOLONE ACETONIDE

PHARMADERM	100,000 UNITS/GM; 0.1%	A062603 001 Oct 09, 1985
------------	------------------------	--------------------------

OCTREOTIDE ACETATE

INJECTABLE;INJECTION

OCTREOTIDE ACETATE

SUN PHARM INDS	EQ 0.05MG BASE/ML	A077329 001 Mar 04, 2008
	EQ 0.1MG BASE/ML	A077329 002 Mar 04, 2008
	EQ 0.2MG BASE/ML	A077330 001 Mar 04, 2008
	EQ 0.5MG BASE/ML	A077329 003 Mar 04, 2008
	EQ 1MG BASE/ML	A077331 001 Mar 04, 2008

WOCKHARDT USA	EQ 0.2MG BASE/ML	A090986 001 May 11, 2011
	EQ 1MG BASE/ML	A090986 002 May 11, 2011

OCTREOTIDE ACETATE PRESERVATIVE FREE

WOCKHARDT USA	EQ 0.05MG BASE/ML	A090985 001 May 11, 2011
	EQ 0.1MG BASE/ML	A090985 002 May 11, 2011
	EQ 0.5MG BASE/ML	A090985 003 May 11, 2011

OFLOXACIN

INJECTABLE;INJECTION

FLOXIN

ORTHO MCNEIL PHARM	20MG/ML	N020087 002 Mar 31, 1992
	40MG/ML	N020087 003 Mar 31, 1992

FLOXIN IN DEXTROSE 5%

ORTHO MCNEIL PHARM	400MG/100ML	N020087 001 Mar 31, 1992
--------------------	-------------	--------------------------

FLOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

ORTHO MCNEIL PHARM	4MG/ML	N020087 004 Mar 31, 1992
	400MG/100ML	N020087 005 Mar 31, 1992

OFLOXACIN

BEDFORD	40MG/ML	A075762 001 Jan 16, 2002
---------	---------	--------------------------

SOLUTION/DROPS;OPHTHALMIC

OFLOXACIN

APOTEX INC	0.3%	A076513 001 May 14, 2004
SANDOZ	0.3%	A076848 001 Nov 25, 2008

SOLUTION/DROPS;OTIC

FLOXIN OTIC

+ DAIICHI	0.3% **	N020799 001 Dec 16, 1997
-----------	---------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-273(of 375)

\*\* See List Footnote

**OFLOXACIN**TABLET;ORAL  
FLOXIN

JANSSEN PHARMS	200MG ** 300MG ** 400MG **	N019735 001 Dec 28, 1990 N019735 002 Dec 28, 1990 N019735 003 Dec 28, 1990
OFLOXACIN		
LARKEN LABS	200MG 300MG	A076093 001 Sep 02, 2003 A076093 002 Sep 02, 2003
RANBAXY LABS LTD	200MG 300MG 400MG	A076220 001 Sep 02, 2003 A076220 002 Sep 02, 2003 A076220 003 Sep 02, 2003

**OLANZAPINE**TABLET;ORAL  
OLANZAPINE

AJANTA PHARMA LTD	2.5MG 5MG 7.5MG 10MG 15MG 20MG	A206711 001 Aug 30, 2016 A206711 002 Aug 30, 2016 A206711 003 Aug 30, 2016 A206711 004 Aug 30, 2016 A206711 005 Aug 30, 2016 A206711 006 Aug 30, 2016
MYLAN PHARMS INC	2.5MG 5MG 7.5MG 10MG 15MG 20MG	A076866 001 Apr 23, 2012 A076866 002 Apr 23, 2012 A076866 003 Apr 23, 2012 A076866 004 Apr 23, 2012 A076866 005 Apr 23, 2012 A076866 006 Apr 23, 2012

**OLIVE OIL; SOYBEAN OIL**

INJECTABLE;INJECTION

CLINOLIPID 20%  
+ BAXTER HLTHCARE CORP 16%(160GM/1000ML);4% (40GM/1000ML)

N204508 001 Oct 03, 2013

**OMEGA-3-ACID ETHYL ESTERS TYPE A**

CAPSULE;ORAL

OMTRYG  
+ OSMOTICA  
1.2GM CONTAINS AT LEAST 900MG OF THE ETHYL ESTERS OF OMEGA-3 FATTY ACIDS

N204977 001 Apr 23, 2014

**OMEPRAZOLE**

CAPSULE, DELAYED REL PELLETS;ORAL

PRILOSEC  
+ ASTRAZENECA PHARMS 10MG \*\*  
+ 20MG \*\*  
+ 40MG \*\*N019810 003 Oct 05, 1995  
N019810 001 Sep 14, 1989  
N019810 002 Jan 15, 1998**ONDANSETRON**TABLET, ORALLY DISINTEGRATING;ORAL  
ONDANSETRON

CHARTWELL MOLECULES	4MG 8MG 16MG 24MG	A076506 001 Dec 26, 2006 A076506 002 Dec 26, 2006 A077406 001 Dec 26, 2006 A077406 002 Dec 26, 2006
NESHER PHARMS	4MG 8MG	A077717 001 Jun 25, 2007 A077717 002 Jun 25, 2007

**ONDANSETRON HYDROCHLORIDE**

INJECTABLE;INJECTION

ONDANSETRON HYDROCHLORIDE  
APOTEX INC EQ 2MG BASE/ML  
HOSPIRA EQ 2MG BASE/ML  
LANNETT EQ 2MG BASE/ML  
EQ 2MG BASE/ML  
LUITPOLD EQ 2MG BASE/ML  
MYLAN LABS LTD EQ 2MG BASE/ML  
PLIVA HRVATSKA DOO EQ 2MG BASE/ML  
SAGENT PHARMS EQ 2MG BASE/ML  
SUN PHARM IND (IN) EQ 2MG BASE/ML  
ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER  
HOSPIRA EQ 0.64MG BASE/MLA077368 001 Dec 26, 2006  
A076695 001 Dec 26, 2006  
A090116 001 Apr 14, 2010  
A090883 001 Aug 05, 2010  
A077582 001 Dec 26, 2006  
A078257 001 Apr 23, 2008  
A077544 001 Dec 26, 2006  
A078180 001 Mar 26, 2007  
A077172 001 Dec 26, 2006  
A076978 001 Feb 26, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-274(of 375)

\*\* See List Footnote

ONDANSETRON HYDROCHLORIDE

## INJECTABLE; INJECTION

## ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

APOTEX INC	EQ 2MG BASE/ML	A077343 001	Dec 26, 2006
HOSPIRA	EQ 2MG BASE/ML	A076696 001	Dec 26, 2006
LUITPOLD	EQ 2MG BASE/ML	A077387 001	Dec 26, 2006
MYLAN LABS LTD	EQ 2MG BASE/ML	A078244 001	Apr 23, 2008
TARO PHARMS IRELAND	EQ 2MG BASE/ML	A078014 001	Mar 21, 2008
ZOFRAN			
+ NOVARTIS PHARMS CORP	EQ 2MG BASE/ML **	N020007 001	Jan 04, 1991
ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER			
+ GLAXOSMITHKLINE	EQ 0.64MG BASE/ML **	N020403 001	Jan 31, 1995
ZOFRAN PRESERVATIVE FREE			
+ NOVARTIS PHARMS CORP	EQ 2MG BASE/ML **	N020007 003	Dec 10, 1993
TABLET; ORAL			
ONDANSETRON HYDROCHLORIDE			
CHARTWELL MOLECULES	EQ 4MG BASE	A077303 001	Jun 25, 2007
	EQ 8MG BASE	A077303 002	Jun 25, 2007
	EQ 24MG BASE	A077303 004	Jun 25, 2007
HIKMA INTL PHARMS	EQ 4MG BASE	A077545 001	Sep 06, 2007
	EQ 8MG BASE	A077545 002	Sep 06, 2007
	EQ 24MG BASE	A077545 003	Sep 06, 2007
TARO	EQ 4MG BASE	A077729 001	Mar 28, 2011
	EQ 8MG BASE	A077729 002	Mar 28, 2011
	EQ 24MG BASE	A077729 003	Mar 28, 2011

ORPHENADRINE CITRATE

## INJECTABLE; INJECTION

## NORFLEX

TELIGENT 30MG/ML N013055 001

## ORPHENADRINE CITRATE

WATSON LABS 30MG/ML A087062 001

## TABLET, EXTENDED RELEASE; ORAL

## NORFLEX

+ MEDICIS 100MG N012157 001

## ORPHENADRINE CITRATE

ASCOT 100MG A088067 001 Apr 06, 1983  
SANDOZ 100MG A085046 001  
WATSON LABS 100MG A084303 001ORPHENADRINE HYDROCHLORIDE

## TABLET; ORAL

## DISIPAL

3M 50MG N010653 001

OSELTAMIVIR PHOSPHATE

## FOR SUSPENSION; ORAL

## TAMIFLU

ROCHE EQ 12MG BASE/ML N021246 001 Dec 14, 2000

OXACILLIN SODIUM

## CAPSULE; ORAL

## BACTOCILL

GLAXOSMITHKLINE EQ 250MG BASE A061336 001  
EQ 250MG BASE A062241 001  
EQ 500MG BASE A061336 002  
EQ 500MG BASE A062241 002

## OXACILLIN SODIUM

ANI PHARMS INC EQ 250MG BASE A062222 001  
EQ 500MG BASE A062222 002

APOTHECON EQ 250MG BASE A061450 002

EQ 500MG BASE A061450 001

## PROSTAPHLIN

APOTHECON EQ 500MG BASE N050118 002

## FOR SOLUTION; ORAL

## BACTOCILL

GLAXOSMITHKLINE EQ 250MG BASE/5ML A062321 001

## OXACILLIN SODIUM

APOTHECON EQ 250MG BASE/5ML A061457 001

TEVA EQ 250MG BASE/5ML A062252 001

**DISCONTINUED DRUG PRODUCT LIST**

6-275(of 375)

\*\* See List Footnote

OXACILLIN SODIUM

FOR SOLUTION;ORAL

PROSTAPHLIN

APOTHECON

EQ 250MG BASE/5ML

N050194 001

INJECTABLE;INJECTION

BACTOCILL

GLAXOSMITHKLINE

EQ 500MG BASE/VIAL \*\*  
EQ 1GM BASE/VIAL \*\*  
EQ 1GM BASE/VIAL \*\*  
EQ 2GM BASE/VIAL \*\*  
EQ 2GM BASE/VIAL \*\*  
EQ 4GM BASE/VIAL \*\*  
EQ 10GM BASE/VIAL \*\*A061334 009 Mar 26, 1982  
A061334 006 Mar 26, 1982  
A062736 001 Dec 19, 1986  
A061334 007 Mar 26, 1982  
A062736 002 Dec 19, 1986  
A061334 008 Mar 26, 1982  
A061334 010

OXACILLIN SODIUM

+ APOTHECON

+

+

+

+

EQ 250MG BASE/VIAL \*\*  
EQ 500MG BASE/VIAL \*\*  
EQ 1GM BASE/VIAL \*\*  
EQ 2GM BASE/VIAL \*\*  
EQ 4GM BASE/VIAL \*\*N050195 001  
N050195 002  
N050195 003  
N050195 004  
N050195 005

ELKINS SINN

EQ 250MG BASE/VIAL  
EQ 500MG BASE/VIAL  
EQ 1GM BASE/VIAL  
EQ 2GM BASE/VIAL  
EQ 4GM BASE/VIAL  
EQ 10GM BASE/VIALA062711 001 Feb 03, 1989  
A062711 002 Feb 03, 1989  
A062711 003 Feb 03, 1989  
A062711 004 Feb 03, 1989  
A062711 005 Feb 03, 1989  
A062711 006 Feb 03, 1989

ISTITUTO BIO ITA SPA

EQ 125MG BASE/VIAL  
EQ 250MG BASE/VIAL  
EQ 500MG BASE/VIAL  
EQ 1GM BASE/VIAL  
EQ 2GM BASE/VIALA062798 003 Dec 11, 1995  
A062798 004 Dec 11, 1995  
A062798 005 Dec 11, 1995  
A062798 001 Dec 11, 1995  
A062798 002 Dec 11, 1995

MYLAN LABS LTD

EQ 1GM BASE/VIAL

A091486 001 Aug 25, 2014

EQ 2GM BASE/VIAL

A091486 002 Aug 25, 2014

SANDOZ

EQ 250MG BASE/VIAL

A061490 001

EQ 500MG BASE/VIAL

A061490 002

WATSON LABS INC

EQ 250MG BASE/VIAL  
EQ 500MG BASE/VIAL  
EQ 1GM BASE/VIAL  
EQ 2GM BASE/VIAL  
EQ 4GM BASE/VIAL  
EQ 10GM BASE/VIALA062856 001 Oct 26, 1988  
A062856 002 Oct 26, 1988  
A062856 003 Oct 26, 1988  
A062856 004 Oct 26, 1988  
A062856 005 Oct 26, 1988  
A062984 001 Sep 29, 1988OXALIPLATIN

INJECTABLE;IV (INFUSION)

ELOXATIN

+ SANOFI AVENTIS US  
+  
+50MG/VIAL \*\*  
100MG/VIAL \*\*  
200MG/40ML (5MG/ML) \*\*N021492 001 Aug 09, 2002  
N021492 002 Aug 09, 2002  
N021759 003 Nov 17, 2006

OXALIPLATIN

SANDOZ

50MG/VIAL

A090849 001 Apr 28, 2011

100MG/VIAL

A090849 002 Apr 28, 2011

SANDOZ INC

50MG/10ML (5MG/ML)  
100MG/20ML (5MG/ML)A078812 001 Aug 07, 2009  
A078812 002 Aug 07, 2009OXAMNIQUINE

CAPSULE;ORAL

VANSIL

PFIZER

250MG

N018069 001

OXANDROLONE

TABLET;ORAL

OXANDROLONE

ROXANE

2.5MG

A077249 001 Jul 10, 2007

10MG

A077249 002 Jul 10, 2007

SANDOZ

2.5MG

A076897 001 Dec 01, 2006

10MG

A076897 002 Dec 01, 2006

**DISCONTINUED DRUG PRODUCT LIST**

6-276(of 375)

\*\* See List Footnote

**OXAPROZIN**TABLET;ORAL  
OXAPROZIN

ACTAVIS ELIZABETH	600MG	A075843 001 Oct 03, 2001
MYLAN	600MG	A075851 001 Aug 17, 2001
MYLAN PHARMS INC	600MG	A075847 001 Feb 28, 2001
SANDOZ	600MG	A075842 001 Apr 12, 2001
	600MG	A075850 001 Apr 27, 2001
WATSON LABS	600MG	A075848 001 Feb 09, 2001

**OXAPROZIN POTASSIUM**TABLET;ORAL  
DAYPRO ALTA  
GD SEARLE

600MG N020776 001 Oct 17, 2002

**OXAZEPAM**CAPSULE;ORAL  
OXAZEPAM

AM THERAP	10MG	A071955 001 Mar 03, 1988
	15MG	A071956 001 Mar 03, 1988
	30MG	A071957 001 Mar 03, 1988
ANDA REPOSITORY	10MG	A071026 002 Aug 10, 1987
	15MG	A071026 003 Aug 10, 1987
	30MG	A071026 001 Aug 10, 1987
IVAX SUB TEVA PHARMS	10MG	A070943 001 Aug 03, 1987
	15MG	A070944 001 Aug 03, 1987
	30MG	A070945 001 Aug 03, 1987
MYLAN	10MG	A071713 001 Oct 20, 1987
	15MG	A071714 001 Oct 20, 1987
	30MG	A071715 001 Oct 20, 1987
WATSON LABS	15MG	A072953 001 Sep 28, 1990
	30MG	A072954 001 Sep 28, 1990
WATSON LABS TEVA	10MG	A072952 001 Sep 28, 1990
SERAX		
ALPHARMA US PHARMS	10MG **	N015539 002
	15MG **	N015539 004
	30MG **	N015539 006
ZAXOPAM		
QUANTUM PHARMICS	10MG	A070650 001 Mar 01, 1988
	15MG	A070640 001 Mar 01, 1988
	30MG	A070641 001 Mar 01, 1988

TABLET;ORAL

OXAZEPAM

PARKE DAVIS	15MG	A071508 001 Feb 02, 1987
SUN PHARM INDUSTRIES	15MG	A070683 001 Jan 16, 1987
WATSON LABS	15MG	A071494 001 Apr 21, 1987

SERAX

ALPHARMA US PHARMS 15MG \*\*

N015539 008

**OXCARBAZEPINE**TABLET;ORAL  
OXCARBAZEPINE

JUBILANT CADISTA	150MG	A090239 001 Jan 25, 2010
	300MG	A090239 002 Jan 25, 2010
	600MG	A090239 003 Jan 25, 2010

**OXPRENOLOL HYDROCHLORIDE**CAPSULE;ORAL  
TRASICOR

NOVARTIS	20MG	N018166 001 Dec 28, 1983
	40MG	N018166 002 Dec 28, 1983
	80MG	N018166 003 Dec 28, 1983
	160MG	N018166 004 Dec 28, 1983

**OXTRIPHYLLINE**

SOLUTION;ORAL

CHOLEDYL

PARKE DAVIS 100MG/5ML

N009268 012 Nov 27, 1984

OXTRIPHYLLINE

MORTON GROVE 100MG/5ML

A088243 001 Dec 05, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-277(of 375)

\*\* See List Footnote

**OXTRIPHYLLINE**

SYRUP;ORAL CHOLEDYL PARKE DAVIS	50MG/5ML	N009268 011
OXTRIPHYLLINE PEDIATRIC MORTON GROVE	50MG/5ML	A088242 001 Dec 05, 1983
TABLET, DELAYED RELEASE;ORAL CHOLEDYL PARKE DAVIS	100MG 200MG	N009268 003 N009268 007
OXTRIPHYLLINE WATSON LABS	100MG 200MG	A087866 001 Aug 25, 1983 A087835 001 Aug 25, 1983
TABLET, EXTENDED RELEASE;ORAL CHOLEDYL SA WARNER CHILCOTT LLC	600MG	A086742 001

**OXYBUTYNIN**

GEL, METERED;TRANSDERMAL GELNIQUE 3% + ALLERGAN SALES LLC	3%	N202513 001 Dec 07, 2011
---	----	--------------------------

**OXYBUTYNIN CHLORIDE**

SYRUP;ORAL DITROPAN + ORTHO MCNEIL JANSSEN	5MG/5ML **	N018211 001
OXYBUTYNIN CHLORIDE APOTEX INC	5MG/5ML	A074997 001 Oct 15, 1997
MIKART	5MG/5ML	A075039 001 Jan 29, 1999
TABLET;ORAL DITROPAN + JANSSEN PHARMS	5MG **	N017577 001
OXYBUTYNIN CHLORIDE QUANTUM PHARMICS	5MG	A072296 001 Dec 08, 1988
USL PHARMA	5MG	A070746 001 Mar 10, 1988
WATSON LABS	5MG	A072485 001 Apr 19, 1989

**OXYCODONE HYDROCHLORIDE**

TABLET, EXTENDED RELEASE;ORAL ROXICODONE ROXANE	10MG 30MG	N020932 001 Oct 26, 1998 N020932 002 Oct 26, 1998
---	--------------	--

**OXYMETAZOLINE HYDROCHLORIDE**

SOLUTION/DROPS;OPHTHALMIC OCUCLEAR BAYER HEALTHCARE LLC	0.025%	N018471 001 May 30, 1986
---	--------	--------------------------

**OXYMORPHONE HYDROCHLORIDE**

INJECTABLE;INJECTION OPANA + ENDO PHARMS	1MG/ML 1.5MG/ML	N011707 002 N011707 001
SUPPOSITORY;RECTAL NUMORPHAN ENDO PHARMS	5MG	N011738 004
TABLET, EXTENDED RELEASE;ORAL OPANA ER ENDO PHARMS	5MG ** 5MG 7.5MG **	N021610 001 Jun 22, 2006 N201655 001 Dec 09, 2011 N021610 005 Feb 29, 2008
+ +	7.5MG 7.5MG	N021655 002 Dec 09, 2011
+ +	10MG ** 10MG 15MG **	N021610 002 Jun 22, 2006 N201655 003 Dec 09, 2011 N021610 006 Feb 29, 2008
+ +	15MG 20MG **	N021655 004 Dec 09, 2011 N021610 003 Jun 22, 2006
+ +	20MG 30MG **	N201655 005 Dec 09, 2011 N021610 007 Feb 29, 2008
+ +	30MG 40MG **	N201655 006 Dec 09, 2011 N021610 004 Jun 22, 2006
+ +	40MG	N201655 007 Dec 09, 2011

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-278(of 375)

\*\* See List Footnote

OXYMORPHONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE;ORAL  
OXYMORPHONE HYDROCHLORIDE

PAR PHARM	5MG	A200792 001	Oct 24, 2014
	7.5MG	A200792 002	Oct 24, 2014
	10MG	A200792 003	Oct 24, 2014
	15MG	A200792 004	Oct 24, 2014
	20MG	A200792 005	Oct 24, 2014
	30MG	A200792 006	Oct 24, 2014
	40MG	A200792 007	Oct 24, 2014

OXPHENBUTAZONE

TABLET;ORAL

OXYPHENBUTAZONE

WATSON LABS	100MG	A088399 001	Sep 17, 1984
TANDEARIL	100MG	N012542 004	Sep 03, 1982

NOVARTIS

OXPHENCYCLIMINE HYDROCHLORIDE

TABLET;ORAL

DARICON

PFIZER 10MG

N011612 001

OXPHENONIUM BROMIDE

TABLET;ORAL

ANTRENYL

NOVARTIS 5MG

N008492 002

OXYTETRACYCLINE

TABLET;ORAL

TERRAMYCIN

PFIZER 250MG

N050287 001

OXYTETRACYCLINE CALCIUM

SYRUP;ORAL

TERRAMYCIN

PFIZER EQ 125MG BASE/5ML

A060595 001

OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE;ORAL

OXY-KESSO-TETRA

FERRANTE EQ 250MG BASE

A060179 001

OXYTETRACYCLINE HYDROCHLORIDE

HIKMA PHARMS EQ 250MG BASE

A060770 001

IMPAK LABS EQ 250MG BASE

A060760 001

PROTER EQ 250MG BASE

A060869 001

PUREPAC PHARM EQ 250MG BASE

A060634 001

TERRAMYCIN

PFIZER EQ 125MG BASE

N050286 001

EQ 250MG BASE

N050286 002

INJECTABLE;INJECTION

TERRAMYCIN

PFIZER EQ 250MG BASE/VIAL

A060586 001

EQ 500MG BASE/VIAL

A060586 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT;OTIC

TERRAMYCIN W/ POLYMYXIN

PFIZER EQ 5MG BASE/GM;10,000 UNITS/GM

A061841 001

TABLET;VAGINAL

TERRAMYCIN-POLYMYXIN

PFIZER EQ 100MG BASE;100,000 UNITS

A061009 001

OXYTOCIN

INJECTABLE;INJECTION

OXYTOCIN

TEVA PHARMS USA 10USP UNITS/ML (10USP UNITS/ML)

A077453 001 Jan 24, 2008

100USP UNITS/10ML (10USP UNITS/ML)

A077453 002 Jan 24, 2008

OXYTOCIN 10 USP UNITS IN DEXTROSE 5%

+ ABBOTT 1USP UNITS/100ML \*\*

N019185 004 Mar 29, 1985

+ 2USP UNITS/100ML \*\*

N019185 003 Mar 29, 1985

OXYTOCIN 20 USP UNITS IN DEXTROSE 5%

+ ABBOTT 2USP UNITS/100ML \*\*

N019185 002 Mar 29, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-279(of 375)

\*\* See List Footnote

OXYTOCIN

INJECTABLE; INJECTION

OXYTOCIN 5 USP UNITS IN DEXTROSE 5%	1USP UNITS/100ML **	N019185 001 Mar 29, 1985
+ ABBOTT		
SYNTOCINON		
NOVARTIS	10USP UNITS/ML	N018245 001
SOLUTION; NASAL		
SYNTOCINON		
RTRX	40USP UNITS/ML	N012285 001

PACLITAXEL

INJECTABLE; INJECTION

PACLITAXEL		
ACCORD HLTHCARE	6MG/ML	A075436 001 Nov 12, 2004
HOSPIRA	6MG/ML	A076233 001 Aug 01, 2002
MYLAN	6MG/ML	A075278 001 Jan 25, 2002
PLIVA LACHEMA	6MG/ML	A077413 001 Mar 12, 2008
TEVA PHARMS USA	6MG/ML	A075297 001 Jan 25, 2002

PALIPERIDONE

TABLET, EXTENDED RELEASE; ORAL

INVEGA		
+ JANSSEN PHARMS	12MG **	N021999 004 Dec 19, 2006

PALONOSETRON HYDROCHLORIDE

CAPSULE; ORAL

ALOXI		
+ HELSINN HLTHCARE	EQ 0.5MG BASE **	N022233 001 Aug 22, 2008
SOLUTION; INTRAVENOUS		
PALONOSETRON HYDROCHLORIDE		
DR REDDYS LABS LTD	EQ 0.075MG BASE/1.5ML (EQ 0.05MG BASE/ML)	N203050 001 Mar 01, 2016
	EQ 0.25MG BASE/5ML (EQ 0.05MG BASE/ML)	N203050 002 Mar 01, 2016

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

AREDIA		
+ NOVARTIS	30MG/VIAL **	N020036 001 Oct 31, 1991
	60MG/VIAL	N020036 003 May 06, 1993
	90MG/VIAL	N020036 004 May 06, 1993
PAMIDRONATE DISODIUM		
AESGEN	30MG/VIAL	A075594 001 May 06, 2002
	90MG/VIAL	A075594 002 May 06, 2002
MN PHARMS	30MG/VIAL	A078300 001 Mar 10, 2009
	90MG/VIAL	A078300 002 Mar 10, 2009

PANCRELIPASE (AMYLASE; LIPASE; PROTEASE)

CAPSULE; ORAL

COTAZYM		
ORGANON USA INC	30,000USP UNITS; 8,000USP UNITS; 30,000USP UNITS	N020580 001 Dec 09, 1996

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM BROMIDE		
ELKINS SINK	1MG/ML	A072058 001 Mar 23, 1988
	2MG/ML	A072059 001 Mar 23, 1988
	2MG/ML	A072060 001 Mar 23, 1988
HOSPIRA	2MG/ML	A072321 001 Jan 19, 1989
IGI LABS INC	1MG/ML	A072210 001 Mar 31, 1988
	2MG/ML	A072211 001 Mar 31, 1988
	2MG/ML	A072212 001 Mar 31, 1988
	2MG/ML	A072213 001 Mar 31, 1988
PAVULON		
+ ORGANON USA INC	1MG/ML	N017015 002
+	2MG/ML	N017015 001

**DISCONTINUED DRUG PRODUCT LIST**

6-280(of 375)

\*\* See List Footnote

PANTOPRAZOLE SODIUMTABLET, DELAYED RELEASE;ORAL  
PANTOPRAZOLE SODIUMSUN PHARM INDS LTD EQ 20MG BASE A077058 001 Sep 10, 2007  
EQ 40MG BASE A077058 002 Sep 10, 2007PARAMETHADIONECAPSULE;ORAL  
PARADIONEABBVIE 150MG N006800 003  
300MG N006800 001

SOLUTION;ORAL

PARADIONE

ABBVIE 300MG/ML N006800 002

PARAMETHASONE ACETATETABLET;ORAL  
HALDRONELILLY 1MG N012772 005  
2MG N012772 006PARGYLINE HYDROCHLORIDETABLET;ORAL  
EUTONYLABBOTT 10MG N013448 002  
25MG N013448 003  
50MG N013448 004PARICALCITOLCAPSULE;ORAL  
ZEMPLAR

+ ABBVIE 4MCG \*\* N021606 003 May 26, 2005

PAROMOMYCIN SULFATECAPSULE;ORAL  
HUMATINKING PFIZER EQ 250MG BASE A062310 001  
PARKADEALE EQ 250MG BASE A060521 001

SYRUP;ORAL

HUMATIN

PARKE DAVIS EQ 125MG BASE/5ML A060522 001

PAROXETINE HYDROCHLORIDECAPSULE;ORAL  
PAXIL+ APOTEX TECHNOLOGIES EQ 10MG BASE \*\* N020885 001 Oct 09, 1998  
+ EQ 20MG BASE \*\* N020885 002 Oct 09, 1998  
+ EQ 30MG BASE \*\* N020885 003 Oct 09, 1998  
+ EQ 40MG BASE \*\* N020885 004 Oct 09, 1998

SUSPENSION;ORAL

PAROXETINE HYDROCHLORIDE

APOTEX INC EQ 10MG BASE/5ML A077395 001 Dec 05, 2006

TABLET;ORAL

PAROXETINE HYDROCHLORIDE

MYLAN PHARMS INC EQ 10MG BASE A075716 001 Mar 08, 2004  
EQ 20MG BASE A075716 002 Mar 08, 2004  
EQ 30MG BASE A075716 003 Mar 08, 2004  
EQ 40MG BASE A075716 004 Mar 08, 2004ROXANE EQ 10MG BASE A078026 001 Jun 29, 2007  
EQ 20MG BASE A078026 002 Jun 29, 2007  
EQ 30MG BASE A078026 003 Jun 29, 2007  
EQ 40MG BASE A078026 004 Jun 29, 2007TEVA PHARMS EQ 10MG BASE A077082 001 Jun 29, 2007  
EQ 20MG BASE A077082 002 Jun 29, 2007  
EQ 30MG BASE A077082 003 Jun 29, 2007  
EQ 40MG BASE A077082 004 Jun 29, 2007UPSHER-SMITH LABS EQ 10MG BASE A075566 001 Mar 08, 2004  
EQ 20MG BASE A075566 002 Mar 08, 2004  
EQ 30MG BASE A075566 003 Mar 08, 2004  
EQ 40MG BASE A075566 004 Mar 08, 2004

PAXIL APOTEX TECHNOLOGIES EQ 50MG BASE N020031 004 Dec 29, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-281(of 375)

\*\* See List Footnote

PAZOPANIB HYDROCHLORIDETABLET;ORAL  
VOTRIENT

NOVARTIS PHARMS CORP EQ 400MG BASE

N022465 002 Oct 19, 2009

PEGINESATIDE ACETATESOLUTION;INTRAVENOUS, SUBCUTANEOUS  
OMONTYS

TAKEDA PHARMS USA	EQ 10MG BASE/ML (EQ 10MG BASE/ML)	N202799 007 Mar 27, 2012
	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N202799 008 Mar 27, 2012
OMONTYS PRESERVATIVE FREE		
TAKEDA PHARMS USA	EQ 1MG BASE/0.5ML (EQ 1MG BASE/0.5ML)	N202799 001 Mar 27, 2012
	EQ 2MG BASE/0.5ML (EQ 2MG BASE/0.5ML)	N202799 002 Mar 27, 2012
	EQ 3MG BASE/0.5ML (EQ 3MG BASE/0.5ML)	N202799 003 Mar 27, 2012
	EQ 4MG BASE/0.5ML (EQ 4MG BASE/0.5ML)	N202799 004 Mar 27, 2012
	EQ 5MG BASE/0.5ML (EQ 5MG BASE/0.5ML)	N202799 005 Mar 27, 2012
	EQ 6MG BASE/0.5ML (EQ 6MG BASE/0.5ML)	N202799 006 Mar 27, 2012

PEMIROLAST POTASSIUMSOLUTION/DROPS;OPHTHALMIC  
ALAMAST  
SANTEN

0.1%

N021079 001 Sep 24, 1999

PEMOLINETABLET;ORAL  
CYLERT

ABBOTT	18.75MG	N016832 001
	37.5MG	N016832 002
	75MG	N016832 003

## PEMOLINE

ACTAVIS ELIZABETH	18.75MG	A075595 001 Feb 28, 2000
	37.5MG	A075595 002 Feb 28, 2000
	75MG	A075595 003 Feb 28, 2000
MALLINCKRODT	18.75MG	A075726 003 Mar 30, 2001
	37.5MG	A075726 002 Mar 30, 2001
	75MG	A075726 001 Mar 30, 2001
SANDOZ	18.75MG	A075286 001 Dec 27, 1999
	37.5MG	A075286 002 Jun 30, 1999
	75MG	A075286 003 Jun 30, 1999
TEVA PHARMS	18.75MG	A075030 003 Feb 22, 2000
	37.5MG	A075030 001 Jan 29, 1999
	75MG	A075030 002 Jan 29, 1999
VINTAGE PHARMS	18.75MG	A075328 001 Apr 19, 2000
	37.5MG	A075328 002 Apr 19, 2000
	75MG	A075328 003 Apr 19, 2000
WATSON LABS	18.75MG	A075287 001 Jun 13, 2001
	37.5MG	A075287 002 Sep 18, 2000
	75MG	A075287 003 Sep 18, 2000

TABLET, CHEWABLE;ORAL

## CYLERT

ABBOTT	37.5MG	N017703 001
PEMOLINE		
ACTAVIS ELIZABETH	37.5MG	A075678 001 Jul 26, 2000

TEVA PHARMS 37.5MG A075555 001 Feb 18, 2000

PENBUTOLOL SULFATETABLET;ORAL  
LEVATOL

+ AUXILIUM PHARMS LLC	10MG **	N018976 001 Dec 30, 1987
+	20MG **	N018976 004 Jan 05, 1989

PENICILLAMINECAPSULE;ORAL  
CUPRIMINE  
ATON

125MG

N019853 002

**DISCONTINUED DRUG PRODUCT LIST**

6-282(of 375)

\*\* See List Footnote

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION

BICILLIN L-A

WYETH AYERST

300,000 UNITS/ML

N050131 001

SUSPENSION; ORAL

BICILLIN

WYETH AYERST

300,000 UNITS/5ML

N050126 002

TABLET; ORAL

BICILLIN

WYETH AYERST

200,000 UNITS

N050128 001

PENICILLIN G POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN

TEVA

200,000 UNITS/5ML

A060307 002

400,000 UNITS/5ML

A060307 004

PENICILLIN G POTASSIUM

MYLAN

200,000 UNITS/5ML

A060752 003

250,000 UNITS/5ML

A060752 002

400,000 UNITS/5ML

A060752 001

PUREPAC PHARM

250,000 UNITS/5ML

A061740 001

400,000 UNITS/5ML

A061740 002

PENICILLIN-2

TEVA

250,000 UNITS/5ML

A060307 003

PENTIDS '200'

APOTHECON

200,000 UNITS/5ML

A062149 001

PENTIDS '400'

APOTHECON

400,000 UNITS/5ML

A062149 002

PFIZERPEN G

PFIZER

400,000 UNITS/5ML

A060587 001

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

APOTHECON

1,000,000 UNITS/VIAL

A060362 001

5,000,000 UNITS/VIAL

A060362 003

10,000,000 UNITS/VIAL

A060362 004

20,000,000 UNITS/VIAL

A060362 002

CONSOLIDATED PHARM

500,000 UNITS/VIAL

A060806 001

1,000,000 UNITS/VIAL

A060806 002

5,000,000 UNITS/VIAL

A060806 003

10,000,000 UNITS/VIAL

A060806 004

LILLY

200,000 UNITS/VIAL

A060384 004

500,000 UNITS/VIAL

A060384 003

1,000,000 UNITS/VIAL

A060384 002

5,000,000 UNITS/VIAL

A060384 001

20,000,000 UNITS/VIAL

A060384 005

20,000,000 UNITS/VIAL

A060601 001

PARKE DAVIS

1,000,000 UNITS/VIAL

A062003 001

5,000,000 UNITS/VIAL

A062003 002

PFIZER

20,000,000 UNITS/VIAL

A060074 003

SANDOZ

1,000,000 UNITS/VIAL \*\*

A065079 001 Aug 30, 2002

WATSON LABS INC

1,000,000 UNITS/VIAL

A062991 001 Sep 13, 1988

5,000,000 UNITS/VIAL

A062991 002 Sep 13, 1988

10,000,000 UNITS/VIAL

A062991 003 Sep 13, 1988

20,000,000 UNITS/VIAL

A062991 004 Sep 13, 1988

PFIZERPEN

PFIZER

1,000,000 UNITS/VIAL \*\*

A060657 001

TABLET; ORAL

PENICILLIN G POTASSIUM

APOTHECON

250,000 UNITS

A060392 003

IVAX SUB TEVA PHARMS

400,000 UNITS

A060073 004

LILLY

250,000 UNITS

A060403 001

MYLAN

200,000 UNITS

A060781 001

250,000 UNITS

A060781 002

400,000 UNITS

A060781 003

500,000 UNITS

A060781 005

800,000 UNITS

A060781 004

PUREPAC PHARM

200,000 UNITS

A061588 001

250,000 UNITS

A061588 002

400,000 UNITS

A061588 003

TEVA

200,000 UNITS

A060306 001

**DISCONTINUED DRUG PRODUCT LIST**

6-283(of 375)

\*\* See List Footnote

PENICILLIN G POTASSIUM

TABLET;ORAL

PENICILLIN G POTASSIUM

	250,000 UNITS	A060306 002
	400,000 UNITS	A060306 003
	500,000 UNITS	A060306 004
WYETH AYERST	200,000 UNITS	A060413 001
	250,000 UNITS	A060413 002
	400,000 UNITS	A060413 003
PENTIDS '200'		
APOTHECON	200,000 UNITS	A062155 001
PENTIDS '250'		
APOTHECON	250,000 UNITS	A062155 002
PENTIDS '400'		
APOTHECON	400,000 UNITS	A060392 004
	400,000 UNITS	A062155 003
PENTIDS '800'		
APOTHECON	800,000 UNITS	A060392 005
	800,000 UNITS	A062155 004
PFIZERPEN G		
PFIZER	50,000 UNITS	A060075 001
	100,000 UNITS	A060075 002
	200,000 UNITS	A060075 003
	250,000 UNITS	A060075 004
	400,000 UNITS	A060075 005
	800,000 UNITS	A060075 006

PENICILLIN G PROCAINE

INJECTABLE;INJECTION

DURACILLIN A.S.

LILLY	300,000 UNITS/ML	A060093 001
PENICILLIN G PROCAINE		
CONSOLIDATED PHARM	300,000 UNITS/ML	A060800 001
	600,000 UNITS/1.2ML	A060800 002
PARKE DAVIS	300,000 UNITS/ML	A062029 001
PFIZER	300,000 UNITS/VIAL	A060099 001
	1,500,000 UNITS/VIAL	A060099 002
PFIZERPEN-AS		
PFIZER	300,000 UNITS/ML	A060286 001
	600,000 UNITS/ML	A060286 002

PENICILLIN G SODIUM

INJECTABLE;INJECTION

PENICILLIN G SODIUM

BRISTOL MYERS SQUIBB	5,000,000 UNITS/VIAL	A061935 001
COPANOS	5,000,000 UNITS/VIAL	A061051 001
PHARMACIA AND UPJOHN	1,000,000 UNITS/VIAL	A061046 001

INJECTABLE;INTRAMUSCULAR, INTRAVENOUS

PENICILLIN G SODIUM

WATSON LABS INC	5,000,000 UNITS/VIAL	A063014 001 Sep 13, 1988
-----------------	----------------------	--------------------------

PENICILLIN V

FOR SUSPENSION;ORAL

V-CILLIN

LILLY	125MG/0.6ML	A060002 001
-------	-------------	-------------

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

BEEPEN-VK

GLAXOSMITHKLINE	EQ 125MG BASE/5ML	A062270 001
	EQ 250MG BASE/5ML	A062270 002

BETAPEN-VK

APOTHECON	EQ 125MG BASE/5ML	A061149 001
	EQ 250MG BASE/5ML	A061149 002

LEDERCILLIN VK

LEDERLE	EQ 125MG BASE/5ML	A060136 001
	EQ 250MG BASE/5ML	A060136 002

PEN-VEE K

WYETH AYERST	EQ 125MG BASE/5ML	A060007 001
	EQ 250MG BASE/5ML	A060007 002

**DISCONTINUED DRUG PRODUCT LIST**

6-284(of 375)

\*\* See List Footnote

PENICILLIN V POTASSIUM

FOR SOLUTION;ORAL

PENAPAR-VK

PARKE DAVIS

EQ 125MG BASE/5ML

A062002 001

EQ 250MG BASE/5ML

A062002 002

PENICILLIN V POTASSIUM

AM ANTIBIOTICS

EQ 125MG BASE/5ML

A061529 001

EQ 250MG BASE/5ML

A061529 002

MYLAN

EQ 125MG BASE/5ML

A061624 002

EQ 250MG BASE/5ML

A061624 001

PUREPAC PHARM

EQ 125MG BASE/5ML

A061758 001

EQ 250MG BASE/5ML

A061758 002

PFIZERPEN VK

PFIZER

EQ 125MG BASE/5ML

A061815 001

EQ 250MG BASE/5ML

A061815 002

V-CILLIN K

LILLY

EQ 125MG BASE/5ML

A060004 001

EQ 250MG BASE/5ML

A060004 002

VEETIDS

APOTHECON

EQ 125MG BASE/5ML

A061410 001

EQ 250MG BASE/5ML

A061410 002

VEETIDS '125'

APOTHECON

EQ 125MG BASE/5ML

A061206 001

EQ 125MG BASE/5ML

A062153 001

VEETIDS '250'

APOTHECON

EQ 250MG BASE/5ML

A061206 002

EQ 250MG BASE/5ML

A062153 002

TABLET;ORAL

BEEPEN-VK

GLAXOSMITHKLINE

EQ 250MG BASE

A062273 001

EQ 500MG BASE

A062273 002

BETAPEN-VK

BRISTOL

EQ 250MG BASE

A061150 001

EQ 500MG BASE

A061150 002

LEDERCILLIN VK

LEDERLE

EQ 250MG BASE

A060134 001

EQ 500MG BASE

A060134 002

PEN-VEE K

WYETH AYERST

EQ 125MG BASE

A060006 001

EQ 250MG BASE

A060006 002

EQ 500MG BASE

A060006 003

PENAPAR-VK

PARKE DAVIS

EQ 250MG BASE

A062001 001

EQ 500MG BASE

A062001 002

PENICILLIN V POTASSIUM

AM ANTIBIOTICS

EQ 250MG BASE

A061528 001

EQ 500MG BASE

A061528 002

IVAX SUB TEVA PHARMS

EQ 125MG BASE

A060518 001

EQ 250MG BASE

A060518 002

EQ 500MG BASE

A060518 003

MYLAN

EQ 250MG BASE

A061530 001

EQ 500MG BASE

A061530 002

PUREPAC PHARM

EQ 125MG BASE

A061571 001

EQ 250MG BASE

A061571 002

EQ 500MG BASE

A061571 003

PFIZERPEN VK

PFIZER

EQ 250MG BASE

A061836 001

EQ 500MG BASE

A061836 002

UTICILLIN VK

PHARMACIA AND UPJOHN

EQ 250MG BASE

A061651 001

EQ 500MG BASE

A061651 002

V-CILLIN K

LILLY

EQ 125MG BASE \*\*

A060003 001

EQ 250MG BASE \*\*

A060003 002

EQ 500MG BASE \*\*

A060003 003

VEETIDS

APOTHECON

EQ 250MG BASE

A061411 001

EQ 500MG BASE

A061411 002

**DISCONTINUED DRUG PRODUCT LIST**

6-285(of 375)

\*\* See List Footnote

PENICILLIN V POTASSIUM

TABLET;ORAL VEETIDS '250'	APOTHECON	EQ 250MG BASE EQ 250MG BASE	A061164 001 A062156 002
VEETIDS '500'	APOTHECON	EQ 500MG BASE EQ 500MG BASE	A061164 002 A062156 001

PENTAGASTRIN

INJECTABLE;INJECTION PEPTAVLON + WYETH AYERST		0.25MG/ML **	N017048 001
---	--	--------------	-------------

PENTAMIDINE ISETHIONATE

FOR SOLUTION;INHALATION NEBUPENT	FRESENIUS KABI USA	600MG/VIAL	N019887 002 Mar 22, 1996
INJECTABLE;INJECTION PENTACARINAT	ARMOUR PHARM	300MG/VIAL	A073447 001 Apr 28, 1994
PENTAMIDINE ISETHIONATE	BAXTER HLTHCARE	300MG/VIAL	A073617 001 Dec 18, 1995
HOSPIRA	300MG/VIAL	A073479 001 Jun 30, 1992	
WATSON LABS	300MG/VIAL	A074303 001 Aug 17, 1995	

PENTAZOCINE HYDROCHLORIDE

TABLET;ORAL TALWIN 50	SANOFI AVENTIS US	EQ 50MG BASE	N016732 001
--------------------------	-------------------	--------------	-------------

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE;INJECTION YTTERBIUM YB 169 DTPA	3M	2mCi/ML	N017518 001
---	----	---------	-------------

PENTOBARBITAL

ELIXIR;ORAL NEMBUTAL	OAK PHARMS	18.2MG/5ML	A083244 001
-------------------------	------------	------------	-------------

PENTOBARBITAL SODIUM

CAPSULE;ORAL NEMBUTAL SODIUM	OAK PHARMS	30MG 50MG 100MG	A084095 001 A084093 001 A083245 001
PENTOBARBITAL SODIUM	LANNETT	50MG 100MG	A085937 001 A085915 001
VITARINE	100MG	A083284 001	
WHITEWORTH TOWN PLSN	100MG	A083338 001	
SODIUM PENTOBARBITAL			
ANABOLIC	100MG	A084590 001	
ELKINS SINN	100MG	A083368 001	
EVERYLIFE	100MG	A083259 001	
HALSEY	100MG	A084677 001	
IVAX SUB TEVA PHARMS	50MG 100MG	A083461 001 A083461 002	
PARKE DAVIS	100MG	A084156 001	
PERRIGO	100MG	A084560 001	
PUREPAC PHARM	100MG	A083301 001	
VALEANT PHARM INTL	100MG	A083264 001	
WATSON LABS	100MG	A085791 001	
WYETH AYERST	100MG	A083239 001	

## INJECTABLE;INJECTION

PENTOBARBITAL SODIUM	ELKINS SINN	50MG/ML	A083270 001
SODIUM PENTOBARBITAL	WYETH AYERST	50MG/ML	A083261 001

**DISCONTINUED DRUG PRODUCT LIST**

6-286(of 375)

\*\* See List Footnote

PENTOBARBITAL SODIUMSUPPOSITORY;RECTAL  
NEMBUTAL

OAK PHARMS	30MG	A083247 001	Jan 25, 1982
	60MG	A083247 002	Jan 25, 1982
	120MG	A083247 003	Jan 25, 1982
	200MG	A083247 004	Jan 25, 1982

TABLET;ORAL

PENTOBARBITAL SODIUM			
VITARINE	100MG	A083285 001	
SODIUM PENTOBARBITAL			
NEXGEN PHARMA INC	100MG	A084238 001	

PENTOLINIUM TARTRATEINJECTABLE;INJECTION  
ANSOLYSEN

WYETH AYERST	10MG/ML	N009372 001	
--------------	---------	-------------	--

PENTOXIFYLLINETABLET, EXTENDED RELEASE;ORAL  
PENTOXIFYLLINE

ACTAVIS ELIZABETH	400MG	A074878 001	Jul 09, 1997
HERITAGE PHARMS INC	400MG	A074877 001	Jul 08, 1997
IMPAX LABS	400MG	A075093 001	Aug 10, 1999
PLIVA	400MG	A074874 001	May 25, 1999
TEVA	400MG	A075199 001	Sep 03, 1999
WATSON LABS	400MG	A075107 001	Sep 04, 1998
TRENTAL			
+ US PHARM HOLDINGS	400MG **	N018631 001	Aug 30, 1984

PERFLUBRONLIQUID;ORAL  
IMAGENT

ALLIANCE PHARM	100%	N020091 001	Aug 13, 1993
----------------	------	-------------	--------------

PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENEPASTE;TOPICAL  
SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS  
US ARMY 50%;50%

N021084 001 Feb 17, 2000

PERGOLIDE MESYLATETABLET;ORAL  
PERGOLIDE MESYLATE

IVAX SUB TEVA PHARMS	EQ 0.05MG BASE	A076094 001	Sep 04, 2003
	EQ 0.25MG BASE	A076094 002	Sep 04, 2003
	EQ 1MG BASE	A076094 003	Sep 04, 2003
PAR PHARM	EQ 0.05MG BASE	A076061 001	Nov 27, 2002
	EQ 0.25MG BASE	A076061 002	Nov 27, 2002
	EQ 1MG BASE	A076061 003	Nov 27, 2002
PERMAX			
VALEANT PHARM INTL	EQ 0.05MG BASE	N019385 001	Dec 30, 1988
	EQ 0.25MG BASE	N019385 002	Dec 30, 1988
	EQ 1MG BASE	N019385 003	Dec 30, 1988

PERINDOPRIL ERBUMINE

TABLET;ORAL

ACEON			
+ SYMPLMED PHARMS LLC	2MG	N020184 001	Dec 30, 1993
+	4MG	N020184 002	Dec 30, 1993
+	8MG	N020184 003	Dec 30, 1993

PERINDOPRIL ERBUMINE

LUPIN LTD	2MG	A078263 001	Jan 27, 2010
	4MG	A078263 002	Jan 27, 2010
	8MG	A078263 003	Jan 27, 2010

PERMETHRINLOTION;TOPICAL  
NIX

GLAXOSMITHKLINE	1%	N019435 001	Mar 31, 1986
-----------------	----	-------------	--------------

**DISCONTINUED DRUG PRODUCT LIST**

6-287(of 375)

\*\* See List Footnote

PERPHENAZINE

CONCENTRATE;ORAL			
PERPHENAZINE			
PHARM ASSOC	16MG/5ML		A040360 001 May 25, 2001
TRILAFON			
SCHERING	16MG/5ML		N011557 001
INJECTABLE;INJECTION			
TRILAFON			
SCHERING	5MG/ML		N011213 002
SYRUP;ORAL			
TRILAFON			
SCHERING	2MG/5ML		N011294 002
TABLET;ORAL			
PERPHENAZINE			
ANI PHARMS INC	2MG		A089707 001 Sep 10, 1987
	4MG		A089708 001 Sep 10, 1987
	8MG		A089456 001 Sep 10, 1987
	16MG		A089457 001 Sep 10, 1987
TRILAFON			
+ SCHERING	2MG **		N010775 001
+	4MG **		N010775 002
+	8MG **		N010775 003
+	16MG **		N010775 004
TABLET, EXTENDED RELEASE;ORAL			
TRILAFON			
SCHERING	8MG		N011361 002

PHENACEMIDE

TABLET;ORAL			
PHENURONE			
+ ABBVIE	500MG **		N007707 001

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE

TABLET;ORAL			
AZO GANTANOL			
+ ROCHE	100MG,500MG **		N013294 001 Sep 10, 1987

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET;ORAL			
SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PHENAZOPYRIDINE HYDROCHLORIDE			
ABLE	200MG,N/A,N/A;N/A,800MG,160MG		N021105 001 Jun 26, 2001

PHENAZOPYRIDINE HYDROCHLORIDE; SULFISOXAZOLE

TABLET;ORAL			
AZO GANTRISIN			
+ ROCHE	50MG;500MG **		N019358 001 Aug 31, 1990

PHENDIMETRAZINE TARTRATE

CAPSULE;ORAL			
PHENAZINE			
MAST MM	35MG		A086523 001
	35MG		A086524 001
	35MG		A086525 001
PHENDIMETRAZINE TARTRATE			
SANDOZ	35MG		A085633 001
	35MG		A085694 001
	35MG		A085695 001
	35MG		A085702 001
VITARINE	35MG		A085634 001
	35MG		A085645 001
	35MG		A085670 001
	35MG		A086403 001
	35MG		A086408 001
	35MG		A086410 001
	35MG		A087424 001
SPRX-3			
SOLVAY	35MG		A085897 001
STATOBEX			
TEVA	35MG		A085507 001
X-TROZINE			
SHIRE RICHWOOD	35MG		A087394 001 Sep 22, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-288(of 375)

\*\* See List Footnote

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE;ORAL

BONTRIL			
VALEANT	105MG	A088021	001 Sep 21, 1982
MELFIAT-105		A087487	001 Oct 13, 1982
NUMARK	105MG		
PHENDIMETRAZINE TARTRATE			
GRAHAM DM	105MG	A087214	001 May 26, 1982
	105MG	A088020	001 Aug 16, 1982
	105MG	A088028	001 Aug 16, 1982
	105MG	A088062	001 Sep 13, 1982
	105MG	A088063	001 Sep 10, 1982
	105MG	A088111	001 Oct 18, 1982
SANDOZ	105MG	A087378	001
SPRX-105			
NUMARK	105MG	A088024	001 Dec 22, 1982
X-TROZINE L.A.			
SHIRE RICHWOOD	105MG	A087371	001 Aug 24, 1982
TABLET;ORAL			
ADPHEN			
FERNDALE LABS	35MG	A083655	001
ALPHAZINE			
SANDOZ	35MG	A085034	001
CAM-METRAZINE			
ABC HOLDING	35MG	A085511	001
CAMALL	35MG	A085756	001
CHARTWELL RX	35MG	A083922	001
	35MG	A085318	001
	35MG	A085320	001
	35MG	A085321	001
DI-METREX			
PVT FORM	35MG	A085698	001
MELFIAT			
NUMARK	35MG	A083790	002
METRA			
FOREST PHARMS	35MG	A083754	001
PHENAZINE			
MAST MM	35MG	A087305	001
PHENAZINE-35			
ABC HOLDING	35MG	A085512	001
PHENDIMETRAZINE TARTRATE			
BARR	35MG	A083644	001
	35MG	A083684	001
	35MG	A083686	001
	35MG	A083687	001
	35MG	A084831	001
	35MG	A084834	001
	35MG	A084835	001
CHARTWELL RX	35MG	A085761	001
	35MG	A085941	001 Jun 27, 1983
FERNDALE LABS	35MG	A086834	001 Sep 15, 1983
INWOOD LABS	35MG	A084740	001
	35MG	A084741	001
	35MG	A084742	001
	35MG	A084743	001
IVAX PHARMS	35MG	A085611	001
	35MG	A085612	001
IVAX SUB TEVA PHARMS	35MG	A083682	001
KV PHARM	35MG	A084138	001
	35MG	A084141	001
	35MG	A085525	001
MFG CHEMISTS	35MG	A085914	001
NEXGEN PHARMA INC	35MG	A086020	001
NUMARK	35MG	A083790	001
PVT FORM	35MG	A085199	001
	35MG	A085697	001
SANDOZ	35MG	A085402	001
	35MG	A085497	001
	35MG	A085830	001

**DISCONTINUED DRUG PRODUCT LIST**

6-289(of 375)

\*\* See List Footnote

PHENDIMETRAZINE TARTRATE

TABLET;ORAL

PHENDIMETRAZINE TARTRATE

SOLVAY	35MG	A086365 001
USL PHARMA	35MG	A086370 001
VITARINE	35MG	A083993 001
	35MG	A083805 001
	35MG	A084398 001
	35MG	A084399 001
WATSON LABS	35MG	A085519 001
	35MG	A086005 001
	35MG	A086106 001
PLEGINE	35MG	A085767 001
WYETH AYERST	35MG **	N012248 001
STATOBEX	35MG	A086013 001
TEVA	35MG	A085095 001
TEVA	35MG	A086550 001
X-TROZINE	35MG	A086551 001
SHIRE RICHWOOD	35MG	A086552 001
	35MG	A086553 001
	35MG	A086554 001

PHENINDIONE

TABLET;ORAL

HEDULIN

SANOFI AVENTIS US	50MG	N008767 002
-------------------	------	-------------

PHENMETRAZINE HYDROCHLORIDE

TABLET;ORAL

PRELUDIN

BOEHRINGER INGELHEIM	25MG	N010460 005
TABLET, EXTENDED RELEASE;ORAL		
PRELUDIN		
BOEHRINGER INGELHEIM	50MG	N011752 004
	75MG	N011752 003

PHENPROCOUMON

TABLET;ORAL

LIQUAMAR

ORGANON USA INC	3MG	N011228 001
-----------------	-----	-------------

PHENSUXIMIDE

CAPSULE;ORAL

MILONTIN

PARKE DAVIS	500MG	N008855 004
-------------	-------	-------------

PHENTERMINE HYDROCHLORIDE

CAPSULE;ORAL

FASTIN

GLAXOSMITHKLINE	30MG **	N017352 001
-----------------	---------	-------------

OBESTIN-30

FERNDALE LABS	30MG	A087144 001
---------------	------	-------------

OBY-TRIM

SHIRE RICHWOOD	30MG	A087764 001 Mar 18, 1982
----------------	------	--------------------------

ONA-MAST

MAST MM	30MG	A086511 001
	30MG	A086516 001

PHENTERMINE HYDROCHLORIDE

ABC HOLDING	30MG	A085411 001
ABLE	15MG	A040497 001 Mar 13, 2003
	30MG	A040403 001 Aug 30, 2001
	30MG	A040427 001 Aug 30, 2001

CAMALL	15MG	A086735 001
	30MG	A087226 001

CHARTWELL RX	18.75MG	A088576 001 May 23, 1984
--------------	---------	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-290(of 375)

\*\* See List Footnote

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

	30MG	A085417 001
	30MG	A086732 002
	30MG	A087215 001
	37.5MG	A087915 001 Dec 22, 1983
	37.5MG	A087918 001 Dec 22, 1983
	37.5MG	A087930 001 Oct 14, 1983
	37.5MG	A088610 001 Jun 04, 1984
	37.5MG	A088611 001 Jun 04, 1984
	37.5MG	A088625 001 Aug 23, 1984
DURAMED PHARMS BARR	30MG	A088948 001 Apr 25, 1986
ELITE LABS INC	30MG	A040227 001 Jun 18, 1997
	30MG	A040448 001 Jan 22, 2003
IVAX PHARMS	30MG	A086329 001
MIKAH PHARMA	15MG	A040460 001 Jan 14, 2003
SANDOZ	30MG	A087208 001
	30MG	A087223 001
	37.5MG	A088414 001 Oct 19, 1983
SUN PHARM INDUSTRIES	37.5MG	A040527 001 Oct 23, 2003
TEVA	30MG	A086911 001
	30MG	A087126 001
	30MG	A087777 001 Nov 01, 1985
	30MG	A088612 001 Apr 04, 1984
	30MG	A088613 001 Apr 09, 1984
	30MG	A088614 001 Apr 09, 1984
TG UNITED INC	30MG	A040083 001 Mar 07, 1997
UPSHER-SMITH LABS	30MG	A084487 001 Apr 09, 1982
	30MG	A088430 001 Mar 27, 1984
USL PHARMA	30MG	A088797 001 Dec 10, 1984
VITARINE	30MG	A087202 001
	30MG	A087235 001
WATSON LABS	30MG	A086740 001 Mar 21, 1985
TABLET; ORAL		
ONA-MAST		
MAST MM	8MG	A086260 001
PHENTERMINE HYDROCHLORIDE		
ABLE	37.5MG	A040402 001 Aug 30, 2001
ACTAVIS ELIZABETH	37.5MG	A040276 001 Nov 25, 1998
CHARTWELL RX	8MG	A083923 001
	8MG	A085319 001
	37.5MG	A087805 001 Dec 06, 1982
	37.5MG	A088596 001 Apr 04, 1984
IVAX PHARMS	8MG	A085553 001
SANDOZ	8MG	A085671 001
	8MG	A085689 001
SANDOZ INC	30MG	A088605 001 Sep 28, 1987
USL PHARMA	8MG	A083804 001
	37.5MG	A088910 001 Jul 17, 1985
	37.5MG	A088917 001 Jul 17, 1985
VITARINE	8MG	A086453 001
	8MG	A086456 001
WATSON LABS	8MG	A085739 001
TORA		
SOLVAY	8MG	A084035 001
WILPO		
+ SANDOZ	8MG **	N012737 001
TABLET, ORALLY DISINTEGRATING; ORAL		
SUPRENZA		
CITIUS PHARMS	15MG	N202088 001 Jun 13, 2011
	30MG	N202088 002 Jun 13, 2011
	37.5MG	N202088 003 Mar 27, 2012

**DISCONTINUED DRUG PRODUCT LIST**

6-291(of 375)

\*\* See List Footnote

PHENTERMINE RESIN COMPLEXCAPSULE, EXTENDED RELEASE;ORAL  
IONAMIN

UCB INC	EQ 15MG BASE **	N011613 004
	EQ 30MG BASE **	N011613 002
PHENTERMINE RESIN 30		
QUANTUM PHARMICS	EQ 30MG BASE	A089120 001 Feb 04, 1988
PHENTERMINE RESIN COMPLEX		
LANNETT HOLDINGS INC	EQ 15MG BASE	A040872 001 Jul 28, 2011
	EQ 30MG BASE	A040872 002 Jul 28, 2011

PHENTOLAMINE MESYLATEINJECTABLE; INJECTION  
REGITINE

+ NOVARTIS	5MG/VIAL	N008278 003
------------	----------	-------------

PHENYL AMINOSALICYLATE

POWDER;ORAL

PHENY-PAS-TEBAMIN		
PHARM RES ASSOC	50%	N011695 002
TABLET;ORAL		
PHENY-PAS-TEBAMIN		
PHARM RES ASSOC	500MG	N011695 003

PHENYLBUTAZONE

CAPSULE;ORAL

AZOLID		
SANOFI AVENTIS US	100MG	A087260 001
BUTAZOLIDIN		
NOVARTIS	100MG	N008319 009
PHENYLBUTAZONE		
IVAX PHARMS	100MG	A088218 001 Jun 24, 1983
SANDOZ	100MG	A087774 001 Jun 16, 1982
SUN PHARM INDUSTRIES	100MG	A088994 001 Dec 04, 1985
WATSON LABS	100MG	A087756 001 Dec 17, 1982
TABLET;ORAL		
AZOLID		
SANOFI AVENTIS US	100MG	A087091 001
BUTAZOLIDIN		
NOVARTIS	100MG	N008319 008
PHENYLBUTAZONE		
SANDOZ	100MG	A084339 001
SUN PHARM INDUSTRIES	100MG	A088863 001 Dec 04, 1985
WATSON LABS	100MG	A086151 001
	100MG	A087674 001 Apr 21, 1982

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PHENERGAN VC		
+ ANI PHARMS	5MG/5ML; 6.25MG/5ML **	N008604 003 Apr 02, 1984
PHERAZINE VC		
HALSEY	5MG/5ML; 6.25MG/5ML	A088868 001 Mar 02, 1987
PROMETHAZINE VC PLAIN		
CENCI	5MG/5ML; 6.25MG/5ML	A088815 001 Nov 22, 1985
WOCKHARDT	5MG/5ML; 6.25MG/5ML	A088897 001 Jan 04, 1985

PHENYLEPHRINE HYDROCHLORIDE; PYRILAMINE MALEATE

SOLUTION/DROPS;OPHTHALMIC

PREFRIN-A		
ALLERGAN	0.12%; 0.1%	N007953 001

PHENYTOIN

SUSPENSION;ORAL

DILANTIN-30		
PARKE DAVIS	30MG/5ML	N008762 002
PHENYTOIN		
ACTAVIS MID ATLANTIC	125MG/5ML	A089892 001 Sep 25, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-292(of 375)

\*\* See List Footnote

PHENYTOIN SODIUM

CAPSULE;ORAL

DIPHENYLAN SODIUM

LANNETT

30MG PROMPT

A080857 001

100MG PROMPT

A080857 002

EXTENDED PHENYTOIN SODIUM

ANI PHARMS INC

100MG EXTENDED

A040435 001 Jun 20, 2003

100MG EXTENDED

A089441 001 Dec 18, 1986

WOCKHARDT

30MG EXTENDED

A040759 001 Dec 18, 2007

WOCKHARDT USA

100MG EXTENDED

A040732 001 Jan 30, 2008

PHENYTEX

WATSON LABS

100MG EXTENDED

A088711 001 Dec 21, 1984

PHENYTOIN SODIUM

PHARMERAL

100MG PROMPT

A085435 001

WATSON LABS

100MG PROMPT

A085894 001

PROMPT PHENYTOIN SODIUM

ANI PHARMS INC

100MG PROMPT

A080259 001

WATSON LABS

100MG PROMPT

A080905 001

INJECTABLE;INJECTION

DILANTIN

PARKE DAVIS

50MG/ML

N010151 001

PHENYTOIN SODIUM

FRESENIUS KABI USA

50MG/ML

A089003 001 May 31, 1985

HOSPIRA

50MG/ML

A089521 001 Mar 17, 1987

50MG/ML

A089744 001 Dec 18, 1987

MARSAM PHARMS LLC

50MG/ML

A089501 001 Oct 13, 1987

50MG/ML

A089779 001 Nov 27, 1992

SMITH AND NEPHEW

50MG/ML

A088519 001 Dec 19, 1984

50MG/ML

A088521 001 Dec 18, 1984

SOLOPAK

50MG/ML

A088520 001 Dec 17, 1984

WARNER CHILCOTT

50MG/ML

A089900 001 Mar 30, 1990

WATSON LABS

50MG/ML

A085434 001

PHYTONADIONE

INJECTABLE;INJECTION

AQUAMEPHYTON

+ TELIGENT

1MG/0.5ML \*\*

N012223 002

+ +

10MG/ML \*\*

N012223 001

KONAKION

ROCHE

1MG/0.5ML

N011745 001

10MG/ML

N011745 003

PHYTONADIONE

GLAXOSMITHKLINE

1MG/0.5ML

A084060 001

10MG/ML

A084060 002

VITAMIN K1

HOSPIRA

10MG/ML

A087956 001 Jul 25, 1983

PILOCARPINE

INSERT, EXTENDED RELEASE;OPHTHALMIC

OCUSERT PILO-20

AKORN

5MG

N017431 001

OCUSERT PILO-40

AKORN

11MG

N017548 001

PILOCARPINE HYDROCHLORIDE

GEL;OPHTHALMIC

PILOPINE HS

ALCON

4%

N018796 001 Oct 01, 1984

PINACIDIL

CAPSULE, EXTENDED RELEASE;ORAL

PINDAC

LEO PHARM

12.5MG

N019456 001 Dec 28, 1989

25MG

N019456 002 Dec 28, 1989

PINDOLOL

TABLET;ORAL

PINDOLOL

G AND W LABS INC

5MG

A073661 001 Oct 31, 1993

5MG

A073687 001 Feb 26, 1993

5MG

A074123 001 Apr 17, 1997

10MG

A073661 002 Oct 31, 1993

**DISCONTINUED DRUG PRODUCT LIST**

6-293(of 375)

\*\* See List Footnote

PINDOLOLTABLET;ORAL  
PINDOLOL

	10MG	A073687 002	Feb 26, 1993
	10MG	A074123 002	Apr 17, 1997
MYLAN PHARMS INC	5MG	A074013 001	Sep 24, 1992
	10MG	A074018 001	Sep 24, 1992
NOSTRUM LABS	5MG	A074474 001	Oct 28, 1996
	10MG	A074474 002	Oct 28, 1996
PUREPAC PHARM	5MG	A074125 001	Apr 28, 1993
	10MG	A074125 002	Apr 28, 1993
WATSON LABS	5MG	A074437 001	Feb 27, 1995
	10MG	A074437 002	Feb 27, 1995
VISKEN			
+ NOVARTIS	5MG **	N018285 001	Sep 03, 1982
+	10MG **	N018285 002	Sep 03, 1982

PIPECURONIUM BROMIDEINJECTABLE;INJECTION  
ARDUAN

ORGANON USA INC	10MG/VIAL	N019638 001	Jun 26, 1990
-----------------	-----------	-------------	--------------

PIPERACETAZINETABLET;ORAL  
QUIDE

DOW PHARM	10MG	N013615 001
	25MG	N013615 002

PIPERACILLIN SODIUMINJECTABLE;INJECTION  
PIPRACIL

WYETH PHARMS INC	EQ 2GM BASE/VIAL	A062750 001	Oct 13, 1987
+	EQ 2GM BASE/VIAL **	N050545 002	
	EQ 3GM BASE/VIAL	A062750 002	Oct 13, 1987
+	EQ 3GM BASE/VIAL **	N050545 003	
	EQ 4GM BASE/VIAL	A062750 003	Oct 13, 1987
+	EQ 4GM BASE/VIAL **	N050545 004	
+	EQ 40GM BASE/VIAL **	N050545 006	Sep 30, 1985

PIPERAZINE CITRATESYRUP;ORAL  
ANTEPAR

GLAXOSMITHKLINE	EQ 500MG BASE/5ML	N009102 001
-----------------	-------------------	-------------

BRYREL SANOFI AVENTIS US EQ 500MG BASE/5ML N017796 001

MULTIFUGE

BLULINE	EQ 500MG BASE/5ML	N009452 001
---------	-------------------	-------------

PIPERAZINE CITRATE

ALPHARMA US PHARMS	EQ 500MG BASE/5ML	A080774 001
--------------------	-------------------	-------------

LANNETT	EQ 500MG BASE/5ML	A080963 001
---------	-------------------	-------------

LUITPOLD	EQ 500MG BASE/5ML	A080671 001
----------	-------------------	-------------

VERMIDOL

SOLVAY	EQ 500MG BASE/5ML	A080992 001
--------	-------------------	-------------

TABLET;ORAL

ANTEPAR GLAXOSMITHKLINE EQ 500MG BASE N009102 003

PIPERAZINE CITRATE IMPAX LABS EQ 250MG BASE A080874 001

PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL;TOPICAL

RID MOUSSE BAYER HEALTHCARE LLC 4%;EQ 0.33% BASE N021043 001 Mar 07, 2000

PIPOBROMANTABLET;ORAL  
VERCYTE

ABBOTT	10MG	N016245 001
	25MG	N016245 002

**DISCONTINUED DRUG PRODUCT LIST**

6-294(of 375)

\*\* See List Footnote

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION  
MAXAIR

MEDICIS	EQ 0.2MG BASE/INH	N020014 001 Nov 30, 1992
VALEANT PHARMS	EQ 0.2MG BASE/INH	N019009 001 Dec 30, 1986

PIRFENIDONE

TABLET; ORAL  
ESBRIET

+ GENENTECH INC	534MG	N208780 002 Jan 11, 2017
-----------------	-------	--------------------------

PIROXICAM

CAPSULE; ORAL  
PIROXICAM

CYCLE PHARMS LTD	10MG	A073651 001 Feb 26, 1993
	20MG	A073651 002 Feb 26, 1993
EGIS	10MG	A074808 001 Jul 08, 1997
	20MG	A074808 002 Jul 08, 1997
IVAX SUB TEVA PHARMS	10MG	A074148 001 Jun 03, 1996
	20MG	A074148 002 Jun 03, 1996
MYLAN	10MG	A074043 001 Sep 22, 1992
	10MG	A074102 001 Jul 31, 1992
	20MG	A074043 002 Sep 22, 1992
	20MG	A074102 002 Jul 31, 1992
SCS	10MG	A074036 001 May 29, 1992
	20MG	A074036 002 May 29, 1992
TEVA	10MG	A073637 001 Jan 28, 1994
	20MG	A073638 001 Jan 28, 1994
TEVA PHARMS	10MG	A074103 001 Aug 28, 1992
	20MG	A074103 002 Aug 28, 1992
WATSON LABS	10MG	A074287 001 May 16, 1996
	10MG	A074460 001 Sep 29, 1995
	20MG	A074287 002 May 16, 1996
	20MG	A074460 002 Sep 29, 1995

PITAVASTATIN SODIUM

TABLET; ORAL

NIKITA		
+ LUPIN LTD	EQ 1MG BASE	N209875 001 Aug 04, 2017
+	EQ 2MG BASE	N209875 002 Aug 04, 2017
+	EQ 4MG BASE	N209875 003 Aug 04, 2017

PLICAMYCIN

INJECTABLE; INJECTION

MITHRACIN		
PFIZER	2.5MG/VIAL	N050109 001

PODOFILOX

SOLUTION; TOPICAL  
PODOFILOX

BAUSCH AND LOMB INC	0.5%	A090184 001 Jul 21, 2010
---------------------	------	--------------------------

POLYESTRADOL PHOSPHATE

INJECTABLE; INJECTION

ESTRADURIN		
WYETH AYERST	40MG/AMP	N010753 001

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

POLYETHYLENE GLYCOL 3350		
PADDICK LLC	17GM/SCOOPFUL	A090567 001 Oct 15, 2009
TEVA PHARMS	17GM/SCOOPFUL	A077445 001 May 04, 2006

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE		
MYLAN	420GM/BOT; 1.48GM/BOT; 5.72GM/BOT; 11.2GM/BOT	A090409 001 Apr 02, 2010

**DISCONTINUED DRUG PRODUCT LIST**

6-295(of 375)

\*\* See List Footnote

**POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE**

FOR SOLUTION;ORAL

CLENZ-LYTE

PADDOCK LLC

236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/  
BOT;22.74GM/BOT

A090769 001 Jun 07, 2010

SOLUTION;ORAL

OCL

HOSPIRA

6GM/100ML;75MG/100ML;168MG/100ML;146MG/  
100ML;1.29GM/100ML

N019284 001 Apr 30, 1986

**POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE  
ANHYDROUS**

FOR SOLUTION;ORAL

COLYTE

MYLAN SPECIALITY LP 120GM/PACKET;1.49GM/PACKET;3.36GM/PACKE  
T;2.92GM/PACKET;11.36GM/PACKET  
227.1GM/PACKET;2.82GM/PACKET;6.36GM/PAC  
KET;5.53GM/PACKET;21.5GM/PACKET  
227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53G  
M/BOT;21.5GM/BOT  
240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/  
BOT;22.72GM/BOT  
360GM/PACKET;4.47GM/PACKET;10.08GM/PACK  
ET;8.76GM/PACKET;34.08GM/PACKET

N018983 005 Oct 26, 1984

N018983 004 Oct 26, 1984

N018983 010 Jan 31, 1989

N018983 007 Jun 12, 1987

N018983 006 Oct 26, 1984

COLYTE-FLAVORED

MYLAN SPECIALITY LP 227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53G  
M/BOT;21.5GM/BOT  
240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/  
BOT;22.72GM/BOT

N018983 008 Nov 14, 1991

N018983 009 Nov 14, 1991

PEG 3350 AND ELECTROLYTES

MYLAN 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/  
BOT;22.74GM/BOT

A090928 001 Jan 28, 2010

POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

PADDOCK LLC 240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/  
BOT;22.72GM/BOT

A090712 001 Feb 25, 2010

FOR SUSPENSION;ORAL

CO-LAV

VINTAGE PHARMS 240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/  
BOT;22.72GM/BOT

A073428 001 Jan 28, 1992

COLOVAGE

DYNAPHARM 227.1GM/PACKET;2.82GM/PACKET;6.36GM/PAC  
KET;5.53GM/PACKET;21.5GM/PACKET

A071320 001 Apr 20, 1988

E-Z-EM PREP LYTE

E Z EM 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/  
BOT;22.74GM/BOT

A071278 001 Nov 21, 1988

GLYCOPREP

GOLDLINE 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/  
BOT;22.74GM/BOT

A072319 001 Dec 23, 1988

GO-EVAC

VINTAGE PHARMS 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/  
BOT;22.74GM/BOT

A073433 001 Apr 28, 1992

PEG-LYTE

SANDOZ 236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/  
BOT;22.74GM/BOT

A073098 001 Aug 31, 1993

**POLYMYXIN B SULFATE**

INJECTABLE;INJECTION

AEROSPORIN

GLAXOSMITHKLINE EQ 500,000 U BASE/VIAL

A062036 001

POWDER;FOR RX COMPOUNDING

POLY-RX

X GEN PHARMS 100,000,000 UNITS/BOT

A061578 001

POLYMYXIN B SULFATE

PADDOCK LLC 100,000,000 UNITS/BOT

A062455 001 Jul 27, 1983

**POLYTHIAZIDE**

TABLET;ORAL

RENESE

PFIZER 1MG  
2MG  
4MG

N012845 001

N012845 002

N012845 003

**DISCONTINUED DRUG PRODUCT LIST**

6-296(of 375)

\*\* See List Footnote

POLYTHIAZIDE; PRAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

MINIZIDE

PFIZER

0.5MG;EQ 1MG BASE

N017986 001

0.5MG;EQ 2MG BASE

N017986 002

0.5MG;EQ 5MG BASE

N017986 003

POLYTHIAZIDE; RESERPINE

TABLET;ORAL

RENESE-R

PFIZER

2MG;0.25MG

N013636 001

POTASSIUM AMINOSALICYLATE

CAPSULE;ORAL

PASKALIUM

GLENWOOD

500MG

N009395 004

POWDER;ORAL

POTASSIUM AMINOSALICYLATE

HEXCEL

100%

A080098 001

TABLET;ORAL

PASKALIUM

GLENWOOD

1GM

N009395 003

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

K-LEASE

SAVAGE LABS

8MEQ

A073398 001 Jan 28, 1992

10MEQ

A072427 001 Mar 28, 1990

POTASSIUM CHLORIDE

NESHER PHARMS

10MEQ

A070980 001 Feb 17, 1987

TEVA

8MEQ

A073531 001 Apr 26, 1996

10MEQ

A073532 001 Apr 26, 1996

FOR SUSPENSION, EXTENDED RELEASE;ORAL

MICRO-K LS

KV PHARM

20MEQ/PACKET

N019561 003 Aug 26, 1988

INJECTABLE;INJECTION

POTASSIUM CHLORIDE

ABRAXIS PHARM

2MEQ/ML

A080204 001

2MEQ/ML

A084290 001

2MEQ/ML

A086713 001

2MEQ/ML

A086714 001

2MEQ/ML

A087787 001 Apr 20, 1982

2MEQ/ML

A087885 001 Feb 03, 1983

AKORN

2MEQ/ML

A088286 001 Sep 05, 1985

BAXTER HLTHCARE

2MEQ/ML

A080203 001

2MEQ/ML

A085499 001

FRESENIUS KABI USA

2MEQ/ML

A087817 001 Oct 20, 1982

GD SEARLE LLC

1MEQ/ML

A086219 001

2MEQ/ML

A086219 002

2MEQ/ML

A086220 002

3MEQ/ML

A086219 003

3MEQ/ML

A086220 001

4MEQ/ML

A086219 004

HOSPIRA

1MEQ/ML

A080205 003

1MEQ/ML

A083345 003

1.5MEQ/ML

A083345 001

2MEQ/ML

A083345 002

2.4MEQ/ML

A080205 004

3.2MEQ/ML

A080205 005

INTL MEDICATION

2MEQ/ML

A083163 001

LILLY

2MEQ/ML

N007865 002

LUITPOLD

2MEQ/ML

A080221 001

2MEQ/ML

A080736 001

2MEQ/ML

A087584 001

2MEQ/ML

A087585 001

MILES

1MEQ/ML

A080195 002

2MEQ/ML

A080195 001

3MEQ/ML

A080195 003

4MEQ/ML

A080195 004

PHARMA SERVE NY

2MEQ/ML

A086297 001

2MEQ/ML

A087362 001 Mar 08, 1983

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-297(of 375)

\*\* See List Footnote

POTASSIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

WATSON LABS

2MEQ/ML

A086208 001

2MEQ/ML

A089163 001 Mar 10, 1988

2MEQ/ML

A089421 001 Jan 02, 1987

3MEQ/ML

A086210 001

TABLET, EXTENDED RELEASE; ORAL

K+10

FUTURE PAK

10MEQ

A070999 001 Oct 22, 1987

K+8

FUTURE PAK

8MEQ

A070998 001 Jan 25, 1993

KAON CL

SAVAGE LABS

6.7MEQ

N017046 001

KAON CL-10

SAVAGE LABS

10MEQ

N017046 002

KLOTRIX

APOTHECON

10MEQ

N017850 001

POTASSIUM CHLORIDE

COPELY PHARM

8MEQ

A070618 001 Sep 09, 1987

NESHER PHARMS

20MEQ

A076044 001 Apr 05, 2002

+ SCHERING

10MEQ \*\*

N019439 002 Jun 13, 1986

+ SCHERING

20MEQ \*\*

N019439 001 Jun 13, 1986

SLOW-K

NOVARTIS

8MEQ

N017476 002

TEN-K

NOVARTIS

10MEQ

N019381 001 Apr 16, 1986

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.037% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 37MG/100ML; 900MG/100ML

N019708 001 Sep 29, 1989

POTASSIUM CHLORIDE 0.075% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 75MG/100ML; 900MG/100ML

N019708 002 Sep 29, 1989

POTASSIUM CHLORIDE 0.11% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 110MG/100ML; 900MG/100ML

N019708 003 Sep 29, 1989

POTASSIUM CHLORIDE 0.22% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 220MG/100ML; 900MG/100ML

N019708 005 Sep 29, 1989

POTASSIUM CHLORIDE 0.3% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN 300MG/100ML; 900MG/100ML

N019708 006 Sep 29, 1989

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER

B BRAUN 75MG/100ML; 900MG/100ML

N018722 001 Nov 09, 1982

BAXTER HLTHCARE 75MG/100ML; 900MG/100ML

N017648 004

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

B BRAUN 150MG/100ML; 900MG/100ML

N018722 002 Nov 09, 1982

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER

B BRAUN 220MG/100ML; 900MG/100ML

N018722 003 Nov 09, 1982

SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN 300MG/100ML; 900MG/100ML

N018722 004 Nov 09, 1982

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE; INJECTION

THAM-E

HOSPIRA

370MG/VIAL; 1.75GM/VIAL; 36GM/VIAL

N013025 001

POTASSIUM CITRATE

FOR SOLUTION; ORAL

POTASSIUM CITRATE

+ UT SW MEDCTR

10MEQ/PACKET \*\*

N019647 002 Oct 13, 1988

+

20MEQ/PACKET \*\*

N019647 001 Oct 13, 1988

POTASSIUM IODIDE

SOLUTION; ORAL

POTASSIUM IODIDE

ROXANE

1GM/ML

N018551 001 Feb 19, 1982

TABLET; ORAL

THYRO-BLOCK

MEDA PHARMS

130MG

N018307 001

**DISCONTINUED DRUG PRODUCT LIST**

6-298(of 375)

\*\* See List Footnote

POTASSIUM PERCHLORATE

CAPSULE; ORAL PERCHLORACAP MALLINCKRODT	200MG	N017551 001
---	-------	-------------

POVIDONE-IODINE

SOLUTION; TOPICAL E-Z PREP CLINIPAD	10%	N019382 001 Jul 25, 1989
SPONGE; TOPICAL E-Z PREP CLINIPAD	5%	N019382 002 Jul 25, 1989
E-Z PREP 220 CLINIPAD	5%	N019382 003 Jul 25, 1989

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION PRALIDOXIME CHLORIDE BAXTER HLTHCARE CORP	300MG/ML	N018799 001 Dec 13, 1982
TABLET; ORAL PROTOPAM CHLORIDE WYETH AYERST	500MG	N014122 002

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL MIRAPEX BOEHRINGER INGELHEIM	1.25MG	N020667 004 Jul 01, 1997
PRAMIPEXOLE DIHYDROCHLORIDE ACTAVIS GRP PTC	0.125MG	A091254 001 Nov 30, 2010
	0.25MG	A091254 002 Nov 30, 2010
	0.5MG	A091254 003 Nov 30, 2010
	0.75MG	A091254 004 Nov 30, 2010
	1MG	A091254 005 Nov 30, 2010
	1.5MG	A091254 006 Nov 30, 2010
SANDOZ	0.125MG	A090190 001 Jul 06, 2010
	0.25MG	A090190 002 Jul 06, 2010
	0.5MG	A090190 003 Jul 06, 2010
	0.75MG	A090190 006 Oct 08, 2010
	1MG	A090190 004 Jul 06, 2010
	1.5MG	A090190 005 Jul 06, 2010
WATSON LABS	0.125MG	A078551 001 Oct 08, 2010
	0.25MG	A078551 002 Oct 08, 2010
	0.5MG	A078551 003 Oct 08, 2010
	1MG	A078551 004 Oct 08, 2010
	1.5MG	A078551 005 Oct 08, 2010

PRAMILINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS SYMLIN ASTRAZENECA AB	EQ 3MG BASE/5ML (EQ 600MCG BASE/ML)	N021332 001 Mar 16, 2005
--	-------------------------------------	--------------------------

PRAVASTATIN SODIUM

TABLET; ORAL PRAVACHOL + BRISTOL MYERS SQUIBB	10MG **	N019898 002 Oct 31, 1991
PRAVASTATIN SODIUM MYLAN	10MG	A077013 001 Oct 23, 2006
	20MG	A077013 002 Oct 23, 2006
	40MG	A077013 003 Oct 23, 2006
	80MG	A077013 004 Dec 28, 2007
PLIVA HRVATSKA DOO	10MG	A077730 001 Nov 21, 2006
	20MG	A077730 002 Nov 21, 2006
	30MG	A077730 003 Nov 21, 2006
	40MG	A077730 005 Nov 21, 2006
RANBAXY LABS LTD	10MG	A076445 001 Apr 23, 2007
	20MG	A076445 002 Apr 23, 2007
	40MG	A076445 003 Apr 23, 2007
	80MG	A076445 004 Apr 23, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-299(of 375)

\*\* See List Footnote

PRAZEPAM

CAPSULE;ORAL

CENTRAX

PARKE DAVIS

5MG

N018144 001

10MG

N018144 002

20MG

N018144 003 May 10, 1982

PRAZEPAM

USL PHARMA

5MG

A070427 001 Nov 06, 1987

10MG

A070428 001 Nov 06, 1987

TABLET;ORAL

CENTRAX

PARKE DAVIS

10MG

N017415 001

PRAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

PRAZOSIN HYDROCHLORIDE

AM THERAP

EQ 1MG BASE

A072782 001 May 16, 1989

EQ 2MG BASE

A072783 001 May 16, 1989

EQ 5MG BASE

A072784 001 May 16, 1989

DAVA PHARMS INC

EQ 1MG BASE

A072705 001 May 16, 1989

EQ 2MG BASE

A072706 001 May 16, 1989

EQ 5MG BASE

A072707 001 May 16, 1989

IDT AUSTRALIA LTD

EQ 1MG BASE

A072577 002 May 16, 1989

EQ 2MG BASE

A072577 001 May 16, 1989

EQ 5MG BASE

A072577 003 May 16, 1989

PUREPAC PHARM

EQ 1MG BASE

A072991 001 May 16, 1989

EQ 2MG BASE

A072921 001 May 16, 1989

EQ 5MG BASE

A072992 001 May 16, 1989

WATSON LABS

EQ 1MG BASE

A072352 001 May 16, 1989

EQ 2MG BASE

A072333 001 May 16, 1989

EQ 5MG BASE

A072609 001 May 16, 1989

TABLET, EXTENDED RELEASE;ORAL

MINIPRESS XL

PFIZER

2.5MG

N019775 001 Jan 29, 1992

5MG

N019775 002 Jan 29, 1992

PREDNISOLONE

CREAM;TOPICAL

METI-DERM

SCHERING

0.5%

N010209 002

SYRUP;ORAL

PREDNISOLONE

APOTEX INC

5MG/5ML

A040570 001 Aug 25, 2005

15MG/5ML

A040571 001 Aug 25, 2005

IVAX SUB TEVA PHARMS

15MG/5ML

A040287 001 May 28, 1999

NESHER PHARMS

5MG/5ML

A040423 001 Oct 22, 2001

15MG/5ML

A040364 001 Apr 10, 2002

TEVA PHARMS

15MG/5ML

A040322 001 Jan 19, 2000

WE PHARMS

15MG/5ML

A040192 001 May 28, 1998

PRELONE

MURO

5MG/5ML

A089654 001 Jan 17, 1989

TABLET;ORAL

CORTALONE

HALSEY

1MG

A080304 003

2.5MG

A080304 002

5MG

A080304 001

DELTA-CORTEF

PHARMACIA AND UPJOHN

5MG

N009987 004

FERNISOLONE-P

FERNDALE LABS

5MG

A083941 001

PREDNISOLONE

AUROLIFE PHARMA LLC

5MG

A084773 001

BARR

5MG

A084426 002

BUNDY

5MG

A083675 001

CHARTWELL RX

5MG

A084542 001

ELKINS SINK

5MG

A080625 001

EVERYLIFE

1MG

A084439 001

2.5MG

A084439 002

5MG

A084439 003

FERRANTE

2.5MG

A080562 001

**DISCONTINUED DRUG PRODUCT LIST**

6-300(of 375)

\*\* See List Footnote

PREDNISOLONE

TABLET;ORAL

PREDNISOLONE

	5MG	A080562 002
HEATHER	5MG	A080326 001
IMPAKX LABS	5MG	A080780 001
INWOOD LABS	5MG	A080748 001
IVAX SUB TEVA PHARMS	5MG	A080378 001
LANNETT	5MG	A080531 002
MARSHALL PHARMA	5MG	A080307 001
PANRAY	1MG	A080351 001
	5MG	A080351 002
PHOENIX LABS NY	5MG	A080322 001
PUREPAC PHARM	5MG	A080325 001
PVT FORM	5MG	A080211 001
ROXANE	5MG	A080327 002
SANDOZ	5MG	A080339 001
SPERTI	1MG	A080358 001
	2.5MG	A080358 002
	5MG	A080358 003
SUPERPHARM	5MG	A088892 001 Feb 26, 1985
TABLICAPS	5MG	A085170 001
TEVA	5MG	A080398 001
UDL	5MG	A087987 001 Jan 18, 1983
VALEANT PHARM INTL	5MG	A080236 001
VITARINE	5MG	A080534 001
WATSON LABS	5MG	A085085 002
	5MG	A085415 001
	5MG	A085416 001
WEST WARD	5MG	A080324 001
WHITEWORTH TOWN PLSN	5MG	A080342 001
STERANE		
PFIZER	5MG	N009996 001

PREDNISOLONE ACETATE

INJECTABLE;INJECTION

METICORTELONE

SCHERING 25MG/ML

N010255 002

PREDNISOLONE ACETATE

AKORN 25MG/ML

A083032 001

50MG/ML

A084492 001

BEL MAR 25MG/ML

A083738 001

50MG/ML

A083738 002

CENT PHARMS 25MG/ML

A084717 001

50MG/ML

A084717 002

WATSON LABS 25MG/ML

A083398 001

25MG/ML

A083654 001

40MG/ML

A083767 001

50MG/ML

A083764 001

50MG/ML

A085781 001

STERANE

PFIZER 25MG/ML

N011446 001

SUSPENSION;ORAL

FLO-PRED

TARO EQ 5MG BASE/5ML

N022067 001 Jan 17, 2008

EQ 15MG BASE/5ML

N022067 002 Jan 17, 2008

SUSPENSION/DROPS;OPHTHALMIC

ECONOPRED

ALCON 0.125%

N017468 001

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

CETAPRED

ALCON 0.25%;10%

A087771 001 Aug 06, 1993

METIMYD

SCHERING 0.5%;10%

N010210 002 Sep 09, 1984

PREDSULFAIR

PHARMAFAIR 0.5%;10%

A088032 001 Apr 15, 1983

VASOCIDIN

NOVARTIS 0.5%;10%

A088791 001 Oct 05, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-301(of 375)

\*\* See List Footnote

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

SUSPENSION;OPHTHALMIC		
ISOPTO CETAPRED		
ALCON	0.25%;10%	A087547 001
SUSPENSION/DROPS;OPHTHALMIC		
METIMYD		
SCHERING	0.5%;10%	N010210 001
PREDAMIDE		
AKORN	0.5%;10%	A088059 001 Jul 29, 1983
PREDSULFAIR		
PHARMAFAIR	0.5%;10%	A088007 001 Apr 19, 1983
PREDSULFAIR II		
PHARMAFAIR	0.2%;10%	A088837 001 Dec 24, 1985
SULPHRIN		
BAUSCH AND LOMB	0.5%;10%	A088089 001 Dec 28, 1982

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE;INJECTION		
HYDELTRASOL		
MERCK	EQ 20MG PHOSPHATE/ML	N011583 002
PREDNISOLONE SODIUM PHOSPHATE		
WATSON LABS	EQ 20MG PHOSPHATE/ML	A080517 001
OINTMENT;OPHTHALMIC, OTIC		
HYDELTRASOL		
MERCK	EQ 0.25% PHOSPHATE	N011028 001
SOLUTION;ORAL		
ORAPRED		
CONCORDIA PHARMS INC	EQ 15MG BASE/5ML **	A075117 001 Dec 14, 2000
PREDNISOLONE SODIUM PHOSPHATE		
AMNEAL PHARMS	EQ 15MG BASE/5ML	A078345 001 Mar 10, 2009
MEDICIS PHARMS	EQ 15MG BASE/5ML	A075250 001 Jul 12, 2002
NESHER PHARMS	EQ 5MG BASE/5ML	A076982 001 May 24, 2005
	EQ 15MG BASE/5ML	A076988 001 May 24, 2005
PHARM ASSOC	EQ 5MG BASE/5ML	A076123 001 Dec 23, 2002
VINTAGE PHARMS	EQ 5MG BASE/5ML	A078416 001 Oct 31, 2007
WE PHARMS	EQ 5MG BASE/5ML	A075181 001 Dec 23, 2002
SOLUTION/DROPS;OPHTHALMIC		
INFLAMASE FORTE		
NOVARTIS	EQ 0.9% PHOSPHATE	A080751 002
INFLAMASE MILD		
NOVARTIS	EQ 0.11% PHOSPHATE	A080751 001
METRETONE		
SCHERING	EQ 0.5% PHOSPHATE	A083834 001
PREDAIR		
PHARMAFAIR	EQ 0.11% PHOSPHATE	A088415 001 Feb 29, 1984
PREDAIR FORTE		
PHARMAFAIR	EQ 0.9% PHOSPHATE	A088165 001 Mar 28, 1983
PREDNISOLONE SODIUM PHOSPHATE		
AKORN	EQ 0.11% PHOSPHATE	A083358 001
	EQ 0.9% PHOSPHATE	A083358 002
ALCON PHARMS LTD	EQ 0.11% PHOSPHATE	A081043 001 Oct 24, 1991
	EQ 0.9% PHOSPHATE	A081044 001 Oct 24, 1991
BAUSCH AND LOMB	EQ 0.11% PHOSPHATE	A040065 001 Jul 29, 1994
SOLA BARNES HIND	EQ 0.11% PHOSPHATE	A084171 001
	EQ 0.9% PHOSPHATE	A084168 001
	EQ 0.9% PHOSPHATE	A084169 001
	EQ 0.9% PHOSPHATE	A084172 001

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC		
SULSTER		
AKORN	EQ 0.23% PHOSPHATE;10%	A074511 001 Jul 30, 1996
VASOCIDIN		
+ NOVARTIS	EQ 0.23% PHOSPHATE;10% **	N018988 001 Aug 26, 1988

**DISCONTINUED DRUG PRODUCT LIST**

6-302(of 375)

\*\* See List Footnote

PREDNISOLONE TEBUTATE

INJECTABLE; INJECTION		
HYDELTRA-TBA		
MERCK	20MG/ML	N010562 001
PREDNISOLONE TEBUTATE		
WATSON LABS	20MG/ML	A083362 001 Feb 17, 1984

PREDNISONE

SOLUTION;ORAL		
PREDNISONE		
WOCKHARDT	5MG/5ML	A089726 001 Aug 02, 1988
SYRUP;ORAL		
LIQUID PRED		
MURO	5MG/5ML	A087611 002 Sep 07, 1982
TABLET;ORAL		
CORTAN		
HALSEY	20MG	A087480 001
DELTA-DOME		
BAYER PHARMS	5MG	A080293 001
DELTASONE		
+ PHARMACIA AND UPJOHN	2.5MG **	N009986 005
+	5MG **	N009986 002
+	10MG **	N009986 006
+	20MG **	N009986 007
+	50MG **	N009986 008
FERNISONE		
FERNDALE LABS	5MG	A083364 001
METICORTEN		
+ SCHERING	1MG **	N009766 002
+	5MG **	N009766 001
ORASONE		
SOLVAY	1MG	A083009 001
	5MG	A083009 002
	10MG	A083009 003
	20MG	A083009 004
	50MG	A085999 001
PARACORT		
PARKE DAVIS	5MG	N010962 002
PREDNICEN-M		
SCHWARZ PHARMA	5MG	A084655 001
PREDNISONE		
AM THERAP	5MG	A089387 001 Nov 06, 1986
	10MG	A089388 001 Nov 06, 1986
	20MG	A089389 001 Nov 06, 1986
AMNEAL PHARMS NY	5MG	A089597 001 Oct 05, 1987
	10MG	A089598 001 Oct 05, 1987
	20MG	A089599 001 Oct 05, 1987
AUROLIFE PHARMA LLC	5MG	A084774 001
	10MG	A089983 001 Jan 12, 1989
	20MG	A085813 001
	50MG	A089984 001 Jan 12, 1989
BUNDY	5MG	A083676 001
CHARTWELL RX	5MG	A083059 001
CONTRACT PHARMACAL	5MG	A080209 001
DURAMED PHARMS BARR	5MG	A088394 001 Oct 04, 1983
	10MG	A088395 001 Oct 04, 1983
	20MG	A088396 001 Oct 04, 1983
ELKINS SINK	5MG	A080491 001
	20MG	A085811 001
EVERYLIFE	1MG	A084440 001
	2.5MG	A084440 002
	5MG	A084440 003
FERRANTE	2.5MG	A080563 001
	5MG	A080563 002
HALSEY	5MG	A080300 001
HEATHER	5MG	A080320 001
	10MG	A084341 001
	20MG	A084417 001
	20MG	A085543 001
	50MG	A086946 001

**DISCONTINUED DRUG PRODUCT LIST**

6-303(of 375)

\*\* See List Footnote

PREDNISONE

TABLET;ORAL

## PREDNISONE

HIKMA PHARMS	1MG	A040890 001	Nov 01, 2010
IMPAKX LABS	5MG	A080782 001	
INWOOD LABS	1MG	A080328 001	
	2.5MG	A080306 001	
	5MG	A080279 001	
IVAX SUB TEVA PHARMS	5MG	A080283 001	
	10MG	A084133 001	
	20MG	A084134 001	
KV PHARM	5MG	A084236 001	
LANNETT	5MG	A080514 001	
	20MG	A084275 001	
LEDERLE	5MG	A086968 001	
MARSHALL PHARMA	5MG	A080301 001	
MUTUAL PHARM	5MG	A080701 001	
	10MG	A086595 001	
	20MG	A084634 001	
NYLOS	5MG	A085115 001	
PANRAY	1MG	A080350 001	
	2.5MG	A080350 002	
	5MG	A080350 003	
PHARMAVITE	5MG	A084662 002	
PHOENIX LABS NY	5MG	A080321 001	
	20MG	A083807 001	
PUREPAC PHARM	5MG	A080353 001	
	10MG	A086062 001	
	20MG	A086061 001	
PVT FORM	20MG	A085151 001	
REXALL	5MG	A080232 001	
ROXANE	20MG	N017109 001	
	25MG	A087833 001	May 04, 1982
SANDOZ	5MG	A080336 002	
SCHERER LABS	5MG	A080371 001	
SPERTI	1MG	A080359 001	
	2.5MG	A080359 002	
	5MG	A080359 003	
SUN PHARM INDUSTRIES	50MG	A086596 001	
SUPERPHARM	5MG	A088865 001	Oct 25, 1984
	10MG	A088866 001	Oct 25, 1984
	20MG	A088867 001	Oct 25, 1984
TEVA	5MG	A080397 001	
UDL	5MG	A087984 001	Jan 18, 1983
	10MG	A087985 001	Jan 18, 1983
	20MG	A087986 001	Jan 18, 1983
UPSHER SMITH	5MG	A087471 001	
	20MG	A087470 001	
VALEANT PHARM INTL	5MG	A080237 001	
VANGARD	5MG	A087682 001	Jan 15, 1982
	20MG	A087701 001	Jan 15, 1982
VITARINE	5MG	A080334 001	
	5MG	A080506 001	
WATSON LABS	5MG	A085084 002	
	10MG	A087773 001	Jul 13, 1982
	20MG	A086813 001	
	50MG	A086867 001	
	50MG	A087772 001	Jul 13, 1982
WHITEWORTH TOWN PLSN	2.5MG	A084913 001	
	5MG	A080343 001	
	10MG	A089028 001	Jul 24, 1986
	20MG	A084913 002	
SERVISONE			
LEDERLE	5MG	A080223 001	

**DISCONTINUED DRUG PRODUCT LIST**

6-304(of 375)

\*\* See List Footnote

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST

+ ASTRazeneca	1% **	N014763 004
+	2% **	N014763 005
+	3% **	N014763 003
CITANEST PLAIN		
+ ASTRazeneca	4% **	N014763 007
CITANEST PLAIN DENTAL		
DENTSPLY PHARM	4%	N021382 001

PRIMIDONE

SUSPENSION; ORAL

MYSOLINE

NURO PHARMA	250MG/5ML	N010401 001
-------------	-----------	-------------

TABLET; ORAL

PRIMIDONE

DR REDDYS LABS LTD	50MG	A040862 001 Oct 03, 2008
	250MG	A040862 002 Oct 03, 2008
HIKMA INTL PHARMS	50MG	A040667 001 Jul 27, 2006
IMPAX LABS	50MG	A040717 001 Feb 12, 2008
	250MG	A040717 002 Feb 12, 2008
WATSON LABS	250MG	A085052 001

PROBENECID

TABLET; ORAL

BENEMID

+ MERCK	500MG **	N007898 004
---------	----------	-------------

PROBENECID

IVAX SUB TEVA PHARMS	500MG	A083740 001 May 09, 1984
LEDERLE	500MG	A086917 001
WATSON LABS	500MG	A086150 002 Apr 23, 1982

PROBUCOL

TABLET; ORAL

LORELCO

SANOFI AVENTIS US	250MG	N017535 001
	500MG	N017535 002 Jul 06, 1988

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ASCOT	250MG	A087542 001 Jan 08, 1982
	375MG	A087697 001 Mar 01, 1983
	500MG	A087543 001 Jan 08, 1982
IDT AUSTRALIA LTD	250MG	A089219 001 Jul 01, 1986
	375MG	A089219 002 Jul 01, 1986
	500MG	A089219 003 Jul 01, 1986
IVAX SUB TEVA PHARMS	250MG	A084604 001
	375MG	A084595 001
	500MG	A084606 001
LANNETT	250MG	A083693 001
	500MG	A084696 001
LEDERLE	250MG	A086942 001
	375MG	A086952 001
	500MG	A086943 001
ROXANE	250MG	A088989 001 Apr 26, 1985
	500MG	A088990 001 Apr 26, 1985
VANGARD	250MG	A087643 001 Jun 01, 1982
	500MG	A087875 001 Jun 01, 1982
WATSON LABS	250MG	A083287 001
	250MG	A083795 001
	250MG	A085167 001
	375MG	A084403 001
	375MG	A087020 001
	500MG	A084280 001
	500MG	A084357 001
	500MG	A087021 001
PROCAN		
PARKE DAVIS	250MG	A085804 001
	375MG	A087502 001

**DISCONTINUED DRUG PRODUCT LIST**

6-305(of 375)

\*\* See List Footnote

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAN

500MG

A085079 001

PROCAPAN

PANRAY

250MG

A083553 002

PRONESTYL

+ APOTHECON

250MG \*\*

N007335 001

+

375MG \*\*

N007335 004

+

500MG \*\*

N007335 003

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

ABRAXIS PHARM

100MG/ML

A089415 001 Nov 17, 1986

500MG/ML

A089416 001 Nov 17, 1986

HOSPIRA

500MG/ML

A089537 001 Aug 25, 1987

INTL MEDICATION

500MG/ML

A088637 001 Jul 31, 1984

PHARMAFAIR

100MG/ML

A088824 001 Nov 20, 1985

500MG/ML

A088830 001 Nov 20, 1985

SMITH AND NEPHEW

100MG/ML

A088530 001 Mar 04, 1985

500MG/ML

A088531 001 Mar 04, 1985

SOLOPAK

500MG/ML

A088532 001 Mar 04, 1985

WARNER CHILCOTT

100MG/ML

A088528 001 May 03, 1988

500MG/ML

A089529 001 May 03, 1988

WATSON LABS

100MG/ML

A087079 001

500MG/ML

A087080 001

WEST-WARD PHARMS INT

100MG/ML

A089029 001 Apr 17, 1986

500MG/ML

A089030 001 Apr 17, 1986

PRONESTYL

+ APOTHECON

100MG/ML \*\*

N007335 002

+

500MG/ML \*\*

N007335 005

TABLET; ORAL

PRONESTYL

APOTHECON

250MG

N017371 001

375MG

N017371 002

500MG

N017371 003

TABLET, EXTENDED RELEASE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ANI PHARMS INC

250MG

A088958 001 Dec 02, 1985

500MG

A088959 001 Dec 02, 1985

500MG

A088974 001 Jul 22, 1985

750MG

A089438 001 Mar 23, 1987

1GM

A040111 001 Dec 13, 1996

IDT AUSTRALIA LTD

250MG

A089369 001 Aug 14, 1987

500MG

A089369 002 Jan 09, 1987

750MG

A089369 003 Aug 14, 1987

INWOOD LABS

500MG

A089840 001 Mar 06, 1989

SANDOZ

500MG

A089284 001 Jun 23, 1986

WATSON LABS

250MG

A088533 001 Dec 03, 1984

250MG

A089026 001 Oct 22, 1985

500MG

A088534 001 Dec 03, 1984

500MG

A089027 001 Oct 22, 1985

750MG

A088535 001 Nov 03, 1984

750MG

A089042 001 Oct 22, 1985

1GM

A089520 001 Jan 15, 1987

PROCAN SR

PARKE DAVIS

250MG

A086468 001

PARKEDALE

500MG

A086065 001

750MG

A087510 001 Apr 01, 1982

1GM

A088489 001 Jan 16, 1985

PROCANBID

KING PHARMS

500MG

N020545 001 Jan 31, 1996

1GM

N020545 002 Jan 31, 1996

PRONESTYL-SR

APOTHECON

500MG

A087361 001

**DISCONTINUED DRUG PRODUCT LIST**

6-306(of 375)

\*\* See List Footnote

PROCAINE HYDROCHLORIDEINJECTABLE; INJECTION  
NOVOCAIN

HOSPIRA	1%	A085362 003
	2%	A085362 004
	10%	A086797 001
<b>PROCAINE HYDROCHLORIDE</b>		
ABRAXIS PHARM	1%	A080384 002
	1%	A080421 001
	2%	A080384 003
	2%	A080421 002
BEL MAR	1%	A080711 001
	2%	A080756 001
ELKINS SINK	1%	A083315 001
	2%	A083315 002
GD SEARLE LLC	1%	A086202 001
	2%	A086202 002
HOSPIRA	1%	A080416 001
	2%	A080416 002
MILES	1%	A080415 001
	2%	A080415 002
WATSON LABS	1%	A080658 001
	1%	A083535 001
	2%	A080658 002
	2%	A083535 002

PROCAINE HYDROCHLORIDE; TETRACYCLINE HYDROCHLORIDEINJECTABLE; INJECTION  
ACHROMYCIN

LEDERLE	40MG/VIAL;100MG/VIAL	N050276 001
	40MG/VIAL;250MG/VIAL	N050276 003
<b>TETRACYN</b>		
PFIZER	40MG/VIAL;100MG/VIAL	A060285 002
	40MG/VIAL;250MG/VIAL	A060285 003

PROCAINE MERETHOXYLLINE; THEOPHYLLINEINJECTABLE; INJECTION  
DICURIN PROCAINE

LILLY	100MG/ML;50MG/ML	N008869 001
-------	------------------	-------------

PROCHLORPERAZINESUPPOSITORY; RECTAL  
COMPATINE

GLAXOSMITHKLINE	2.5MG **	N011127 003
	5MG **	N011127 001
	25MG **	N011127 002
<b>PROCHLORPERAZINE</b>		
ABLE	2.5MG	A040407 001 Jul 11, 2001
	5MG	A040407 002 Jul 11, 2001
	25MG	A040407 003 Jul 11, 2001

PROCHLORPERAZINE EDISYLADE

CONCENTRATE; ORAL

COMPATINE		
GLAXOSMITHKLINE	EQ 10MG BASE/ML	N011276 001
<b>PROCHLORPERAZINE</b>		
ALPHARMA US PHARMS	EQ 10MG BASE/ML	A087153 001 Jun 08, 1982
<b>PROCHLORPERAZINE EDISYLADE</b>		
MORTON GROVE	EQ 10MG BASE/ML	A088598 001 Oct 25, 1984
<b>INJECTABLE; INJECTION</b>		
COMPATINE		
+ GLAXOSMITHKLINE	EQ 5MG BASE/ML **	N010742 002
<b>PROCHLORPERAZINE</b>		
BAXTER HLTHCARE	EQ 5MG BASE/ML	A087759 001 Oct 01, 1982
PROCHLORPERAZINE EDISYLADE		
ATHENEX INC	EQ 5MG BASE/ML	A040540 001 May 28, 2004
HOSPIRA	EQ 5MG BASE/ML	A089703 001 Apr 07, 1988
MARSAM PHARMS LLC	EQ 5MG BASE/ML	A089675 001 Dec 05, 1988
SMITH AND NEPHEW	EQ 5MG BASE/ML	A089251 001 Dec 04, 1986
TEVA PARENTERAL	EQ 5MG BASE/ML	A040505 001 May 30, 2003
WATSON LABS	EQ 5MG BASE/ML	A089530 001 Jul 08, 1987

**DISCONTINUED DRUG PRODUCT LIST**

6-307(of 375)

\*\* See List Footnote

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

WEST-WARD PHARMS INT	EQ 5MG BASE/ML	A089605 001 Jul 08, 1987
WYETH AYERST	EQ 5MG BASE/ML	A089606 001 Jul 08, 1987
	EQ 5MG BASE/ML	A089523 001 May 03, 1988
	EQ 5MG BASE/ML	A086348 001
SYRUP; ORAL COMPATZINE		
GLAXOSMITHKLINE	EQ 5MG BASE/5ML	N011188 001
PROCHLORPERAZINE EDISYLATE		
ALPHARMA US PHARMS	EQ 5MG BASE/5ML	A087154 001 Sep 01, 1982
MORTON GROVE	EQ 5MG BASE/5ML	A088597 001 Oct 25, 1984

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPATZINE

GLAXOSMITHKLINE	EQ 10MG BASE	N011000 001
	EQ 10MG BASE	N021019 001 Oct 06, 1999
	EQ 15MG BASE	N011000 002
	EQ 15MG BASE	N021019 002 Oct 06, 1999
	EQ 30MG BASE	N011000 003
	EQ 75MG BASE	N011000 004

TABLET; ORAL

COMPATZINE

GLAXOSMITHKLINE	EQ 5MG BASE **	N010571 001
	EQ 10MG BASE **	N010571 002
	EQ 25MG BASE **	N010571 003

PROCHLORPERAZINE

WATSON LABS	EQ 5MG BASE	A085580 001
	EQ 10MG BASE	A085178 001
	EQ 25MG BASE	A085579 001

PROCHLORPERAZINE MALEATE

DURAMED PHARMS BARR	EQ 5MG BASE	A040207 001 May 01, 1997
	EQ 5MG BASE	A089484 001 Jan 20, 1987
	EQ 10MG BASE	A040207 002 May 01, 1997
	EQ 10MG BASE	A089485 001 Jan 20, 1987
	EQ 25MG BASE	A089486 001 Jan 20, 1987
IVAX SUB TEVA PHARMS	EQ 5MG BASE	A040162 001 Jan 20, 1998
	EQ 10MG BASE	A040162 002 Jan 20, 1998
SANDOZ	EQ 25MG BASE	A040101 003 Jul 19, 1996

PROCYCLIDINE HYDROCHLORIDE

TABLET; ORAL

KEMADRIN

MONARCH PHARMS	2MG	N009818 005
	5MG	N009818 003

PROGESTERONE

CAPSULE; ORAL

PROMETRIUM

VIRTUS PHARMS	300MG	N019781 003 Oct 15, 1999
---------------	-------	--------------------------

INJECTABLE; INJECTION

PROGESTERONE

LILLY	25MG/ML	N009238 002
	50MG/ML	N009238 001

INSERT, EXTENDED RELEASE; INTRAUTERINE

PROGESTASERT

ALZA	38MG	N017553 001
------	------	-------------

PROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

SPARINE

WYETH AYERST	30MG/ML	N010942 001
	100MG/ML	N010942 004

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE

WATSON LABS	25MG/ML	A084510 001
	50MG/ML	A084517 001

SPARINE

BAXTER HLTHCARE CORP	25MG/ML	N010349 008
----------------------	---------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-308(of 375)

\*\* See List Footnote

PROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

SPARINE

50MG/ML

N010349 006

SYRUP; ORAL

SPARINE

WYETH AYERST

10MG/5ML

N010942 003

TABLET; ORAL

SPARINE

WYETH AYERST

10MG

N010348 006

25MG

N010348 001

50MG

N010348 002

100MG

N010348 003

200MG

N010348 004

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PHENERGAN

WYETH AYERST

25MG/ML

N008857 002

50MG/ML

N008857 003

PROMETHAZINE HYDROCHLORIDE

ABBOTT

25MG/ML

A084223 001

50MG/ML

A084222 001

AKORN

25MG/ML

A083955 002

50MG/ML

A083955 001

BEDFORD LABS

25MG/ML

A040524 001 Mar 17, 2004

50MG/ML

A040524 002 Mar 17, 2004

HOSPIRA

25MG/ML

A040372 001 Jun 08, 2000

50MG/ML

A040372 002 Jun 08, 2000

50MG/ML

A083838 002

LUITPOLD

25MG/ML

A040515 001 Mar 19, 2003

MARSAM PHARMS LLC

25MG/ML

A089463 001 May 02, 1988

50MG/ML

A089477 001 May 02, 1988

MYLAN INSTITUTIONAL

25MG/ML

A040471 001 Nov 21, 2002

SANDOZ

25MG/ML

A040593 001 Nov 08, 2006

50MG/ML

A040593 002 Nov 08, 2006

TEVA PHARMS USA

25MG/ML \*\*

A040454 001 Aug 22, 2002

50MG/ML \*\*

A040454 002 Aug 22, 2002

WATSON LABS

25MG/ML

A083532 001

25MG/ML

A084591 001

50MG/ML

A080629 002

50MG/ML

A083532 002

WOCKHARDT

25MG/ML

A040785 001 Sep 26, 2008

50MG/ML

A040785 002 Sep 26, 2008

ZIPAN-25

ALTANA

25MG/ML

A083997 001

ZIPAN-50

ALTANA

50MG/ML

A083997 002

SUPPOSITORY; RECTAL

PHENERGAN

+ MYLAN PHARMS INC

12.5MG \*\*

N010926 002

+

25MG \*\*

N010926 001

+

50MG \*\*

N011689 001

PROMETHACON

POLYMEDICA

25MG

A084901 001

50MG

A084902 001

PROMETHAZINE HYDROCHLORIDE

ABLE

12.5MG

A040504 001 Apr 11, 2003

25MG

A040504 002 Apr 11, 2003

50MG

A040449 001 Feb 27, 2003

SYRUP; ORAL

MYMETHAZINE FORTIS

USL PHARMA

25MG/5ML

A087996 001 Jan 18, 1983

PROMETH FORTIS

ALPHARMA US PHARMS

25MG/5ML

A084772 001

PROMETH PLAIN

ACTAVIS MID ATLANTIC

6.25MG/5ML

A085953 001

PROMETHAZINE

CENCI

6.25MG/5ML

A089013 001 Sep 20, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-309(of 375)

\*\* See List Footnote

PROMETHAZINE HYDROCHLORIDE

SYRUP;ORAL

PROMETHAZINE HYDROCHLORIDE

KV PHARM	6.25MG/5ML	A085388 001
	25MG/5ML	A085385 001
PHARM ASSOC	6.25MG/5ML	A087518 001
WHITEWORTH TOWN PLSN	6.25MG/5ML	A086395 001
PROMETHAZINE HYDROCHLORIDE	PLAIN	
+ ANI PHARMS	6.25MG/5ML **	N008381 004 Apr 18, 1984
+	25MG/5ML **	N008381 003

TABLET;ORAL

PHENERGAN

+ DELCOR ASSET CORP	12.5MG **	N007935 002
+	25MG **	N007935 003
+	50MG **	N007935 004

PROMETHAZINE HYDROCHLORIDE

ABBOTT	12.5MG	A084160 001
	25MG	A084166 001
	50MG	A084539 001
ABLE	12.5MG	A040558 001 Jul 01, 2004
	25MG	A040558 002 Jul 01, 2004
	50MG	A040558 003 Jul 01, 2004
IMPAX LABS	25MG	A084214 002 Jul 07, 1982
	50MG	A040791 001 May 20, 2008
IVAX SUB TEVA PHARMS	12.5MG	A083604 001
	25MG	A083603 001
	50MG	A083613 001
LANNETT	12.5MG	A080949 001
	25MG	A080949 002
	50MG	A080949 003
MYLAN	12.5MG	A091054 001 Aug 30, 2011
	25MG	A091054 002 Aug 30, 2011
	50MG	A091054 003 Aug 30, 2011
PVT FORM	12.5MG	A083214 001
	25MG	A083658 001
SANDOZ	12.5MG	A084176 002 May 22, 2009
	12.5MG	A084233 001
	25MG	A085146 001
	50MG	A085146 002
SUN PHARM INDUSTRIES	12.5MG	A084555 001
	25MG	A084554 001
	50MG	A084557 001
TABLICAPS	12.5MG	A084080 001
	25MG	A084027 001
TEVA	25MG	A089109 001 Sep 10, 1985
WATSON LABS	12.5MG	A083401 001
	12.5MG	A083712 001
	25MG	A085986 001
	25MG	A083204 001
	50MG	A085684 001
	50MG	A083403 001
	50MG	A085664 001
REMSED		
BRISTOL MYERS SQUIBB	25MG	A083176 002
	50MG	A083176 001

PROPAFENONE HYDROCHLORIDE

TABLET;ORAL

PROPAFENONE HYDROCHLORIDE

NESHER PHARMS	150MG	A076193 001 Feb 07, 2002
	225MG	A076193 002 Feb 07, 2002
	300MG	A076193 003 Feb 07, 2002

PROPANTHELINE BROMIDE

INJECTABLE; INJECTION

PRO-BANTHINE

GD SEARLE LLC	30MG/VIAL	N008843 001
---------------	-----------	-------------

TABLET;ORAL

PRO-BANTHINE

+ SHIRE	7.5MG **	N008732 003
---------	----------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-310(of 375)

\*\* See List Footnote

PROPANTHELINE BROMIDE

TABLET;ORAL		
PRO-BANTHINE		
+ 15MG **		N008732 002
PROPANTHELINE BROMIDE		
ASCOT 15MG		A087663 001 Oct 25, 1982
HEATHER 15MG		A085780 001
IMPAX LABS 15MG		A084541 002
MYLAN 15MG		A083706 001
PAR PHARM 15MG		A088377 001 Dec 08, 1983
PVT FORM 15MG		A080977 001
SANDOZ 15MG		A080928 001
TABLICAPS 15MG		A084428 001
WATSON LABS 15MG		A083029 002
WEST-WARD PHARMS INT 15MG		A083151 001
WEST-WARD PHARMS INT 7.5MG		A080927 001

PROPARACAIN HYDROCHLORIDE

SOLUTION/DROPS;OPHTHALMIC		
KAINAIR		
PHARMAFAIR 0.5%		A088087 001 Jun 07, 1983
OPHTHAINE		
+ APOTHECON 0.5% **		N008883 001
OPHTHETIC		
+ ALLERGAN 0.5% **		N012583 001
PARACAIN		
OPTOPICS 0.5%		A087681 001 Aug 05, 1982
PROPARACAIN HYDROCHLORIDE		
SOLA BARNES HIND 0.5%		A084144 001
	0.5%	A084151 001

PROPIOLACTONE

SOLUTION;IRRIGATION		
BETAPRONE		
FOREST LABS N/A		N011657 001

PROTIOMAZINE HYDROCHLORIDE

INJECTABLE;INJECTION		
LARGON		
WEST-WARD PHARMS INT 20MG/ML		N012382 002

PROPOFOL

INJECTABLE;INJECTION		
DIPRIVAN		
FRESENIUS KABI USA 10MG/ML		N019627 001 Oct 02, 1989
PROPOFOL		
TEVA PARENTERAL 10MG/ML		A075392 001 Sep 19, 2000
WEST-WARD PHARMS INT 10MG/ML		A074848 001 Apr 19, 2005

PROPOXYPHENE HYDROCHLORIDE

CAPSULE;ORAL		
DARVON		
XANODYNE PHARM 32MG		N010997 001
	65MG	N010997 003
DOLENE		
HERITAGE PHARMS INC 65MG		A080530 001
KESSO-GESIC		
MK LABS 65MG		A083544 001
PROPHENE 65		
HALSEY 65MG		A083538 002
PROPOXYPHENE HYDROCHLORIDE		
ALRA 65MG		A083184 001
IMPAX LABS 65MG		A083317 001
IVAX SUB TEVA PHARMS 32MG		A083597 001
MUTUAL PHARM 65MG		A083186 001
MYLAN 32MG		A083528 001
	65MG	A040569 001 Dec 16, 2004
	65MG	A083299 001
NEXGEN PHARMA INC 65MG		A083185 001
PAR PHARM 65MG		A080269 001
PUREPAC PHARM 65MG		A083278 001
PVT FORM 32MG		A083464 001

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-311(of 375)

\*\* See List Footnote

PROPOXYPHENONE HYDROCHLORIDE

CAPSULE; ORAL

PROPOXYPHENONE HYDROCHLORIDE

	65MG	A083113 001
ROXANE	32MG	A083089 001
	65MG	A083089 002
SANDOZ	32MG	A084014 001
	65MG	A083125 002
	65MG	A083688 001
	65MG	A083870 002
	65MG	A086495 001
TEVA	65MG	A088615 001 Oct 22, 1984
VALEANT PHARM INTL	65MG	A080783 001
VINTAGE PHARMS	65MG	A040908 001 Jul 17, 2009
WATSON LABS	65MG	A080908 002
	65MG	A085190 001
WEST WARD	65MG	A083501 001
WHITEWORTH TOWN PLSN	65MG	A084551 001
PROPOXYPHENONE HYDROCHLORIDE	65	
WARNER CHILCOTT	65MG	A083786 001

PROPOXYPHENONE NAPSYLATE

SUSPENSION; ORAL

DARVON-N

AAIPHARMA LLC 50MG/5ML

N016861 001

TABLET; ORAL

DARVON-N

XANODYNE PHARM 100MG

N016862 002

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PROPRANOLOL HYDROCHLORIDE

INWOOD LABS	60MG	A072499 001 Apr 11, 1989
	80MG	A072500 001 Apr 11, 1989
	120MG	A072501 001 Apr 11, 1989
	160MG	A072502 001 Apr 11, 1989
UPSHER-SMITH LABS	60MG	A078311 001 Mar 06, 2009
	80MG	A078311 002 Mar 06, 2009
	120MG	A078311 003 Mar 06, 2009
	160MG	A078311 004 Mar 06, 2009

CONCENTRATE; ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL  
ROXANE 80MG/ML

A071388 001 May 15, 1987

INJECTABLE; INJECTION

PROPRANOLOL HYDROCHLORIDE

SANDOZ INC	1MG/ML	A076400 001 Feb 26, 2003
SMITH AND NEPHEW	1MG/ML	A070135 001 Apr 15, 1986
	1MG/ML	A070137 001 Apr 15, 1986
SOLOPAK	1MG/ML	A070136 001 Apr 15, 1986

SOLUTION; ORAL

PROPRANOLOL HYDROCHLORIDE

MORTON GROVE	20MG/5ML	A071984 001 Mar 03, 1989
	40MG/5ML	A071985 001 Mar 03, 1989

SUSPENSION; ORAL

INDERAL

WYETH AYERST 10MG/ML

N019536 001 Dec 12, 1986

TABLET; ORAL

INDERAL

+ WYETH PHARMS INC	10MG **	N016418 001
+	20MG **	N016418 003
+	40MG **	N016418 002
+	60MG **	N016418 009 Oct 18, 1982
+	80MG **	N016418 004
+	90MG **	N016418 010 Oct 18, 1982

PROPRANOLOL HYDROCHLORIDE ANDA REPOSITORY	10MG	A070319 001 Oct 22, 1985
	20MG	A070320 001 Oct 22, 1985
	40MG	A070103 001 Oct 22, 1985
	60MG	A070321 001 Sep 24, 1986
	80MG	A070322 001 Aug 04, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-312(of 375)

\*\* See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET;ORAL

## PROPRANOLOL HYDROCHLORIDE

ANI PHARMS INC	90MG	A071977 001	Apr 06, 1988
DAVA PHARMS INC	10MG	A070125 001	Jul 30, 1985
	20MG	A070126 001	Jul 30, 1985
	40MG	A070127 001	Jul 30, 1985
	60MG	A071495 001	Dec 31, 1987
	80MG	A070128 001	Jul 30, 1985
	90MG	A071496 001	Dec 31, 1987
DURAMED PHARMS BARR	10MG	A070306 001	Sep 09, 1985
	20MG	A070307 001	Sep 09, 1985
	40MG	A070308 001	Sep 09, 1985
	60MG	A070309 001	Oct 01, 1986
	80MG	A070310 001	Sep 09, 1985
	90MG	A071327 001	Oct 01, 1986
INTERPHARM	10MG	A071368 001	May 05, 1987
	20MG	A071369 001	May 05, 1987
	40MG	A071370 001	May 05, 1987
	80MG	A071371 001	May 05, 1987
IVAX SUB TEVA PHARMS	10MG	A072063 001	Jul 29, 1988
	20MG	A072066 001	Jul 29, 1988
	40MG	A072067 001	Jul 29, 1988
	60MG	A072068 001	Jul 29, 1988
	80MG	A072069 001	Jul 29, 1988
LEDERLE	10MG	A072117 001	Jun 23, 1988
	20MG	A072118 001	Jun 23, 1988
	40MG	A072119 001	Jun 23, 1988
	80MG	A072120 001	Jun 23, 1988
MYLAN	60MG	A072275 001	Jun 09, 1989
PAR PHARM	90MG	A071288 001	Oct 22, 1986
PUREPAC PHARM	10MG	A070814 001	Nov 03, 1986
	20MG	A070815 001	Nov 03, 1986
	40MG	A070816 001	Nov 03, 1986
	60MG	A070817 001	Nov 03, 1986
	80MG	A070757 001	Nov 03, 1986
ROXANE	10MG	A070516 001	Jul 07, 1986
	20MG	A070517 001	Jul 07, 1986
	40MG	A070518 001	Jul 07, 1986
	60MG	A070519 001	Sep 24, 1986
	80MG	A070520 001	Jul 07, 1986
	90MG	A070521 001	Sep 24, 1986
SANDOZ	10MG	A070663 001	Jun 13, 1986
	10MG	A071658 001	Jul 05, 1988
	20MG	A070664 001	Jun 13, 1986
	20MG	A071687 001	Jul 05, 1988
	40MG	A070665 001	Jun 13, 1986
	40MG	A071688 001	Jul 05, 1988
	60MG	A070666 001	Oct 10, 1986
	60MG	A072197 001	Jul 05, 1988
	80MG	A070667 001	Jun 13, 1986
	80MG	A071689 001	Jul 05, 1988
	90MG	A072198 001	Jul 05, 1988
SCHERING	10MG	A070120 001	Aug 06, 1985
	20MG	A070121 001	Aug 06, 1985
	40MG	A070122 001	Aug 06, 1985
	60MG	A070123 001	Oct 29, 1986
	80MG	A070124 001	Aug 06, 1985
SUPERPHARM	10MG	A071515 001	Jun 08, 1988
	20MG	A071516 001	Jun 08, 1988
	40MG	A071517 001	Jun 08, 1988
	80MG	A071518 001	Jun 08, 1988
TEVA	10MG	A070232 001	Oct 07, 1987
	20MG	A070233 001	Jun 23, 1986
	40MG	A070234 001	Jun 23, 1986
WARNER CHILCOTT	10MG	A070438 001	Sep 15, 1986
	20MG	A070439 001	Sep 15, 1986
	40MG	A070440 001	Sep 15, 1986
	60MG	A070441 001	Sep 24, 1986

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-313(of 375)

\*\* See List Footnote

PROPRANOLOL HYDROCHLORIDE

TABLET;ORAL

PROPRANOLOL HYDROCHLORIDE

	80MG	A070442 001	Sep 15, 1986
WATSON LABS	10MG	A070140 001	Jul 30, 1985
	10MG	A070378 001	Mar 19, 1987
	20MG	A070141 001	Jul 30, 1985
	20MG	A070379 001	Mar 19, 1987
	40MG	A070142 001	Jul 30, 1985
	40MG	A070380 001	Mar 19, 1987
	60MG	A070143 001	Jan 15, 1987
	60MG	A070381 001	Mar 19, 1987
	60MG	A071098 001	Oct 06, 1986
	60MG	A071791 001	Jul 15, 1987
	80MG	A070144 001	Jul 30, 1985
	80MG	A070382 001	Mar 19, 1987
	80MG	A070551 001	Jul 10, 1986
	90MG	A071183 001	Oct 06, 1986
	90MG	A071792 001	Jul 15, 1987
WATSON LABS TEVA	10MG	A070548 001	Jul 10, 1986
	20MG	A070549 001	Apr 11, 1986
	40MG	A070550 001	Apr 11, 1986

PROPYLIODONE

SUSPENSION;INTRATRACHEAL

DIONOSIL AQUEOUS

GLAXOSMITHKLINE	50%	N009309 001
DIONOSIL OILY	60%	N009309 002

PROPYLTHIOURACIL

TABLET;ORAL

PROPYLTHIOURACIL

ABBOTT	50MG	A084075 001
ANABOLIC	50MG	A080285 001
ANI PHARMS INC	50MG	A080215 001
CHARTWELL RX	50MG	A084543 001
HALSEY	50MG	A080015 001
HIKMA INTL PHARMS	50MG	A080154 001
IMPAX LABS	50MG	A080159 001
LANNETT	50MG	A080016 001
LILLY	50MG	N006213 001
SUN PHARM INDUSTRIES	50MG	A083982 001
TABLICAPS	50MG	A080840 001
WATSON LABS	50MG	A080932 001
	50MG	A085201 001

PROTAMINE SULFATE

INJECTABLE;INJECTION

PROTAMINE SULFATE

+ LILLY	10MG/ML **	N006460 002
PHARMACIA AND UPJOHN	50MG/VIAL	N007413 001
	250MG/VIAL	N007413 002 Aug 02, 1984
WEST-WARD PHARMS INT	10MG/ML	A089474 001 Nov 05, 1986
	10MG/ML	A089475 001 Nov 05, 1986

PROTEIN HYDROLYSATE

INJECTABLE;INJECTION

AMINOSOL 5%

ABBVIE	5%	N005932 012 Jan 31, 1985
HYPROTIGEN 5%		N006170 003 Jan 10, 1984

B BRAUN

5%

PROTIRELIN

INJECTABLE;INJECTION

THYPINONE

ABBOTT	0.5MG/ML	N017638 001
THYREL TRH		N018087 001

FERRING

0.5MG/ML

**DISCONTINUED DRUG PRODUCT LIST**

6-314(of 375)

\*\* See List Footnote

PROTOKYLOL HYDROCHLORIDE

TABLET;ORAL  
VENTAIRE  
SANOFI AVENTIS US 2MG A083459 001

PROTRIPTYLINE HYDROCHLORIDE

TABLET;ORAL  
VIVACTIL  
TEVA WOMENS 5MG \*\* N016012 001  
10MG \*\* N016012 002

PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL  
NOVAFED  
SANOFI AVENTIS US 120MG N017603 001  
SUDAFED 12 HOUR  
+ GLAXOSMITHKLINE 120MG \*\* N017941 002

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL  
ACTIFED  
GLAXOSMITHKLINE 120MG;5MG N018996 001 Jun 17, 1985  
TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES  
KV PHARM 120MG;5MG A071798 001 Mar 16, 1989  
SYRUP;ORAL  
ACTAHIST  
CENCI 30MG/5ML;1.25MG/5ML A088344 001 Feb 09, 1984  
HISTAFED  
CENCI 30MG/5ML;1.25MG/5ML A088283 001 Apr 20, 1984  
MYFED  
USL PHARMA 30MG/5ML;1.25MG/5ML A088116 001 Mar 04, 1983  
TRILITRON  
NEWTRON PHARMS 30MG/5ML;1.25MG/5ML A088474 001 Feb 12, 1985  
TABLET;ORAL  
ALLERFED  
PVT FORM 60MG;2.5MG A088860 001 Jan 31, 1985  
CORPHED  
SANDOZ 60MG;2.5MG A088602 001 Apr 11, 1985  
PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE  
SANDOZ 60MG;2.5MG A088193 001 May 17, 1983  
TRILITRON  
NEWTRON PHARMS 60MG;2.5MG A088515 001 Jan 09, 1985  
TRIPHED  
TEVA 60MG;2.5MG A088630 001 May 17, 1984  
TRIPROLIDINE AND PSEUDOEPHEDRINE  
WATSON LABS 60MG;2.5MG A088318 002 Jan 13, 1984  
WEST WARD 60MG;2.5MG A088117 001 Apr 19, 1983  
TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE  
IVAX SUB TEVA PHARMS 60MG;2.5MG A085273 001 Dec 12, 1984  
SUPERPHARM 60MG;2.5MG A088578 001 Feb 21, 1985  
TABLET, EXTENDED RELEASE;ORAL  
TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES  
KV PHARM 120MG;5MG A072758 001 Nov 25, 1991

PSEUDOEPHEDRINE POLISTIREX

SUSPENSION, EXTENDED RELEASE;ORAL  
PSEUDO-12  
UCB INC EQ 60MG HCL/5ML N019401 001 Jun 19, 1987

PYRIDOSTIGMINE BROMIDE

TABLET;ORAL  
PYRIDOSTIGMINE BROMIDE  
ANI PHARMS INC 30MG A040512 002 Jul 20, 2005  
60MG A040512 001 Oct 08, 2003  
IMPAX LABS INC 60MG A040457 001 Dec 26, 2002  
SOLVAY 30MG A089572 001 Nov 27, 1990  
US ARMY 30MG N020414 001 Feb 05, 2003

**DISCONTINUED DRUG PRODUCT LIST**

6-315(of 375)

\*\* See List Footnote

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

HEXA-BETALIN

LILLY

100MG/ML

A080854 001

PYRIDOXINE HYDROCHLORIDE

AKORN

100MG/ML

A087967 001 Oct 01, 1982

BEL MAR

100MG/ML

A080761 001

DELL LABS

50MG/ML

A083771 001

100MG/ML

A083772 001

ELKINS SINK

100MG/ML

A080581 001

LUITPOLD

100MG/ML

A080669 001

WATSON LABS

100MG/ML

A080572 001

100MG/ML

A083760 001

PYRILAMINE MALEATE

TABLET; ORAL

PYRILAMINE MALEATE

IMPAK LABS

25MG

A080808 001

WATSON LABS

25MG

A085231 001

PYRIMETHAMINE; SULFADOXINE

TABLET; ORAL

FANSIDAR

ROCHE

25MG; 500MG

N018557 001

PYRITHIONE ZINC

LOTION; TOPICAL

HEAD &amp; SHOULDERS CONDITIONER

WARNER CHILCOTT 0.3%

N019412 002 Mar 10, 1986

PYRVINIUM PAMOATE

SUSPENSION; ORAL

POVAN

PARKE DAVIS

EQ 50MG BASE/5ML

N011964 001

TABLET; ORAL

POVAN

PARKE DAVIS

EQ 50MG BASE

N012485 002

QUAZEPAM

TABLET; ORAL

DORAL

CUTIS HEALTH LLC

7.5MG

N018708 003 Feb 26, 1987

QUETIAPINE FUMARATE

TABLET; ORAL

QUETIAPINE FUMARATE

ACTAVIS GRP PTC

EQ 25MG BASE

A201762 001 Feb 27, 2013

EQ 50MG BASE

A201762 002 Feb 27, 2013

EQ 100MG BASE

A201762 003 Feb 27, 2013

EQ 150MG BASE

A201762 004 Feb 27, 2013

EQ 200MG BASE

A201762 005 Feb 27, 2013

EQ 300MG BASE

A201762 006 Feb 27, 2013

EQ 400MG BASE

A201762 007 Feb 27, 2013

MYLAN PHARMS INC

EQ 25MG BASE

A090323 001 Mar 27, 2012

SEROQUEL

+ ASTRazeneca PHARMS

EQ 150MG BASE \*\*

N020639 004 Dec 20, 1998

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE

ACTAVIS ELIZABETH

EQ 5MG BASE

A076459 001 Dec 22, 2004

EQ 10MG BASE

A076459 002 Dec 22, 2004

EQ 20MG BASE

A076459 003 Dec 22, 2004

EQ 40MG BASE

A076459 004 Dec 22, 2004

ACTAVIS LABS FL INC

EQ 5MG BASE

A076049 001 Jan 14, 2005

EQ 10MG BASE

A076049 002 Jan 14, 2005

EQ 20MG BASE

A076049 003 Jan 14, 2005

EQ 40MG BASE

A076049 004 Jan 14, 2005

APOTEX INC

EQ 5MG BASE

A076240 001 Jan 26, 2006

EQ 10MG BASE

A076240 002 Jan 26, 2006

EQ 20MG BASE

A076240 003 Jan 26, 2006

EQ 40MG BASE

A076240 004 Jan 26, 2006

FOSUN PHARMA

EQ 5MG BASE

A076803 001 Mar 02, 2005

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-316(of 375)

\*\* See List Footnote

QUINAPRIL HYDROCHLORIDE

TABLET;ORAL

QUINAPRIL HYDROCHLORIDE

	EQ 10MG BASE	A076803 002 Mar 02, 2005
	EQ 20MG BASE	A076803 003 Mar 02, 2005
	EQ 40MG BASE	A076803 004 Mar 02, 2005
MYLAN	EQ 5MG BASE	A076036 001 Jan 28, 2005
	EQ 10MG BASE	A076036 002 Jan 28, 2005
	EQ 20MG BASE	A076036 003 Jan 28, 2005
	EQ 40MG BASE	A076036 004 Jan 28, 2005
SUN PHARM INDS LTD	EQ 5MG BASE	A090800 001 Jun 18, 2009
	EQ 10MG BASE	A090800 002 Jun 18, 2009
	EQ 20MG BASE	A090800 003 Jun 18, 2009
	EQ 40MG BASE	A090800 004 Jun 18, 2009

QUINESTROL

TABLET;ORAL

ESTROVIS

PARKE DAVIS	0.1MG	N016768 002
	0.2MG	N016768 003

QUINETHAZONE

TABLET;ORAL

HYDROMOX

LEDERLE	50MG	N013264 001
---------	------	-------------

QUINETHAZONE: RESERPINE

TABLET;ORAL

HYDROMOX R

LEDERLE	50MG; 0.125MG	N013927 001
---------	---------------	-------------

QUINIDINE GLUCONATE

INJECTABLE; INJECTION

QUINIDINE GLUCONATE

+ LILLY

80MG/ML N007529 002 Feb 10, 1989

TABLET;ORAL

QUINACT

BAYER HLTHCARE	266MG	A085978 001
	400MG	A086099 001

TABLET, EXTENDED RELEASE;ORAL

DURAQUIN

WARNER CHILCOTT	330MG	N017917 001
-----------------	-------	-------------

QUINAGLUTE

BAYER HLTHCARE	324MG	N016647 001
----------------	-------	-------------

QUINALAN

LANNETT 324MG A088081 001 Feb 10, 1986

QUINATIME

WATSON LABS	324MG	A087448 001
-------------	-------	-------------

QUINIDINE GLUCONATE

ASCOT	324MG	A088582 001 Jun 17, 1985
AUROLIFE PHARMA LLC	324MG	A089894 001 Dec 15, 1988
CYCLE PHARMS LTD	324MG	A088431 001 Jan 06, 1984
HALSEY	324MG	A089476 001 Apr 10, 1987
SUPERPHARM	324MG	A089164 001 Nov 21, 1985
WATSON LABS	324MG	A087785 001 Jan 24, 1983
	324MG	A087810 001 Sep 29, 1982

QUINIDINE POLYGALACTURONATE

TABLET;ORAL

CARDIOQUIN

PHARM RES ASSOC	275MG	N011642 002
-----------------	-------	-------------

QUINIDINE SULFATE

CAPSULE;ORAL

CIN-QUIN

SOLVAY	200MG	A085296 001
	300MG	A085297 001

QUINIDINE SULFATE

LILLY	200MG	A085103 001
-------	-------	-------------

TABLET;ORAL

CIN-QUIN

SOLVAY	100MG	A085299 001
--------	-------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-317(of 375)

\*\* See List Footnote

QUINIDINE SULFATETABLET;ORAL  
CIN-QUIN

QUINIDINE SULFATE			
BARR	200MG	A084932	001
CONTRACT PHARMACAL	300MG	A085298	001
CYCLE PHARMS LTD			
	200MG	A084177	001
	300MG	A083808	001
		A083640	001
	200MG	A085632	001
DAVA PHARMS INC	200MG	A087011	001
ELKINS SINN	200MG	A083622	001
EVERYLIFE	200MG	A083439	001
HALSEY	200MG	A083583	001
HIKMA PHARMS	200MG	A083862	001
IMPAKX LABS	200MG	A083347	001
IVAX SUB TEVA PHARMS	200MG	A084549	001
KING PHARMS	200MG	A085175	001
KV PHARM	200MG	A085276	001
LANNETT	200MG	A083743	001
LEDERLE	200MG	A086176	001
LILLY	200MG	A085038	001
PERRIGO	200MG	A085322	001
PHARMAVITE	200MG	A084627	001
PUREPAC PHARM	200MG	A084003	001
SANDOZ	200MG	A084631	001
	200MG	A084914	001
	300MG	A089839	001 Sep 29, 1988
SCHERER LABS	200MG	A085068	001
SUN PHARM INDUSTRIES	100MG	A081029	001 Apr 14, 1989
SUPERPHARM	200MG	A088973	001 Apr 10, 1985
USL PHARMA	200MG	A087837	001 Apr 14, 1982
VALEANT PHARM INTL	200MG	A083393	001
VANGARD	200MG	A087909	001 Jul 13, 1982
VINTAGE PHARMS	200MG	A083963	001
WARNER CHILCOTT	200MG	A083879	001
WATSON LABS	100MG	A085584	001
	200MG	A085140	002
WHITEWORTH TOWN PLSN	200MG	A085444	001
QUINORA			
KEY PHARMS	200MG	A083576	001
SCHERING	300MG	A085222	001
TABLET, EXTENDED RELEASE;ORAL			
QUINIDEX			
WYETH PHARMS INC	300MG	N012796	002

RABEPRAZOLE SODIUMTABLET, DELAYED RELEASE;ORAL  
ACIPHEX  
+ EISAI INC 10MG \*\* N020973 001 May 29, 2002RAMIPRIL

CAPSULE;ORAL

RAMIPRIL			
ACTAVIS ELIZABETH	1.25MG	A077513	001 Jun 18, 2008
	2.5MG	A077513	002 Jun 18, 2008
	5MG	A077513	003 Jun 18, 2008
	10MG	A077513	004 Jun 18, 2008
CIPLA	1.25MG	A077004	001 Aug 07, 2008
	2.5MG	A077004	002 Aug 07, 2008
	5MG	A077004	003 Aug 07, 2008
	10MG	A077004	004 Aug 07, 2008
FOSUN PHARMA	1.25MG	A077514	001 Jun 18, 2008
	2.5MG	A077514	002 Jun 18, 2008
	5MG	A077514	003 Jun 18, 2008
	10MG	A077514	004 Jun 18, 2008
RANBAXY LABS LTD	5MG	A078849	001 Mar 06, 2009
	10MG	A078849	002 Mar 06, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-318(of 375)

\*\* See List Footnote

**RAMIPRIL**

TABLET;ORAL

ALTACE

+ KING PFIZER	1.25MG **	N022021 001	Feb 27, 2007
+	2.5MG **	N022021 002	Feb 27, 2007
+	5MG **	N022021 003	Feb 27, 2007
+	10MG **	N022021 004	Feb 27, 2007

RAMIPRIL

MYLAN PHARMS INC	1.25MG	A090650 001	Jun 30, 2011
	2.5MG	A090650 002	Jun 30, 2011
	5MG	A090650 003	Jun 30, 2011
	10MG	A090650 004	Jun 30, 2011
ZYDUS PHARMS USA INC	1.25MG	A090697 001	Sep 24, 2009
	2.5MG	A090697 002	Sep 24, 2009
	5MG	A090697 003	Sep 24, 2009
	10MG	A090697 004	Sep 24, 2009

**RANITIDINE BISMUTH CITRATE**

TABLET;ORAL

TRITEC

GLAXOSMITHKLINE	400MG	N020559 001	Aug 08, 1996
-----------------	-------	-------------	--------------

**RANITIDINE HYDROCHLORIDE**

CAPSULE;ORAL

RANITIDINE HYDROCHLORIDE

MYLAN	EQ 150MG BASE	A075564 001	Oct 27, 2000
	EQ 300MG BASE	A075564 002	Oct 27, 2000
TEVA	EQ 150MG BASE	A075557 001	Oct 31, 2003
	EQ 300MG BASE	A075557 002	Oct 31, 2003

ZANTAC 150

+ GLAXOSMITHKLINE	EQ 150MG BASE **	N020095 001	Mar 08, 1994
-------------------	------------------	-------------	--------------

ZANTAC 300

+ GLAXOSMITHKLINE	EQ 300MG BASE **	N020095 002	Mar 08, 1994
-------------------	------------------	-------------	--------------

GRANULE, EFFERVESCENT;ORAL

ZANTAC 150

GLAXO GRP LTD	EQ 150MG BASE/PACKET	N020251 002	Mar 31, 1994
---------------	----------------------	-------------	--------------

INJECTABLE;INJECTION

RANITIDINE HYDROCHLORIDE

BEDFORD	EQ 25MG BASE/ML	A074764 001	Nov 19, 2004
ZANTAC IN PLASTIC CONTAINER			

TELIGENT	EQ 1MG BASE/ML	N019593 002	Sep 27, 1991
	EQ 50MG BASE/100ML	N019593 001	Dec 17, 1986

SYRUP;ORAL

RANITIDINE HYDROCHLORIDE

APOTEX INC	EQ 15MG BASE/ML	A077602 001	Sep 17, 2007
RANBAXY	EQ 15MG BASE/ML	A078448 001	Dec 13, 2007
WOCKHARDT	EQ 15MG BASE/ML	A079211 001	May 26, 2009
	EQ 15MG BASE/ML	A079212 001	Feb 23, 2009

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

BOEHRINGER INGELHEIM	EQ 150MG BASE	A074662 001	Aug 29, 1997
	EQ 300MG BASE	A074662 002	Aug 29, 1997
CONTRACT PHARMACAL	EQ 75MG BASE	A075094 001	Jun 21, 1999
MYLAN	EQ 150MG BASE	A074023 001	Aug 22, 1997
	EQ 150MG BASE	A074552 001	Jul 30, 1998
	EQ 300MG BASE	A074023 002	Aug 22, 1997
	EQ 300MG BASE	A074552 002	Jul 30, 1998
RANBAXY	EQ 75MG BASE	A075254 001	Jan 14, 2000
	EQ 150MG BASE	A075000 001	Jan 30, 1998
	EQ 300MG BASE	A075000 002	Jan 30, 1998
SANDOZ	EQ 75MG BASE	A075519 001	Sep 26, 2002
SUN PHARM IND'S LTD	EQ 75MG BASE	A075132 001	Jan 14, 2000
	EQ 150MG BASE	A075439 001	Apr 19, 2000
	EQ 300MG BASE	A075439 002	Apr 19, 2000
WATSON LABS	EQ 75MG BASE	A075212 001	Jan 14, 2000
	EQ 150MG BASE	A074864 001	Oct 20, 1997
	EQ 300MG BASE	A074864 002	Oct 20, 1997
WATSON LABS INC	EQ 150MG BASE	A077426 001	Dec 19, 2005
	EQ 300MG BASE	A077426 002	Dec 19, 2005
WOCKHARDT	EQ 75MG BASE	A078884 001	Jul 31, 2008

**DISCONTINUED DRUG PRODUCT LIST**

6-319(of 375)

\*\* See List Footnote

RANITIDINE HYDROCHLORIDE

TABLET;ORAL

RANITIDINE HYDROCHLORIDE

EQ 150MG BASE	A078653 001 Nov 26, 2007
EQ 150MG BASE	A078701 001 Nov 12, 2009
EQ 300MG BASE	A078701 002 Dec 11, 2009

TABLET, EFFERVESCENT;ORAL

ZANTAC 150

GLAXO GRP LTD EQ 150MG BASE N020251 001 Mar 31, 1994

ZANTAC 25

GLAXO GRP LTD EQ 25MG BASE N020251 003 Apr 01, 2004

ZANTAC 75

+ SANOFI US EQ 75MG BASE \*\* N020745 001 Feb 26, 1998

RANOLAZINE

TABLET, EXTENDED RELEASE;ORAL

RANOLAZINE

LUPIN LTD	500MG	A201046 001 Jul 29, 2013
	1GM	A201046 002 Jul 29, 2013

RAPACURONIUM BROMIDE

INJECTABLE;INJECTION

RAPLON

ORGANON USA INC	100MG/VIAL	N020984 001 Aug 18, 1999
	200MG/VIAL	N020984 002 Aug 18, 1999

RAUWOLFIA SERPENTINA ROOT

TABLET;ORAL

HIWOLFIA

BOWMAN PHARMS	50MG	N009276 003
	50MG	N009276 005
	100MG	N009276 004

HYSERPIN

PHYS PRODS VA 50MG N010581 001

KOGLUCOID

PANRAY	50MG	N009278 001
	100MG	N009278 002

RAUDIXIN

APOTHECON	50MG	N008842 001
	100MG	N008842 002

RAUSERPIN

FERNDALE LABS	50MG	N009926 002
	100MG	N009926 004

RAUVAL

PAL PAK	50MG	N009108 002
	100MG	N009108 004

RAUWOLFIA SERPENTINA

BUNDY	50MG	N009477 001
	100MG	N009477 002

HALSEY	50MG	A080498 001
	100MG	A080498 002

IMPAX LABS	50MG	N009273 001
	100MG	N009273 002

IVAX SUB TEVA PHARMS	50MG	N011521 001
	100MG	N011521 002

PUREPAC PHARM	50MG	A080842 001
	100MG	A080842 002

PVT FORM	50MG	A080583 001
	100MG	A080583 002

SOLVAY	50MG	A080500 001
	100MG	A080500 002

TABLICAPS	50MG	A083867 001
	100MG	A083444 001

VALEANT PHARM INTL	50MG	N009668 001
	100MG	N009668 002

WATSON LABS	50MG	A080907 001
	100MG	A080914 001

WOLFINA

FOREST PHARMS	50MG	N009255 008
	100MG	N009255 006

**DISCONTINUED DRUG PRODUCT LIST**

6-320(of 375)

\*\* See List Footnote

RESCINNAMINE

CAPSULE;ORAL		
CINNASIL		
PANRAY	0.5MG	A084736 001
TABLET;ORAL		
MODERIL		
PFIZER	0.25MG	N010686 003
	0.5MG	N010686 006

RESERPINE

ELIXIR;ORAL		
SERPASIL		
NOVARTIS	0.2MG/4ML	N009115 005
INJECTABLE; INJECTION		
SANDRIL		
LILLY	2.5MG/ML	N010012 001
SERPASIL		
NOVARTIS	2.5MG/ML	N009434 002
TABLET;ORAL		
HISERPIA		
BOWMAN PHARMS	0.1MG	N009631 002
	0.25MG	N009631 004
RAU-SED		
BRISTOL MYERS SQUIBB	0.1MG	N009357 001
	0.25MG	N009357 004
	0.5MG	N009357 006
	1MG	N009357 008
RESERPINE		
BARR	0.25MG	A080721 002
BELL PHARMA	0.1MG	A083058 001
	0.25MG	A083058 002
BUNDY	0.1MG	N009663 001
	0.25MG	N009663 003
CYCLE PHARMS LTD	0.1MG	N009859 001
	0.25MG	N009859 002
ELKINS SINK	0.1MG	A083145 001
	0.25MG	A083145 002
EVERYLIFE	0.1MG	N010441 001
	0.25MG	N010441 002
	0.5MG	N010441 003
	1MG	N010441 004
HALSEY	0.1MG	A080457 002
	0.25MG	A080457 001
	1MG	A080457 003
HIKMA INTL PHARMS	0.1MG	A080975 001
	0.25MG	A080975 002
	1MG	A080975 003
IMPAX LABS	0.1MG	N009627 001
	0.25MG	N009627 002
IVAX SUB TEVA PHARMS	0.1MG	N011185 001
	0.25MG	N011185 002
MARSHALL PHARMA	0.1MG	A080492 001
	0.25MG	A080492 002
MK LABS	0.1MG	A080525 002
	0.25MG	A080525 001
MYLAN	1MG	A084974 001
PHARMAVITE	0.25MG	A084663 001
PUREPAC PHARM	0.1MG	A080753 002
	0.25MG	A080753 001
PVT FORM	0.1MG	A086117 001
	0.25MG	A080582 001
	0.25MG	A085775 001
	1MG	A080582 002
REXALL	0.25MG	A080637 001
+ SANDOZ	0.1MG	N009838 001
+ SOLVAY	0.25MG	N009838 002
TABLICAPS	0.25MG	A080446 001
TEVA	0.1MG	A085207 001
	0.25MG	A089020 001 Mar 07, 1985
		A089019 001 Mar 07, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-321(of 375)

\*\* See List Footnote

RESERPINE

TABLET;ORAL

RESERPINE

VALEANT PHARM INTL	0.1MG	N009667 001
	0.25MG	N009667 002
WATSON LABS	0.1MG	A080679 001
	0.25MG	A080393 001
	0.25MG	A085401 001
	1MG	A080749 001
WHITEWORTH TOWN PLSN	0.1MG	A080723 001
	0.25MG	A080723 002
	1MG	A080723 003
SANDRIL		
LILLY	0.1MG	N009376 004
	0.25MG	N009376 001
SERPALAN		
LANNETT	0.1MG	N010124 001
	0.25MG	N010124 002
SERPANRAY		
PANRAY	0.1MG	N009391 001
	0.25MG	N009391 002
	1MG	N009391 004
SERPASIL		
NOVARTIS	0.1MG	N009115 001
	0.25MG	N009115 003
	1MG	N009115 004
SERPATE		
VALE	0.1MG	N009453 001
	0.25MG	N009453 002
SERPIVITE		
VITARINE	0.25MG	N009645 002

RESERPINE; TRICHLORMETHIAZIDE

TABLET;ORAL

METATENSIN #2

SANOFI AVENTIS US	0.1MG; 2MG	N012972 001
METATENSIN #4		
SANOFI AVENTIS US	0.1MG; 4MG	N012972 002
NAQUIVAL		
SCHERING	0.1MG; 4MG	N012265 003
TRICHLORMETHIAZIDE W/ RESERPINE		
WATSON LABS	0.1MG; 4MG	A085248 001

RIBAVIRIN

CAPSULE;ORAL

REBETOL

MERCK SHARP DOHME	200MG**Indicated for use and comarketed with Interferon ALFA-2B, Recombinant (INTRON A), as Rebetron Combination Therapy**	N020903 001 Jun 03, 1998
-------------------	--	--------------------------

TABLET;ORAL

COPEGUS

ROCHE	400MG	N021511 002 Jun 21, 2005
-------	-------	--------------------------

RIMANTADINE HYDROCHLORIDE

SYRUP;ORAL

FLUMADINE

FOREST LABS	50MG/5ML	N019650 001 Sep 17, 1993
-------------	----------	--------------------------

TABLET;ORAL

RIMANTADINE HYDROCHLORIDE

ACTAVIS ELIZABETH	100MG	A076375 001 Jan 14, 2003
IMPAX LABS INC	100MG	A075916 001 Nov 02, 2001

RIMEXOLONE

SUSPENSION/DROPS;OPHTHALMIC

VEXOL

ALCON	1%	N020474 001 Dec 30, 1994
-------	----	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-322(of 375)

\*\* See List Footnote

RISEDRONATE SODIUM

TABLET;ORAL ACTONEL + APIL	75MG **	N020835 004 Apr 16, 2007
----------------------------------	---------	--------------------------

RISPERIDONE

SOLUTION;ORAL RISPERIDONE SILARX PHARMS INC WOCKHARDT	1MG/ML 1MG/ML	A202386 001 Jan 12, 2015 A078744 001 Oct 08, 2009
TABLET;ORAL RISPERDAL JANSSEN PHARMS RISPERIDONE JUBILANT CADISTA	5MG 0.25MG 0.5MG 1MG 2MG 3MG 4MG	N020272 005 Dec 29, 1993 A078828 001 Mar 23, 2009 A078828 002 Mar 23, 2009 A078828 003 Mar 23, 2009 A078828 004 Mar 23, 2009 A078828 005 Mar 23, 2009 A078828 006 Mar 23, 2009
RATIOPHARM	0.25MG 0.5MG 1MG 2MG 3MG 4MG	A077784 001 Jun 08, 2010 A077784 002 Jun 08, 2010 A077784 003 Jun 08, 2010 A077784 004 Jun 08, 2010 A077784 005 Jun 08, 2010 A077784 006 Jun 08, 2010
SYNTTHON PHARMS	0.25MG 0.5MG 1MG 2MG 3MG 4MG	A078187 001 Oct 22, 2009 A078187 002 Oct 22, 2009 A078187 003 Oct 22, 2009 A078187 004 Oct 22, 2009 A078187 005 Oct 22, 2009 A078187 006 Oct 22, 2009
WATSON LABS	0.25MG 0.5MG 1MG 2MG 3MG 4MG	A077860 001 Dec 05, 2008 A077860 002 Dec 05, 2008 A077860 003 Dec 05, 2008 A077860 004 Dec 05, 2008 A077860 005 Dec 05, 2008 A077860 006 Dec 05, 2008
WEST WARD PHARMS	0.25MG 0.5MG 1MG 2MG 3MG 4MG	A078740 001 May 29, 2009 A078740 002 May 29, 2009 A078740 003 May 29, 2009 A078740 004 May 29, 2009 A078740 005 May 29, 2009 A078740 006 May 29, 2009
TABLET, ORALLY DISINTEGRATING;ORAL RISPERIDONE MYLAN PHARMS INC	0.25MG 0.5MG 1MG 2MG 3MG 4MG	A091537 006 Feb 12, 2013 A091537 001 Mar 30, 2011 A091537 002 Mar 30, 2011 A091537 003 Mar 30, 2011 A091537 004 Mar 30, 2011 A091537 005 Mar 30, 2011

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION RITODRINE HYDROCHLORIDE ABRAXIS PHARM	10MG/ML 15MG/ML	A071188 001 Jul 23, 1987 A071189 001 Jul 23, 1987
HOSPIRA	10MG/ML 15MG/ML	A071618 001 Feb 28, 1991 A071619 001 Feb 28, 1991
RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER HOSPIRA	30MG/100ML	A071438 001 Jan 22, 1991
YUTOPAR ASTRAZENECA	10MG/ML 15MG/ML	N018580 001 N018580 002
TABLET;ORAL YUTOPAR ASTRAZENECA	10MG	N018555 001

**DISCONTINUED DRUG PRODUCT LIST**

6-323(of 375)

\*\* See List Footnote

RITONAVIR

CAPSULE;ORAL

NORVIR

ABBOTT

100MG

N020680 001 Mar 01, 1996

RIVASTIGMINE TARTRATE

SOLUTION;ORAL

EXELON

NOVARTIS

EQ 2MG BASE/ML

N021025 001 Apr 21, 2000

ROCURONIUM BROMIDE

INJECTABLE;INJECTION

ZEMURON

+ ORGANON USA INC

50MG/5ML (10MG/ML) \*\*

N020214 001 Mar 17, 1994

+

10MG/ML (10MG/ML) \*\*

N020214 002 Mar 17, 1994

+

100MG/10ML (10MG/ML) \*\*

N020214 003 Mar 17, 1994

ROFECOXIB

SUSPENSION;ORAL

VIOXX

MERCK

12.5MG/5ML

N021052 001 May 20, 1999

25MG/5ML

N021052 002 May 20, 1999

TABLET;ORAL

VIOXX

MERCK

12.5MG

N021042 001 May 20, 1999

25MG

N021042 002 May 20, 1999

50MG

N021042 003 Feb 25, 2000

ROPINIROLE HYDROCHLORIDE

TABLET;ORAL

ROPINIROLE HYDROCHLORIDE

EPIC PHARMA LLC

EQ 0.25MG BASE

A078230 001 May 20, 2008

EQ 0.5MG BASE

A078230 002 May 20, 2008

EQ 1MG BASE

A078230 003 May 20, 2008

EQ 2MG BASE

A078230 004 May 20, 2008

EQ 3MG BASE

A078230 005 May 20, 2008

EQ 4MG BASE

A078230 006 May 20, 2008

EQ 5MG BASE

A078230 007 May 20, 2008

TABLET, EXTENDED RELEASE;ORAL

REQUIP XL

+ GLAXOSMITHKLINE LLC

EQ 3MG BASE \*\*

N022008 002 Jun 13, 2008

ROPINIROLE HYDROCHLORIDE

MYLAN PHARMS INC

EQ 3MG BASE

A200462 002 Oct 15, 2012

ROPIVACAINE HYDROCHLORIDE

SOLUTION;INJECTION

NAROPIN

FRESENIUS KABI USA

50MG/10ML (5MG/ML)

N020533 013 May 01, 1998

75MG/10ML (7.5MG/ML)

N020533 012 Sep 24, 1996

ROSE BENGAL SODIUM I-131

INJECTABLE;INJECTION

ROBENGATOPE

BRACCO

0.5mCi/VIAL

N016224 001

1mCi/VIAL

N016224 002

2mCi/VIAL

N016224 003

SODIUM ROSE BENGAL I 131

SORIN

0.5mCi/ML

N017318 001

RUFINAMIDE

TABLET;ORAL

BANZEL

+ EISAI INC

100MG \*\*

N021911 001 Nov 14, 2008

SAFFLOWER OIL

INJECTABLE;INJECTION

LIPOSYN 10%

ABBOTT

10% (10GM/100ML)

N018203 001

LIPOSYN 20%

ABBOTT

20% (20GM/100ML)

N018614 001

**DISCONTINUED DRUG PRODUCT LIST**

6-324(of 375)

\*\* See List Footnote

SAFFLOWER OIL; SOYBEAN OIL

INJECTABLE; INJECTION

LIPOSYN II 10%	5%;5% (5GM/100ML)	N018997 001 Aug 27, 1984
HOSPIRA		
LIPOSYN II 20%	10%;10% (10GM/100ML)	N018991 001 Aug 27, 1984

SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

SEREVENT		
GLAXOSMITHKLINE	EQ 0.021MG BASE/INH	N020236 001 Feb 04, 1994

SAQUINAVIR

CAPSULE; ORAL

FORTOVASE		
+ HOFFMANN LA ROCHE	200MG **	N020828 001 Nov 07, 1997

SARALASIN ACETATE

INJECTABLE; INJECTION

SARENIN		
PROCTER AND GAMBLE	EQ 0.6MG BASE/ML	N018009 001

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECOBARBITAL SODIUM		
ANABOLIC	100MG	A084422 001
BARR	100MG	A084225 001
EVERYLIFE	100MG	A085895 001
HALSEY	100MG	A084676 001
IVAX PHARMS	100MG	A085869 001
KV PHARM	100MG	A085285 001
LANNETT	50MG	A085909 001
	100MG	A085903 001
PARKE DAVIS	100MG	A084762 001
PERRIGO	100MG	A084561 001
PUREPAC PHARM	100MG	A085867 001
VALEANT PHARM INTL	100MG	A085477 001
VITARINE	100MG	A085898 001
	100MG	A086273 001
WATSON LABS	100MG	A085792 001
WEST WARD	100MG	A084926 001
WHITEWORTH TOWN PLSN	100MG	A085798 001
WYETH AYERST	100MG	A086390 001

INJECTABLE; INJECTION

SECOBARBITAL SODIUM		
ELKINS SINK	100MG/VIAL	A083281 001
WYETH AYERST	50MG/ML	A083262 001

SECONAL SODIUM		
LILLY	50MG/ML	N007392 002

SUPPOSITORY; RECTAL

SECONAL SODIUM		
LILLY	30MG	A086530 001
	60MG	A086530 002
	120MG	A086530 003
	200MG	A086530 004

SECRETIN

INJECTABLE; INJECTION		
SECRETIN-FERRING		

FERRING	75CU/VIAL	N018290 001
---------	-----------	-------------

SECRETIN SYNTHETIC PORCINE

FOR SOLUTION; INTRAVENOUS		
SECREFLO		

CHIRHOCLIN	16MCG/VIAL	N021136 001 Apr 04, 2002
------------	------------	--------------------------

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL		
SELEGILINE HYDROCHLORIDE		

LANNETT HOLDINGS INC	5MG	A075145 001 Sep 15, 2003
----------------------	-----	--------------------------

**DISCONTINUED DRUG PRODUCT LIST**

6-325(of 375)

\*\* See List Footnote

SELEGILINE HYDROCHLORIDE

TABLET;ORAL

SELEGILINE HYDROCHLORIDE

CHARTWELL MOLECULES	5MG	A074565 001 Aug 02, 1996
	5MG	A074641 001 Aug 02, 1996
G AND W LABS INC	5MG	A074537 001 Aug 02, 1996
	5MG	A074744 001 Jan 27, 1997
	5MG	A074756 001 Nov 25, 1998
SIEGFRIED	5MG	A074672 001 Apr 01, 1997
+ SOMERSET	5MG **	N019334 001 Jun 05, 1989

SELENIUM SULFIDE

LOTION/SHAMPOO;TOPICAL

EXSEL

ALLERGAN HERBERT	2.5%	A083892 001
SELENIUM SULFIDE		
ACTAVIS MID ATLANTIC	2.5%	A084394 001
G AND W LABS INC	2.5%	A086209 001

IVAX PHARMS 2.5%

SELENOMETHIONINE SE-75

INJECTABLE;INJECTION

SELENOMETHIONINE SE 75

GE HEALTHCARE	250uCi/ML	N017257 001
MALLINCKRODT	100uCi/ML	N017098 001
PHARMALUCENCE	500uCi/ML	N017322 001
SETHOTOPE		

BRACCO 85-550uCi/ML

N017047 001

SERMORELIN ACETATE

INJECTABLE;INJECTION

GEREF

+ EMD SERONO	EQ 0.05MG BASE/AMP **	N019863 001 Dec 28, 1990
+ EMD SERONO INC	EQ 0.5MG BASE/VIAL **	N020443 001 Sep 26, 1997
+	EQ 1MG BASE/VIAL **	N020443 002 Sep 26, 1997

SERTRALINE HYDROCHLORIDE

CONCENTRATE;ORAL

SERTRALINE HYDROCHLORIDE

ALLIED PHARMA INC	EQ 20MG BASE/ML	A076934 001 Jun 30, 2006
RANBAXY LABS LTD	EQ 20MG BASE/ML	A078053 001 Feb 05, 2007

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

ACI HEALTHCARE LTD	EQ 25MG BASE	A076881 001 Feb 06, 2007
	EQ 50MG BASE	A076881 002 Feb 06, 2007
	EQ 100MG BASE	A076881 003 Feb 06, 2007
ACTAVIS ELIZABETH	EQ 25MG BASE	A077345 001 Feb 06, 2007
	EQ 50MG BASE	A077345 002 Feb 06, 2007
	EQ 100MG BASE	A077345 003 Feb 06, 2007
ANDA REPOSITORY	EQ 25MG BASE	A077818 001 Feb 06, 2007
	EQ 50MG BASE	A077818 002 Feb 06, 2007
	EQ 100MG BASE	A077818 003 Feb 06, 2007
CIPLA LTD	EQ 25MG BASE	A077162 001 Feb 06, 2007
	EQ 50MG BASE	A077162 002 Feb 06, 2007
	EQ 100MG BASE	A077162 003 Feb 06, 2007
HIKMA PHARMS	EQ 25MG BASE	A077864 001 Aug 10, 2009
	EQ 50MG BASE	A077864 002 Aug 10, 2009
	EQ 100MG BASE	A077864 003 Aug 10, 2009
IVAX SUB TEVA PHARMS	EQ 25MG BASE	A075719 003 Jun 30, 2006
	EQ 50MG BASE	A075719 001 Jun 30, 2006
	EQ 100MG BASE	A075719 002 Jun 30, 2006
MYLAN	EQ 25MG BASE	A076671 001 Feb 06, 2007
	EQ 50MG BASE	A076671 002 Feb 06, 2007
	EQ 100MG BASE	A076671 003 Feb 06, 2007
MYLAN PHARMS INC	EQ 25MG BASE	A076540 001 Mar 20, 2007
	EQ 50MG BASE	A076540 002 Mar 20, 2007
	EQ 100MG BASE	A076540 003 Mar 20, 2007
PLIVA HRVATSKA DOO	EQ 25MG BASE	A077299 001 Feb 06, 2007
	EQ 50MG BASE	A077299 002 Feb 06, 2007
	EQ 100MG BASE	A077299 003 Feb 06, 2007
SANDOZ	EQ 25MG BASE	A077713 001 Feb 06, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-326(of 375)

\*\* See List Footnote

SERTRALINE HYDROCHLORIDE

TABLET;ORAL

SERTRALINE HYDROCHLORIDE

SCIEGEN PHARMS INC	EQ 50MG BASE EQ 100MG BASE EQ 25MG BASE EQ 50MG BASE EQ 100MG BASE EQ 25MG BASE EQ 50MG BASE EQ 100MG BASE	A077713 002 Feb 06, 2007 A077713 003 Feb 06, 2007 A076442 001 Apr 30, 2007 A076442 002 Apr 30, 2007 A076442 003 Apr 30, 2007 A078108 001 Feb 06, 2007 A078108 002 Feb 06, 2007 A078108 003 Feb 06, 2007
SUN PHARM INDs (IN)	EQ 25MG BASE EQ 50MG BASE EQ 100MG BASE	A078108 001 Feb 06, 2007 A078108 002 Feb 06, 2007 A078108 003 Feb 06, 2007
WATSON LABS TEVA	EQ 25MG BASE EQ 50MG BASE EQ 100MG BASE	A077663 001 Feb 06, 2007 A077663 002 Feb 06, 2007 A077663 003 Feb 06, 2007
ZOLOFT		
+ PFIZER	EQ 150MG BASE **	N019839 003 Dec 30, 1991
+	EQ 200MG BASE **	N019839 004 Dec 30, 1991

SEVELAMER HYDROCHLORIDE

CAPSULE;ORAL

RENAGEL

GENZYME 403MG

N020926 001 Oct 30, 1998

SIBUTRAMINE HYDROCHLORIDE

CAPSULE;ORAL

MERIDIA

ABBOTT 5MG  
10MG  
15MGN020632 001 Nov 22, 1997  
N020632 002 Nov 22, 1997  
N020632 003 Nov 22, 1997SILDENAFIL CITRATE

TABLET;ORAL

SILDENAFIL CITRATE

ACTAVIS GRP PTC EQ 20MG BASE

A200149 001 Feb 25, 2013

SILVER SULFADIAZINE

DRESSING;TOPICAL

SILDAFLO

FRANKLIN PHARMS 1%

N019608 001 Nov 30, 1989

SIMETHICONE-CELLULOSE

SUSPENSION;ORAL

SONORX

BRACCO 7.5MG/ML

N020773 001 Oct 29, 1998

SIMVASTATIN

TABLET;ORAL

SIMVASTATIN

FOSUN PHARMA 5MG  
10MG  
20MG  
40MG  
80MGA077766 001 Dec 20, 2006  
A077766 002 Dec 20, 2006  
A077766 003 Dec 20, 2006  
A077766 004 Dec 20, 2006  
A077766 005 Dec 20, 2006MYLAN PHARMS INC 5MG  
10MG  
20MG  
40MG  
80MGA090868 001 Jun 08, 2010  
A090868 002 Jun 08, 2010  
A090868 003 Jun 08, 2010  
A090868 004 Jun 08, 2010  
A090868 005 Jun 08, 2010SUN PHARM INDs LTD 5MG  
10MG  
20MG  
40MG  
80MGA076285 001 Dec 20, 2006  
A076285 002 Dec 20, 2006  
A076285 003 Dec 20, 2006  
A076285 004 Dec 20, 2006  
A076285 005 Jun 23, 2006

TABLET, ORALLY DISINTEGRATING;ORAL

SIMVASTATIN

SYNTTHON PHARMS 10MG  
20MG  
40MG  
80MGN021961 001 Oct 09, 2007  
N021961 002 Oct 09, 2007  
N021961 003 Oct 09, 2007  
N021961 004 Oct 09, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-327(of 375)

\*\* See List Footnote

SIMVASTATIN; SITAGLIPTIN PHOSPHATE

TABLET;ORAL

JUVISYNC

+ MERCK SHARP DOHME	10MG;EQ 50MG BASE **	N202343 004	Sep 18, 2012
+	10MG;EQ 100MG BASE **	N202343 001	Oct 07, 2011
+	20MG;EQ 50MG BASE **	N202343 005	Sep 18, 2012
+	20MG;EQ 100MG BASE **	N202343 002	Oct 07, 2011
+	40MG;EQ 50MG BASE **	N202343 006	Sep 18, 2012
+	40MG;EQ 100MG BASE **	N202343 003	Oct 07, 2011

SIROLIMUS

TABLET;ORAL

RAPAMUNE

+ PF PRISM CV	5MG **	N021110 003	Feb 23, 2004
---------------	--------	-------------	--------------

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION;ORAL

UCEPHAN

B BRAUN	100MG/ML;100MG/ML	N019530 001	Dec 23, 1987
---------	-------------------	-------------	--------------

SODIUM BICARBONATE

INJECTABLE;INJECTION

SODIUM BICARBONATE IN PLASTIC CONTAINER

+ ABBOTT	0.9MEQ/ML **	N019443 001	Jun 03, 1986
+	1MEQ/ML **	N019443 002	Jun 03, 1986

SODIUM BICARBONATE; TARTARIC ACID

GRANULE, EFFERVESCENT;ORAL

BAROS

MALLINCKRODT INC	460MG/GM;420MG/GM	N018509 001	Aug 07, 1985
------------------	-------------------	-------------	--------------

SODIUM CHLORIDE

INJECTABLE;INJECTION

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABRAXIS PHARM	9MG/ML	A088909 001	Feb 07, 1985
---------------	--------	-------------	--------------

SODIUM CHLORIDE

ABBOTT	20GM/100ML	N017013 001
--------	------------	-------------

B BRAUN	20GM/100ML	N017038 001
---------	------------	-------------

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	450MG/100ML	N018184 001
---------	-------------	-------------

MILES	450MG/100ML	N018503 001
-------	-------------	-------------

SODIUM CHLORIDE 0.9%

MEDEFIL INC	18MG/2ML (9MG/ML) 22.5MG/2.5ML (9MG/ML) 27MG/3ML (9MG/ML) 45MG/5ML (9MG/ML)	N202832 002	Jan 06, 2012
		N202832 003	Jan 06, 2012
		N202832 004	Jan 06, 2012
		N202832 005	Jan 06, 2012

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABBOTT	9MG/ML	N019218 001	Jul 13, 1984
--------	--------	-------------	--------------

MEDEFIL INC	9MG/ML (9MG/ML)	N202832 001	Jan 06, 2012
-------------	-----------------	-------------	--------------

MILES	900MG/100ML	N018502 001
-------	-------------	-------------

SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER

+ ABRAXIS PHARM	234MG/ML **	N019329 001	Apr 22, 1987
-----------------	-------------	-------------	--------------

SOLUTION;IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

BAXTER HLTHCARE	450MG/100ML	N017864 001
-----------------	-------------	-------------

	450MG/100ML	N018497 001	Feb 19, 1982
--	-------------	-------------	--------------

HOSPIRA	450MG/100ML	N017670 001
---------	-------------	-------------

	450MG/100ML	N018380 001
--	-------------	-------------

SODIUM CHLORIDE IN PLASTIC CONTAINER

MILES	900MG/100ML	N018247 001
-------	-------------	-------------

SODIUM CHROMATE CR-51

INJECTABLE;INJECTION

CHROMITOPE SODIUM

BRACCO	2mCi/VIAL	N013993 002
--------	-----------	-------------

	200uCi/ML	N013993 001
--	-----------	-------------

SODIUM CHROMATE CR 51

MALLINKRODT NUCLEAR	100uCi/ML	N016708 001
---------------------	-----------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-328(of 375)

\*\* See List Footnote

SODIUM FLUORIDE F-18

INJECTABLE; INTRAVENOUS

FLUORINE F-18

+ GE HEALTHCARE	2mCi/ML **	N017042 001
SODIUM FLUORIDE F 18 NIH NCI DCTD	10-200mCi/ML **	N022494 001 Jan 26, 2011
SODIUM FLUORIDE F-18 UIHC PET IMAGING	10-200mCi/ML	A204462 001 Nov 17, 2015

SODIUM IODIDE I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

CARDINAL HEALTH 418

GE HEALTHCARE

400uCi

100uCi

N018671 003 May 27, 1982  
N017630 001

SOLUTION; ORAL

SODIUM IODIDE I 123

GE HEALTHCARE

2mCi/ML

N017630 002

SODIUM IODIDE I-131

CAPSULE; ORAL

IODOTOPE

BRACCO

1-130mCi

1-150mCi

N010929 001  
N010929 003

SODIUM IODIDE I 131

CIS

50uCi

100uCi

N017316 001  
N017316 002

JUBILANT DRAXIMAGE

2-200mCi

MALLINKRODT NUCLEAR

0.8-100mCi

15-100uCi

N021305 004 Nov 18, 2004  
N016515 002  
N016517 002

SOLUTION; ORAL

HICON

JUBILANT DRAXIMAGE

1-250mCi/0.25ML

1-500mCi/0.5ML

1-1000mCi/ML

N021305 002 Jan 24, 2003  
N021305 003 Jan 24, 2003  
N021305 005 Apr 04, 2006

IODOTOPE

BRACCO

7-106mCi/BOT

N010929 002

SODIUM IODIDE I 131

CIS

50mCi/ML

N017315 001

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

B BRAUN

1.87GM/100ML

N018186 001

BAXTER HLTHCARE

1.87GM/100ML

N016692 001

HOSPIRA

1.87GM/100ML

N018249 001

SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER

B BRAUN

1.87GM/100ML

N020004 001 Apr 21, 1992

SODIUM MONOFLUOROPHOSPHATE

GEL; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS

1.2%

N019518 002 Aug 06, 1986

PASTE; DENTAL

EXTRA-STRENGTH AIM

CHESEBROUGH PONDS

1.2%

N019518 001 Jun 03, 1987

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NIPRIDE

ROCHE

50MG/VIAL

N017546 001

NITROPRESS

ABBOTT

50MG/VIAL

A071555 001 Nov 16, 1987

+ ABBVIE

50MG/VIAL \*\*

N018450 001

HOSPIRA

50MG/VIAL

A070566 001 Jun 09, 1986

SODIUM NITROPRUSSIDE

ABRAXIS PHARM

50MG/VIAL

A070031 001 Jan 17, 1985

+ BAXTER HLTHCARE

50MG/VIAL \*\*

N018581 001 Jul 28, 1982

TEVA PARENTERAL

25MG/ML

A073465 001 Mar 30, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-329(of 375)

\*\* See List Footnote

SODIUM PHOSPHATE P-32

SOLUTION; INJECTION, ORAL PHOSPHOTOPE BRACCO	1-8mCi/VIAL	N0110927 001
SODIUM PHOSPHATE P 32 MALLINCKRODT	0.67mCi/ML 1.5mCi/VIAL	N011777 001 N011777 002

SODIUM PHOSPHATE, DIBASIC ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL VISICOL SALIX PHARMS	0.398GM; 1.102GM	N021097 001 Sep 21, 2000
---	------------------	--------------------------

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL MONOBASIC SODIUM PHOSPHATE AND DIBASIC SODIUM PHOSPHATE NOVEL LABS INC	0.398GM; 1.102GM	A079247 001 Dec 30, 2011
---	------------------	--------------------------

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL KAYEXALATE + CONCORDIA PHARMS INC	453.6GM/BOT	N011287 001
SODIUM POLYSTYRENE SULFONATE CITRUSPHRMA	454GM/BOT	A040909 001 Dec 03, 2008
WOCKHARDT	453.6GM/BOT	A088786 001 Sep 11, 1984
SUSPENSION; ORAL, RECTAL SODIUM POLYSTYRENE SULFONATE MORTON GROVE	15GM/60ML	A088717 001 Sep 11, 1984
ROXANE	15GM/60ML	A088453 001 Nov 17, 1983

SODIUM SUCCINATE

INJECTABLE; INJECTION SODIUM SUCCINATE ELKINS SINK	30%	A080516 001
--	-----	-------------

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION SOTRADECOL + ELKINS SINK	1% **	N005970 004
	3% **	N005970 005

SODIUM THIOSULFATE

INJECTABLE; INJECTION SODIUM THIOSULFATE US ARMY	250MG/ML	N020166 001 Feb 14, 1992
--	----------	--------------------------

SOMATREM

INJECTABLE; INJECTION PROTROPIN GENENTECH	5MG/VIAL 10MG/VIAL	N019107 001 Oct 17, 1985 N019107 002 Oct 24, 1989
---	-----------------------	--

SOMATROPIN

INJECTABLE; INJECTION ASELLACRIN 10 SERONO	10 IU/VIAL	N017726 001
ASELLACRIN 2 SERONO	2 IU/VIAL	N017726 002 Jul 21, 1983
CRESCORMON GENENTECH	4 IU/VIAL	N017992 001

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION ACCRETROPIN EMERGENT	5MG/ML (5MG/ML)	N021538 001 Jan 23, 2008
BIO-TROPIN FERRING	4.8MG/VIAL	N019774 001 May 25, 1995
HUMATROPE LILLY	2MG/VIAL	N019640 001 Jun 23, 1987
NORDITROPIN NOVO NORDISK INC	4MG/VIAL 5MG/1.5ML 8MG/VIAL 10MG/1.5ML	N019721 001 May 08, 1995 N021148 001 Jun 20, 2000 N019721 002 May 08, 1995 N021148 002 Jun 20, 2000

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-330(of 375)

\*\* See List Footnote

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

NORDITROPIN

15MG/1.5ML

N021148 003 Jun 20, 2000

NORDITROPIN NORDIFLEX

NOVO NORDISK INC

5MG/1.5ML

N021148 004 Oct 01, 2004

10MG/1.5ML

N021148 005 Oct 01, 2004

15MG/1.5ML

N021148 006 Oct 01, 2004

30MG/3ML

N021148 007 Mar 10, 2009

NUTROPIN

GENENTECH

5MG/VIAL

N020168 001 Nov 17, 1993

10MG/VIAL

N020168 002 Nov 17, 1993

NUTROPIN AQ

GENENTECH

10MG/2ML (5MG/ML)

N020522 001 Dec 29, 1995

NUTROPIN DEPOT

GENENTECH

13.5MG/VIAL

N021075 001 Dec 22, 1999

18MG/VIAL

N021075 002 Dec 22, 1999

22.5MG/VIAL

N021075 003 Dec 22, 1999

SAIZEN

EMD SERONO

4MG/VIAL

N019764 005 Jan 16, 2007

6MG/VIAL

N019764 001 Oct 08, 1996

SEROSTIM

EMD SERONO

8.8MG/VIAL

N020604 004 Sep 06, 2001

ZORBTIVE

EMD SERONO

4MG/VIAL

N021597 001 Dec 01, 2003

5MG/VIAL

N021597 002 Dec 01, 2003

6MG/VIAL

N021597 003 Dec 01, 2003

INJECTABLE; SUBCUTANEOUS

SEROSTIM LQ

EMD SERONO

6MG/0.5ML (6MG/0.5ML)

N020604 005 Feb 11, 2005

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE

3GM/100ML

N018512 001 May 27, 1982

SOTALOL HYDROCHLORIDE

TABLET; ORAL

BETAPACE

COVIS PHARMA BV

320MG

N019865 004 Oct 30, 1992

BETAPACE AF

COVIS PHARMA BV

40MG

N021151 006 Apr 02, 2003

60MG

N021151 007 Apr 02, 2003

100MG

N021151 005 Mar 14, 2003

SOTALOL HYDROCHLORIDE

IMPAK PHARMS

80MG

A075663 001 Nov 07, 2000

120MG

A075663 002 Nov 07, 2000

160MG

A075663 003 Nov 07, 2000

240MG

A075663 004 Nov 07, 2000

MYLAN

80MG

A075237 001 May 01, 2000

80MG

A075725 001 Dec 19, 2000

120MG

A075725 002 May 01, 2000

120MG

A075237 003 May 01, 2000

160MG

A075725 003 Dec 19, 2000

160MG

A075237 004 May 01, 2000

240MG

A075725 004 Dec 19, 2000

240MG

A075725 004 Dec 19, 2000

SUN PHARM INDUSTRIES

80MG

A075515 001 Oct 15, 2001

80MG

A076576 001 Apr 08, 2004

120MG

A075515 004 Oct 15, 2001

120MG

A076576 002 Apr 08, 2004

160MG

A075515 002 Oct 15, 2001

160MG

A076576 003 Apr 08, 2004

240MG

A075515 003 Oct 15, 2001

TEVA

80MG

A076883 001 Jul 26, 2004

120MG

A076883 002 Jul 26, 2004

160MG

A076883 003 Jul 26, 2004

WATSON LABS

80MG

A075238 001 Jul 13, 2000

120MG

A075238 002 Jul 13, 2000

160MG

A075238 003 Jul 13, 2000

**DISCONTINUED DRUG PRODUCT LIST**

6-331(of 375)

\*\* See List Footnote

SOTALOL HYDROCHLORIDE

TABLET;ORAL

SOTALOL HYDROCHLORIDE

240MG

A075238 004 Jul 13, 2000

SOYBEAN OIL

INJECTABLE;INJECTION

LIPOSYN III 10%

HOSPIRA

10%

N018969 001 Sep 24, 1984

LIPOSYN III 20%

HOSPIRA

20%

N018970 001 Sep 25, 1984

LIPOSYN III 30%

HOSPIRA

30%

N020181 001 Jan 13, 1998

SOYACAL 10%

ALPHA THERA

10%

N018465 001 Jun 29, 1983

SOYACAL 20%

ALPHA THERA

20%

N018786 001 Jun 29, 1983

TRAVAMULSION 10%

BAXTER HLTHCARE

10%

N018660 001 Feb 26, 1982

TRAVAMULSION 20%

BAXTER HLTHCARE

20%

N018758 001 Feb 15, 1983

SPARFLOXACIN

TABLET;ORAL

ZAGAM

MYLAN

200MG

N020677 001 Dec 19, 1996

SPECTINOMYCIN HYDROCHLORIDE

INJECTABLE;INJECTION

TROBICIN

PFIZER

EQ 2GM BASE/VIAL

N050347 001

EQ 4GM BASE/VIAL

N050347 002

SPIRAPRIL HYDROCHLORIDE

TABLET;ORAL

RENORMAX

SCHERING

3MG

N020240 001 Dec 29, 1994

6MG

N020240 002 Dec 29, 1994

12MG

N020240 003 Dec 29, 1994

24MG

N020240 004 Dec 29, 1994

SPIRONOLACTONE

TABLET;ORAL

SPIRONOLACTONE

ASCOT

25MG

A087687 001 Oct 20, 1982

IVAX PHARMS

25MG

A087108 001

LEDERLE

25MG

A087634 001

MUTUAL PHARM

25MG

A087265 001

MYLAN

25MG

A087086 001

PUREPAC PHARM

25MG

A087998 001 Oct 14, 1983

25MG

A088053 001 Aug 25, 1983

SUPERPHARM

25MG

A089364 001 Nov 07, 1986

UPSHER SMITH

25MG

A087554 001

VANGARD

25MG

A087648 001 Feb 01, 1982

WARNER CHILCOTT

25MG

A087952 001 Nov 18, 1982

WATSON LABS

25MG

A086898 002 Mar 02, 1982

25MG

A087078 001

STANOZOLOL

TABLET;ORAL

WINSTROL

LUNDBECK INC

2MG

N012885 001 May 14, 1984

STAVUDINE

CAPSULE;ORAL

STAVUDINE

MYLAN LABS LTD

30MG

A078775 001 Jan 05, 2009

40MG

A078775 002 Jan 05, 2009

ZERIT

BRISTOL MYERS SQUIBB

5MG

N020412 001 Jun 24, 1994

**DISCONTINUED DRUG PRODUCT LIST**

6-332(of 375)

\*\* See List Footnote

STAVUDINE

CAPSULE, EXTENDED RELEASE;ORAL  
ZERIT XR

BRISTOL MYERS SQUIBB 37.5MG  
50MG  
75MG  
100MG

N021453 001 Dec 31, 2002  
N021453 002 Dec 31, 2002  
N021453 003 Dec 31, 2002  
N021453 004 Dec 31, 2002

FOR SOLUTION;ORAL  
STAVUDINE

AUROBINDO PHARMA 1MG/ML

A077774 001 Dec 29, 2008

STERILE WATER FOR INJECTION

LIQUID;N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

ABRAXIS PHARM 100%  
100%

A089099 001 Dec 29, 1987  
A089100 001 Dec 29, 1987

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

B BRAUN 100%

N019077 001 Mar 02, 1984

STERILE WATER FOR IRRIGATION

LIQUID;IRRIGATION

STERILE WATER IN PLASTIC CONTAINER  
MILES 100%

N018246 001

STREPTOMYCIN SULFATE

INJECTABLE;INJECTION

STREPTOMYCIN SULFATE

COPANOS EQ 500MG BASE/ML  
LILLY EQ 1GM BASE/VIAL  
EQ 1GM BASE/2ML  
EQ 5GM BASE/VIAL  
PFIZER EQ 1GM BASE/VIAL  
EQ 1GM BASE/2.5ML  
EQ 5GM BASE/VIAL

A060684 001  
A060107 001  
A060404 001  
A060107 002  
A060076 001  
A060111 001  
A060076 002

SUCCINYLCHOLINE CHLORIDE

INJECTABLE;INJECTION

ANECTINE

SANDOZ INC 50MG/ML  
500MG/VIAL  
1GM/VIAL

N008453 003  
N008453 001  
N008453 004

QUELICIN PRESERVATIVE FREE

HOSPIRA 50MG/ML  
100MG/ML

N008845 002  
N008845 004

SUCCINYLCHOLINE CHLORIDE

INTL MEDICATION 100MG/VIAL  
ORGANON USA INC 20MG/ML

A085400 001 Feb 04, 1982  
A080997 001

SUCOSTRIN

APOTHECON 20MG/ML  
100MG/ML

N008847 001  
N008847 003

SUFENTANIL CITRATE

INJECTABLE;INJECTION

SUFENTANIL CITRATE

WATSON LABS EQ 0.05MG BASE/ML

A074406 001 Dec 15, 1995

SULFACETAMIDE SODIUM

OINTMENT;OPHTHALMIC

BLEPH-10

ALLERGAN 10%

A084015 001

CETAMIDE

ALCON 10%

A080021 001

SODIUM SULAMYD

+ SCHERING 10% \*\*

N005963 002

SULFAIR 10

PHARMAFAIR 10%

A088000 001 Dec 22, 1982

SOLUTION/DROPS;OPHTHALMIC

BLEPH-30

ALLERGAN 30%

A080028 002

IISOPTO CETAMIDE

ALCON 15%

A080020 002

**DISCONTINUED DRUG PRODUCT LIST**

6-333(of 375)

\*\* See List Footnote

SULFACETAMIDE SODIUM

SOLUTION/DROPS;OPHTHALMIC		
OCUSULF-10		
MIZA PHARMS USA	10%	A080660 001
OCUSULF-30		
MIZA PHARMS USA	30%	A080660 002
SODIUM SULAMYD		
+ SCHERING	10% **	N005963 001
+	30% **	N005963 003
SODIUM SULFACETAMIDE		
AKORN	10%	A083021 001
	15%	A083021 002
	30%	A083021 003
SOLA BARNES HIND	10%	A084143 001
	10%	A084145 001
	30%	A084146 001
	30%	A084147 001
SULF-10		
NOVARTIS	10%	A080025 001
SULF-15		
NOVARTIS	15%	A089047 001 Oct 31, 1995
SULFACEL-15		
OPTOPICS	15%	A080024 001
SULFACETAMIDE SODIUM		
AKORN	30%	A040216 001 May 25, 1999
ALCON PHARMS LTD	30%	A089068 001 May 05, 1987
PHARMAFAIR	10%	A088947 001 May 17, 1985
SULFAIR 10		
PHARMAFAIR	10%	A087949 001 Dec 13, 1982
SULFAIR FORTE		
PHARMAFAIR	30%	A088385 001 Oct 13, 1983
SULFAIR-15		
PHARMAFAIR	15%	A088186 001 May 25, 1983
SULTEN-10		
BAUSCH AND LOMB	10%	A087818 001 Feb 03, 1983

SULFACYTINE

TABLET;ORAL		
RENOQUID		
GLENWOOD	250MG	N017569 001

SULFADIAZINE

TABLET;ORAL		
SULFADIAZINE		
ABBVIE	300MG	N004125 005
EVERYLIFE	500MG	A080088 001
IMPAK LABS	500MG	A080081 001
LANNETT	500MG	A080084 001
LEDERLE	500MG	N004054 001
LILLY	500MG	N004122 002

SULFADIAZINE SODIUM

INJECTABLE; INJECTION		
SULFADIAZINE SODIUM		
LEDERLE	250MG/ML	N004054 002

SULFADIAZINE; SULFAMERAZINE

SUSPENSION;ORAL		
SULFONAMIDES DUPLEX		
LILLY	250MG/5ML;250MG/5ML	N006317 007

SULFAMETER

TABLET;ORAL		
SULLA		
BAYER HLTHCARE	500MG	N016000 002

**DISCONTINUED DRUG PRODUCT LIST**

6-334(of 375)

\*\* See List Footnote

SULFAMETHIZOLE

TABLET;ORAL		
MICROSUL		
FOREST PHARMS	1GM	A086012 001
PROKLAR		
FOREST PHARMS	500MG	A080273 001
THIOSULFIL		
WYETH AYERST	250MG	N008565 001
	500MG	N008565 004

SULFAMETHOXAZOLE

SUSPENSION;ORAL		
GANTANOL		
ROCHE	500MG/5ML	N013664 002
TABLET;ORAL		
GANTANOL		
ROCHE	500MG	N012715 002
GANTANOL-DS		
ROCHE	1GM	N012715 003
SULFAMETHOXAZOLE		
ASCOT	500MG	A087662 001 Oct 20, 1982
AUROLIFE PHARMA LLC	500MG	A085844 001
BARR	500MG	A087189 001 Jul 25, 1983
HEATHER	500MG	A086163 001
WATSON LABS	500MG	A085053 001
	1GM	A086000 001
UROBAK		
SHIONOGI	500MG	A087307 001

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE;INJECTION		
BACTRIM		
+ SUN PHARM INDNS INC	80MG/ML;16MG/ML **	N018374 001
SEPTRA		
MONARCH PHARMS	80MG/ML;16MG/ML	N018452 001
SULFAMETHOXAZOLE AND TRIMETHOPRIM		
ABRAXIS PHARM	80MG/ML;16MG/ML	A070223 001 Dec 29, 1987
BEDFORD	80MG/ML;16MG/ML	A072383 001 Apr 29, 1992
HOSPIRA	80MG/ML;16MG/ML	A073199 001 Sep 11, 1992
WATSON LABS	80MG/ML;16MG/ML	A071556 001 Dec 29, 1987
WEST-WARD PHARMS INT	80MG/ML;16MG/ML	A070627 001 Dec 29, 1987
	80MG/ML;16MG/ML	A070628 001 Dec 29, 1987

## SUSPENSION;ORAL

BACTRIM		
SUN PHARM INDUSTRIES	200MG/5ML;40MG/5ML **	N017560 001
BACTRIM PEDIATRIC		
SUN PHARM INDUSTRIES	200MG/5ML;40MG/5ML **	N017560 002
SEPTRA		
MONARCH PHARMS	200MG/5ML;40MG/5ML **	N017598 001
SEPTRA GRAPE		
MONARCH PHARMS	200MG/5ML;40MG/5ML **	N017598 002 Feb 12, 1986
SULFAMETHOXAZOLE AND TRIMETHOPRIM		
ANI PHARMS INC	200MG/5ML;40MG/5ML	A070028 001 Jun 02, 1987
	200MG/5ML;40MG/5ML **	A077612 001 Nov 13, 2006
TEVA	200MG/5ML;40MG/5ML	N018812 001 Jan 28, 1983
	200MG/5ML;40MG/5ML	N018812 002 Jun 10, 1983
SULFATRIM		
STI PHARMA LLC	200MG/5ML;40MG/5ML	N018615 002 Jan 07, 1983
SULMEPRIM		
USL PHARMA	200MG/5ML;40MG/5ML	A070063 001 Aug 01, 1986
SULMEPRIM PEDIATRIC		
USL PHARMA	200MG/5ML;40MG/5ML	A070064 001 Aug 01, 1986
TRIMETH/SULFA		
ALPHARMA US PHARMS	200MG/5ML;40MG/5ML	A072289 001 May 23, 1988
	200MG/5ML;40MG/5ML	A072398 001 May 23, 1988
NASKA	200MG/5ML;40MG/5ML	A072399 001 May 23, 1988
TABLET;ORAL		
COTRIM		
TEVA	400MG;80MG	A070034 001 May 16, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-335(of 375)

\*\* See List Footnote

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET;ORAL			
COTRIM D.S.			
TEVA	800MG;160MG	A070048	001 Mar 18, 1985
SULFAMETHOPRIM			
NOVEL LABS INC	400MG;80MG	A070022	001 Feb 15, 1985
SULFAMETHOPRIM-DS			
NOVEL LABS INC	800MG;160MG	A070032	001 Feb 15, 1985
SULFAMETHOXAZOLE AND TRIMETHOPRIM			
HEATHER	400MG;80MG	N018946	001 Aug 10, 1984
	800MG;160MG	N018946	002 Aug 10, 1984
INTERPHARM	400MG;80MG	A071299	001 Oct 27, 1987
	800MG;160MG	A071300	001 Oct 27, 1987
MARTEC USA LLC	400MG;80MG	A072408	001 Dec 07, 1988
MUTUAL PHARM	400MG;80MG	A070006	001 Nov 14, 1984
PLIVA	400MG;80MG	A070215	001 Sep 10, 1985
	800MG;160MG	A070216	001 Sep 10, 1985
ROXANE	400MG;80MG	A072768	001 Aug 30, 1991
SANDOZ	400MG;80MG	A070889	001 Nov 13, 1986
	400MG;80MG	N018598	003 May 19, 1982
	800MG;160MG	A070890	001 Nov 13, 1986
TEVA	400MG;80MG	N018242	001
	800MG;160MG	N018242	002
USL PHARMA	400MG;80MG	A070203	001 Nov 08, 1985
	800MG;160MG	A070204	001 Nov 08, 1985
WATSON LABS	400MG;80MG	A070002	001 Nov 07, 1984
	400MG;80MG	N018852	001 May 09, 1983
	800MG;160MG	A070000	001 Nov 07, 1984
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH			
MARTEC USA LLC	800MG;160MG	A072417	001 Dec 07, 1988
MUTUAL PHARM	800MG;160MG	A070007	001 Nov 14, 1984
ROXANE	800MG;160MG	A072769	001 Aug 30, 1991
SANDOZ	800MG;160MG	N018598	004 May 19, 1982
WATSON LABS	800MG;160MG	N018854	001 May 09, 1983
SULFATRIM-DS			
SUPERPHARM	800MG;160MG	A070066	001 Jun 24, 1985
SULFATRIM-SS			
SUPERPHARM	400MG;80MG	A070065	002 Jun 24, 1985
UROPLUS DS			
SHIONOGI	800MG;160MG	A071816	001 Sep 28, 1987
UROPLUS SS			
SHIONOGI	400MG;80MG	A071815	001 Sep 28, 1987

SULFANILAMIDE

CREAM;VAGINAL			
SULFANILAMIDE			
G AND W LABS INC	15%	A088718	001 Sep 19, 1985
SUPPOSITORY;VAGINAL			
AVC			
MYLAN SPECIALITY LP	1.05GM	N006530	004 Jan 27, 1987

SULFAPHENAZOLE

SUSPENSION;ORAL			
SULFABID			
PHARM RES ASSOC	500MG/5ML	N013093	001
TABLET;ORAL			
SULFABID			
PURDUE FREDERICK	500MG	N013092	002

SULFAPYRIDINE

TABLET;ORAL			
SULFAPYRIDINE			
LILLY	500MG	N000159	001

SULFASALAZINE

SUSPENSION;ORAL			
AZULFIDINE			
PHARMACIA AND UPJOHN	250MG/5ML	N018605	001
TABLET;ORAL			
S.A.S.-500			
SOLVAY	500MG	A083450	001

**DISCONTINUED DRUG PRODUCT LIST**

6-336(of 375)

\*\* See List Footnote

SULFASALAZINE

TABLET;ORAL

SULFASALAZINE

HERITAGE PHARMS INC	500MG	A080197 001
SANDOZ	500MG	A086184 001
SUN PHARM INDUSTRIES	500MG	A089590 001 Oct 19, 1987
SUPERPHARM	500MG	A089339 001 Oct 26, 1987
WATSON LABS	500MG	A084964 001
	500MG	A087197 001

TABLET, DELAYED RELEASE;ORAL

SULFASALAZINE

WATSON LABS	500MG	A088052 001 May 24, 1983
-------------	-------	--------------------------

SULFINPYRAZONE

CAPSULE;ORAL

ANTURANE

+ NOVARTIS	200MG **	N011556 004
SULFINPYRAZONE		
BARR	200MG	A087666 001 Sep 17, 1982
IVAX PHARMS	200MG	A087770 001 Nov 19, 1982
PAR PHARM	200MG	A088934 001 Sep 06, 1985
VANGARD	200MG	A088666 001 Feb 17, 1984

TABLET;ORAL

ANTURANE

NOVARTIS	100MG **	N011556 003
SULFINPYRAZONE		
BARR	100MG	A087665 001 Sep 17, 1982
IVAX PHARMS	100MG	A087769 001 Jun 01, 1982
PAR PHARM	100MG	A088933 001 Sep 06, 1985
WATSON LABS	100MG	A087667 001 May 26, 1982

SULFISOXAZOLE

TABLET;ORAL

GANTRISIN

ROCHE	500MG	N006525 001
-------	-------	-------------

SOSOL

MK LABS	500MG	A080036 001
---------	-------	-------------

SOXAZOLE

ALRA	500MG	A080366 001
------	-------	-------------

SULFALAR

PARKE DAVIS	500MG	A084955 001
-------------	-------	-------------

SULFISOXAZOLE

ANI PHARMS INC	500MG	A080142 001
----------------	-------	-------------

AUROLIFE PHARMA LLC	500MG	A085628 001
BARR	500MG	A084031 001
HEATHER	500MG	A080189 001
IMPAX LABS	500MG	A080109 001
LANNETT	500MG	A080085 001
LEDERLE	500MG	A087649 001
PHARMERAL	500MG	A084385 001
PUREPAC PHARM	500MG	A080087 001
ROXANE	500MG	A080082 001
VALEANT PHARM INTL	500MG	A080268 002
VITARINE	500MG	A087332 001
WATSON LABS	500MG	A085534 001
WEST WARD	500MG	A080379 001

SULSOXIN

SOLVAY	500MG	A080040 001
--------	-------	-------------

SULFISOXAZOLE ACETYL

EMULSION;ORAL

LIPO GANTRISIN		
ROCHE	EQ 1GM BASE/5ML	N009182 009

SUSPENSION;ORAL

GANTRISIN PEDIATRIC		
ROCHE	EQ 500MG BASE/5ML	N009182 004

SYRUP;ORAL

GANTRISIN		
ROCHE	EQ 500MG BASE/5ML	N009182 002

**DISCONTINUED DRUG PRODUCT LIST**

6-337(of 375)

\*\* See List Footnote

**SULFISOXAZOLE DIOLAMINE**

INJECTABLE; INJECTION

GANTRISIN

ROCHE

EQ 400MG BASE/ML

N006917 001

OINTMENT; OPHTHALMIC

GANTRISIN

ROCHE

EQ 4% BASE

N008414 002

SOLUTION/DROPS; OPHTHALMIC

GANTRISIN

ROCHE

EQ 4% BASE

N007757 002

SULFISOXAZOLE DIOLAMINE

SOLA BARNES HIND

EQ 4% BASE

A084148 001

**SULFOXONE SODIUM**

TABLET, DELAYED RELEASE; ORAL

DIASONE SODIUM

ABBVIE

165MG

N006044 003

**SULFUR**

POWDER; TOPICAL

BENSULFOID

POYTHRESS

33.32%

N002918 001

**SULINDAC**

TABLET; ORAL

CLINORIL

+ MERCK

150MG \*\*

N017911 001

+ +

200MG \*\*

N017911 002

SULINDAC

ANI PHARMS INC

150MG

A072972 001 Feb 28, 1992

200MG

A072973 001 Feb 28, 1992

SANDOZ

150MG

A072712 001 Aug 30, 1991

200MG

A072713 001 Aug 30, 1991

**SUMATRIPTAN**

SPRAY; NASAL

IMITREX

GLAXOSMITHKLINE

10MG/SPRAY

N020626 002 Aug 26, 1997

**SUMATRIPTAN SUCCINATE**

INJECTABLE; SUBCUTANEOUS

ALSUMA

MERIDIAN MEDCL

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

N022377 001 Jun 29, 2010

SUMATRIPTAN SUCCINATE

FRESENIUS KABI USA

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A079240 002 Sep 18, 2009

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A079240 001 Sep 18, 2009

SANDOZ INC

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A078067 002 Feb 06, 2009

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A078067 001 Feb 06, 2009

TEVA PARENTERAL

EQ 4MG BASE/0.5ML (EQ 8MG BASE/ML)

A078318 001 Feb 06, 2009

EQ 6MG BASE/0.5ML (EQ 12MG BASE/ML)

A078318 002 Feb 06, 2009

SYSTEM; IONTOPHORESIS

ZECURITY

+ TEVA BRANDED PHARM

EQ 6.5MG BASE/4HR

N202278 001 Jan 17, 2013

TABLET; ORAL

SUMATRIPTAN SUCCINATE

HIKMA PHARMS

EQ 25MG BASE

A078298 001 May 21, 2013

EQ 50MG BASE

A078298 002 May 21, 2013

EQ 100MG BASE

A078298 003 May 21, 2013

MYLAN

EQ 25MG BASE

A077163 001 Nov 02, 2009

EQ 50MG BASE

A077163 002 Nov 02, 2009

EQ 100MG BASE

A077163 003 Nov 02, 2009

ROXANE

EQ 25MG BASE

A078241 001 Aug 10, 2009

EQ 50MG BASE

A078241 002 Aug 10, 2009

EQ 100MG BASE

A078241 003 Aug 10, 2009

SANDOZ

EQ 25MG BASE

A076976 001 Aug 10, 2009

EQ 50MG BASE

A076976 002 Aug 10, 2009

EQ 100MG BASE

A076976 003 Aug 10, 2009

TEVA

EQ 25MG BASE

A076840 001 Feb 09, 2009

EQ 50MG BASE

A076840 002 Feb 09, 2009

EQ 100MG BASE

A076840 003 Feb 09, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-338(of 375)

\*\* See List Footnote

SUPROFEN

SOLUTION/DROPS;OPHTHALMIC

PROFENAL

ALCON

1%

N019387 001 Dec 23, 1988

SUTILAINS

OINTMENT;TOPICAL

TRAVASE

+ ABBOTT

82,000 UNITS/GM \*\*

N012828 001

TACRINE HYDROCHLORIDE

CAPSULE;ORAL

COGNEX

SHIONOGI INC

EQ 10MG BASE

N020070 001 Sep 09, 1993

EQ 20MG BASE

N020070 002 Sep 09, 1993

EQ 30MG BASE

N020070 003 Sep 09, 1993

EQ 40MG BASE

N020070 004 Sep 09, 1993

TACROLIMUS

CAPSULE;ORAL

TACROLIMUS

WATSON LABS

EQ 5MG BASE

A090402 001 Jul 01, 2010

TALBUTAL

TABLET;ORAL

LOTUSATE

SANOFI AVENTIS US

120MG

N009410 005

TAMOXIFEN CITRATE

TABLET;ORAL

NOLVADEX

+ ASTRAZENECA

EQ 10MG BASE \*\*

N017970 001

+ ASTRAZENECA

EQ 20MG BASE \*\*

N017970 002 Mar 21, 1994

TAMOXIFEN CITRATE

ACTAVIS LABS FL INC

EQ 10MG BASE

A076179 001 Feb 20, 2003

EQ 20MG BASE

A076179 002 Feb 20, 2003

AEGIS PHARMS

EQ 10MG BASE

A076398 001 Mar 31, 2003

EQ 20MG BASE

A076398 002 Mar 31, 2003

IVAX SUB TEVA PHARMS

EQ 10MG BASE

A075740 001 Feb 20, 2003

EQ 20MG BASE

A075740 002 Feb 20, 2003

PHARMACHEMIE

EQ 10MG BASE

A074539 001 Mar 31, 2003

ROXANE

EQ 10MG BASE

A076027 001 Feb 20, 2003

EQ 20MG BASE

A076027 002 Feb 20, 2003

TEVA

EQ 10MG BASE

A074504 001 Apr 28, 2003

EQ 20MG BASE

A074504 002 Apr 28, 2003

TECHNETIUM TC-99M ALBUMIN AGGREGATED

INJECTABLE;INJECTION

TC 99M-LUNGAGGREGATE

GE HEALTHCARE

5mCi/ML

N017848 001

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE;INJECTION

A-N STANNOUS AGGREGATED ALBUMIN

SYNCOR PHARMS N/A

N017916 001

AN-MAA

PHARMALUCENCE N/A

N017792 001

LUNGAGGREGATE REAGENT

GE HEALTHCARE N/A

N017838 001

MACROTEC

BRACCO N/A

N017833 001

TECHNESCAN MAA

MALLINCKRODT N/A

N017842 001

TECHNETIUM TC 99M MAA

GE HEALTHCARE N/A

N017773 001

TECHNETIUM TC-99M ALBUMIN COLLOID KIT

INJECTABLE;INJECTION

MICROLITE

PHARMALUCENCE N/A

N018263 001 Mar 25, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-339(of 375)

\*\* See List Footnote

TECHNETIUM TC-99M ALBUMIN KIT

INJECTABLE; INJECTION  
 TECHNETIUM TC 99M HSA  
 GE HEALTHCARE N/A N017775 001

TECHNETIUM TC-99M ALBUMIN MICROSPHERES KIT

INJECTABLE; INJECTION  
 INSTANT MICROSPHERES  
 3M N/A N017832 001

TECHNETIUM TC-99M APCITIDE

INJECTABLE; INJECTION  
 ACUTECT  
 CIS BIO INTL SA N/A N020887 001 Sep 14, 1998

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE; INJECTION  
 NEO TECT KIT  
 CIS BIO INTL SA N/A N021012 001 Aug 03, 1999

TECHNETIUM TC-99M ETIDRONATE KIT

INJECTABLE; INJECTION  
 CINTICHEM TECHNETIUM 99M HEDSPA  
 GE HEALTHCARE N/A N017653 001  
 MPI STANNOUS DIPHOSPHONATE  
 GE HEALTHCARE N/A N017667 001  
 OSTEOSCAN  
 MALLINCKRODT N/A N017454 001  
 TECHNETIUM TC 99M DIPHOSPHONATE-TIN KIT  
 GE HEALTHCARE N/A N017562 001

TECHNETIUM TC-99M FERPENTETATE KIT

INJECTABLE; INJECTION  
 RENOTEC  
 BRACCO N/A N017045 001

TECHNETIUM TC-99M GLUCEPTATE KIT

INJECTABLE; INJECTION  
 GLUCOSCAN  
 BRISTOL MYERS SQUIBB N/A N017907 001  
 TECHNESCAN GLUCEPTATE  
 DRAXIMAGE N/A N018272 001 Jan 27, 1982

TECHNETIUM TC-99M LIDOGENIN KIT

INJECTABLE; INJECTION  
 TECHNESCAN HIDA  
 DRAXIMAGE N/A N018489 001 Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE

INJECTABLE; INJECTION  
 DRAXIMAGE MDP-10  
 JUBILANT DRAXIMAGE N/A N018035 001

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION  
 AMERSCAN MDP KIT  
 GE HEALTHCARE N/A N018335 001 Aug 05, 1982  
 OSTEOLITE  
 PHARMALUCENCE N/A N017972 001  
 TECHNETIUM TC 99M MPI MDP  
 GE HEALTHCARE N/A N018141 001  
 N/A N018141 002 Jun 12, 1989

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION  
 AN-DTPA  
 JUBILANT DRAXIMAGE N/A N017714 001  
 MPI DTPA KIT - CHELATE  
 GE HEALTHCARE N/A N017255 001  
 TECHNETIUM TC-99M PENTETATE KIT  
 GE HEALTHCARE N/A N017264 002

**DISCONTINUED DRUG PRODUCT LIST**

6-340(of 375)

\*\* See List Footnote

TECHNETIUM TC-99M POLYPHOSPHATE KIT

INJECTABLE; INJECTION  
 SODIUM POLYPHOSPHATE-TIN KIT  
 GE HEALTHCARE N/A N017664 001

TECHNETIUM TC-99M PYRO/TRIMETA PHOSPHATES KIT

INJECTABLE; INJECTION  
 PYROLITE  
 PHARMALUCENCE N/A N017684 001

TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION  
 PHOSPHOTEC  
 BRACCO N/A N017680 001

TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION  
 RBC-SCAN  
 CADEMA N/A N020063 001 Jun 11, 1992

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION  
 MIRALUMA  
 LANTHEUS MEDCL N/A N019785 003 May 23, 1997

TECHNETIUM TC-99M SODIUM PERTECHNETATE

SOLUTION; INJECTION, ORAL SODIUM PERTECHNETATE TC 99M		
+ GE HEALTHCARE	2-100mCi/ML **	N017471 001
+ MALLINCKRODT	10-60mCi/ML **	N017725 001
PHARMALUCENCE	12mCi/ML	N017321 001
	24mCi/ML	N017321 002
	48mCi/ML	N017321 003

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL  
 MINITEC  
 BRACCO 0.22-2.22 CI/GENERATOR N017339 001

SOLUTION; INTRAVENOUS  
 TECHNELITE  
 LANTHEUS MEDCL 0.0083-2.7 CI/GENERATOR N017771 001

ULTRA-TECHNEKOW FM  
 MALLINKRODT NUCLEAR 0.25-3 CI/GENERATOR N017243 002

SOLUTION; INTRAVENOUS, ORAL  
 TECHNETIUM TC 99M GENERATOR  
 GE HEALTHCARE 830-16600mCi/GENERATOR N017693 001

TECHNETIUM TC-99M SUCCIMER KIT

INJECTABLE; INJECTION  
 MPI DMSA KIDNEY REAGENT  
 GE HEALTHCARE N/A N017944 001 May 18, 1982

TECHNETIUM TC-99M SULFUR COLLOID

SOLUTION; INJECTION, ORAL  
 TECHNETIUM TC 99M SULFUR COLLOID  
 GE HEALTHCARE 4mCi/ML N017456 001

SOLUTION; ORAL  
 TECHNETIUM TC 99M SULFUR COLLOID  
 MALLINCKRODT 3mCi/ML N017724 001

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL TECHNECOLL MALLINCKRODT	N/A	N017059 001
TECHNETIUM TC 99M TSC GE HEALTHCARE	N/A	N017784 001
TESULOID BRACCO	N/A	N016923 001

**DISCONTINUED DRUG PRODUCT LIST**

6-341(of 375)

\*\* See List Footnote

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE; INJECTION

CARDIOTEC

BRACCO

N/A

N0119928 001 Dec 19, 1990

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW

+ GE HEALTHCARE

N/A

N020372 001 Feb 09, 1996

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

US WORLDMEDS LLC

EQ 2MG BASE

N021200 001 Jul 24, 2002

EQ 6MG BASE

N021200 002 Jul 24, 2002

TELAPREVIR

TABLET; ORAL

INCIVEK

VERTEX PHARMS

375MG

N201917 001 May 23, 2011

TELAVANCIN HYDROCHLORIDE

POWDER; IV (INFUSION)

VIBATIV

+ THERAVANCE BIOPHARMA EQ 250MG BASE/VIAL

N022110 001 Sep 11, 2009

TELBIVUDINE

SOLUTION; ORAL

TYZEKA

NOVARTIS

100MG/5ML

N022154 001 Apr 28, 2009

TABLET; ORAL

TYZEKA

+ NOVARTIS

600MG

N022011 001 Oct 25, 2006

TELITHROMYCIN

TABLET; ORAL

KETEK

SANOFI AVENTIS US

300MG

N021144 002 Feb 09, 2005

400MG

N021144 001 Apr 01, 2004

TEMAZEPAM

CAPSULE; ORAL

TEMAZ

QUANTUM PHARMICS

15MG

A070564 001 Oct 15, 1985

30MG

A070547 001 Oct 15, 1985

TEMAZEPAM

DURAMED PHARMS BARR

15MG

A071708 001 Sep 29, 1988

30MG

A071709 001 Sep 29, 1988

SUN PHARM INDUSTRIES

15MG

A071174 001 Jul 10, 1986

30MG

A071175 001 Jul 10, 1986

USL PHARMA

15MG

A070489 001 Jul 07, 1986

30MG

A070490 001 Jul 07, 1986

WATSON LABS

15MG

A070383 001 Mar 23, 1987

15MG

A071446 001 May 21, 1993

30MG

A070384 001 Mar 23, 1987

30MG

A071447 001 May 21, 1993

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

+ HQ SPECLT PHARMA

10MG/ML

N020119 001 Jul 14, 1992

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

HYTRIN

+ ABBOTT

EQ 1MG BASE \*\*

N020347 001 Dec 14, 1994

+

EQ 2MG BASE \*\*

N020347 002 Dec 14, 1994

+

EQ 5MG BASE \*\*

N020347 003 Dec 14, 1994

+

EQ 10MG BASE \*\*

N020347 004 Dec 14, 1994

TERAZOSIN HYDROCHLORIDE

MYLAN TECHNOLOGIES

EQ 1MG BASE

A075384 001 Dec 01, 2000

EQ 2MG BASE

A075384 002 Dec 01, 2000

EQ 5MG BASE

A075384 003 Dec 01, 2000

EQ 10MG BASE

A075384 004 Dec 01, 2000

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-342(of 375)

\*\* See List Footnote

TERAZOSIN HYDROCHLORIDE

CAPSULE;ORAL

TERAZOSIN HYDROCHLORIDE

RANBAXY LABS LTD

EQ 1MG BASE

A076021 001 Aug 22, 2002

EQ 2MG BASE

A076021 002 Aug 22, 2002

EQ 5MG BASE

A076021 003 Aug 22, 2002

EQ 10MG BASE

A076021 004 Aug 22, 2002

SANDOZ

EQ 1MG BASE

A075667 001 Jul 28, 2000

EQ 2MG BASE

A075667 002 Jul 28, 2000

EQ 5MG BASE

A075667 003 Jul 28, 2000

EQ 10MG BASE

A075667 004 Jul 28, 2000

TABLET;ORAL

HYTRIN

ABBOTT

EQ 1MG BASE

N019057 001 Aug 07, 1987

EQ 2MG BASE

N019057 002 Aug 07, 1987

EQ 5MG BASE

N019057 003 Aug 07, 1987

EQ 10MG BASE

N019057 004 Aug 07, 1987

TERAZOSIN HYDROCHLORIDE

IVAX SUB TEVA PHARMS

EQ 1MG BASE

A074530 001 Apr 21, 2000

EQ 2MG BASE

A074530 002 Apr 21, 2000

EQ 5MG BASE

A074530 003 Apr 21, 2000

EQ 10MG BASE

A074530 004 Apr 21, 2000

SANDOZ

EQ 1MG BASE

A074315 001 Dec 31, 1998

EQ 2MG BASE

A074657 001 Apr 28, 2000

EQ 5MG BASE

A074315 002 Dec 31, 1998

EQ 10MG BASE

A074657 003 Apr 28, 2000

EQ 10MG BASE

A074315 004 Dec 31, 1998

TEVA

EQ 1MG BASE

A074657 004 Apr 28, 2000

EQ 2MG BASE

A074446 001 May 18, 2000

EQ 5MG BASE

A074446 002 May 18, 2000

EQ 10MG BASE

A074446 003 May 18, 2000

EQ 10MG BASE

A074446 004 May 18, 2000

TERBINAFINE

GEL;TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS 1%

N020846 001 Apr 29, 1998

TERBINAFINE HYDROCHLORIDE

CREAM;TOPICAL

LAMISIL

NOVARTIS

1%

N020192 001 Dec 30, 1992

GRANULE;ORAL

LAMISIL

+ NOVARTIS

EQ 125MG BASE/PACKET

N022071 001 Sep 28, 2007

+

EQ 187.5MG BASE/PACKET

N022071 002 Sep 28, 2007

SOLUTION;TOPICAL

LAMISIL

GLAXOSMITHKLINE CONS 1%

N020749 001 Oct 17, 1997

TABLET;ORAL

TERBINAFINE HYDROCHLORIDE

GEDEON RICHTER USA

EQ 250MG BASE

A077065 001 Jul 02, 2007

MYLAN

EQ 250MG BASE

A077136 001 Jul 02, 2007

ROXANE

EQ 250MG BASE

A077195 001 Jul 02, 2007

WOCKHARDT

EQ 250MG BASE

A077223 001 Jul 02, 2007

A078229 001 Jul 02, 2007

TERBUTALINE SULFATE

AEROSOL, METERED;INHALATION

BRETHAIRE

NOVARTIS

0.2MG/INH

N018762 001 Aug 17, 1984

BRICANYL

SANOFI AVENTIS US

0.2MG/INH

N018000 001 Mar 19, 1985

INJECTABLE;INJECTION

BRETHINE

+ PHARMACARE

1MG/ML \*\*

N018571 001

BRICANYL

SANOFI AVENTIS US

1MG/ML

N017466 001

**DISCONTINUED DRUG PRODUCT LIST**

6-343(of 375)

\*\* See List Footnote

TERBUTALINE SULFATE

INJECTABLE; INJECTION		
TERBUTALINE SULFATE		
TEVA PHARMS USA	1MG/ML	A076853 001 Jul 20, 2004
TABLET; ORAL		
BRETHINE		
+ ANI PHARMS INC	2.5MG **	N017849 001
+	5MG **	N017849 002
BRICANYL		
SANOFI AVENTIS US	2.5MG	N017618 001
	5MG	N017618 002

TERCONAZOLE

CREAM; VAGINAL		
TERAZOL 3		
+ JANSSEN PHARMS	0.8%	N019964 001 Feb 21, 1991
SUPPOSITORY; VAGINAL		
TERCONAZOLE		
FOUGERA PHARMS	80MG	A076850 001 Jul 12, 2006

TERIPARATIDE ACETATE

INJECTABLE; INJECTION		
PARATHAR		
SANOFI AVENTIS US	200 UNITS/VIAL	N019498 001 Dec 23, 1987

TERIPARATIDE RECOMBINANT HUMAN

INJECTABLE; SUBCUTANEOUS		
FORTEO		
LILLY	0.75MG/3ML (0.25MG/ML)	N021318 001 Nov 26, 2002

TESAMORELIN ACETATE

POWDER; SUBCUTANEOUS		
EGRIFTA		
THERATECHNOLOGIES	EQ 2MG BASE/VIAL	N022505 002 Nov 29, 2011

TESTOLACTONE

INJECTABLE; INJECTION		
TESLAC		
BRISTOL MYERS SQUIBB	100MG/ML	N016119 001
TABLET; ORAL		
TESLAC		
BRISTOL MYERS SQUIBB	50MG	N016118 001
	250MG	N016118 002

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL		
ANDRODERM		
ALLERGAN SALES LLC	2.5MG/24HR	N020489 001 Sep 29, 1995
	5MG/24HR	N020489 002 May 02, 1997
TESTODERM		
ALZA	4MG/24HR	N019762 001 Oct 12, 1993
	6MG/24HR	N019762 002 Oct 12, 1993
TESTODERM TTS		
ALZA	5MG/24HR	N020791 001 Dec 18, 1997
INJECTABLE; INJECTION		
TESTOSTERONE		
WATSON LABS	25MG/ML	A086420 001 May 10, 1983
	50MG/ML	A086419 001 Aug 23, 1983
	100MG/ML	A086417 001 Jul 07, 1983
SOLUTION, METERED; TRANSDERMAL		
AXIRON		
+ ELI LILLY AND CO	30MG/1.5ML ACTUATION	N022504 001 Nov 23, 2010

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION		
DEPO-TESTOSTERONE		
PHARMACIA AND UPJOHN	50MG/ML	A085635 001
TESTOSTERONE CYPIONATE		
WATSON LABS	100MG/ML	A084401 001
	100MG/ML	A086029 001
	200MG/ML	A084401 002

**DISCONTINUED DRUG PRODUCT LIST**

6-344(of 375)

\*\* See List Footnote

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DELATESTRYL

ENDO PHARMS	200MG/ML
TESTOSTERONE ENANTHATE	
WATSON LABS	100MG/ML
	100MG/ML
	200MG/ML

N009165	001
A083667	001
A085599	001
A083667	002

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

BEL MAR	25MG/ML
	50MG/ML
	100MG/ML
ELKINS SINK	25MG/ML
LILLY	50MG/ML
WATSON LABS	25MG/ML
	25MG/ML
	50MG/ML
	50MG/ML
	100MG/ML
	100MG/ML

A080741	001
A080742	001
A080743	001
A080276	001
A080254	002
A080188	001
A085490	001
A080188	002
A085490	002
A080188	003
A083595	003

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

BRISTACYCLINE

BRISTOL	250MG
	250MG
	500MG
	500MG

A061658	001
A061888	001
A061658	002
A061888	002

CYCLOPAR

WARNER CHILCOTT	250MG
	250MG
	250MG
	500MG
	500MG

A061725	001
A062175	001
A062332	001
A061725	002
A062332	002

PANMYCIN

PHARMACIA AND UPJOHN	250MG
----------------------	-------

A060347	001
---------	-----

RETEF

SOLVAY	250MG
	500MG

A061443	001
A061443	002

ROBITET

WYETH AYERST	250MG
	500MG

A061734	001
A061734	002

SUMYCIN

APOTHECON	100MG
	125MG
	250MG
	500MG

A060429	002
A060429	004
A060429	001
A060429	003

TETRACHEL

ANGUS	250MG
	500MG

A060343	001
A060343	003

TETRACYCLINE HYDROCHLORIDE

ABBOTT	250MG
	500MG
ELKINS SINK	250MG
FERRANTE	125MG
	250MG
HEATHER	250MG
	500MG
HIKMA PHARMS	250MG
	500MG
IDT AUSTRALIA LTD	250MG
IMPAX LABS	100MG
	250MG
	500MG
IVAX SUB TEVA PHARMS	250MG
	500MG
MAST MM	250MG

A061802	001
A061802	002
A060059	001
A060173	001
A060173	002
A061148	001
A061148	002
A060768	001
A060768	002
A061471	001
A060469	002
A060469	001
A060469	003
A060704	001
A060704	002
A062085	001

**DISCONTINUED DRUG PRODUCT LIST**

6-345(of 375)

\*\* See List Footnote

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

MYLAN	250MG	A060783 001
	500MG	A060783 002
PUREPAC PHARM	250MG	A060290 001
	500MG	A060290 002
PVT FORM	250MG	A062686 001 Jul 24, 1986
	500MG	A062686 002 Jul 24, 1986
ROXANE	500MG	A061214 002
SUN PHARM INDUSTRIES	250MG	A060736 001
	500MG	A060736 002
SUPERPHARM	250MG	A062540 001 Mar 21, 1985
	500MG	A062540 002 Mar 21, 1985
VALEANT PHARM INTL	250MG	A060471 001
	500MG	A060471 002
WARNER CHILCOTT	250MG	A062300 001
	500MG	A062300 002
WATSON LABS	250MG	A062103 001
	250MG	A062343 001
	500MG	A062103 002
	500MG	A062343 002
WYETH AYERST	250MG	A061685 001
	500MG	A061685 002

TETRACYCIN

PFI-PHARMECS	250MG	A060082 003
	500MG	A060082 004

FIBER, EXTENDED RELEASE; PERIODONTAL

ACTISITE

SCHIFF AND CO	12.7MG/FIBER	N050653 001 Mar 25, 1994
---------------	--------------	--------------------------

FOR SOLUTION; TOPICAL

TOPICYCLINE

SHIRE	2.2MG/ML	N050493 001
-------	----------	-------------

INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	250MG/VIAL	N050273 002
	500MG/VIAL	N050273 003

TETRACYCIN

Pfizer	250MG/VIAL	A060096 001
	500MG/VIAL	A060096 002

OINTMENT; OPHTHALMIC

ACHROMYCIN

STORZ	10MG/GM	N050266 001
-------	---------	-------------

SUSPENSION; ORAL

ACHROMYCIN V

LEDERLE	125MG/5ML	N050263 002
---------	-----------	-------------

SUMYCIN

PAR PHARM	125MG/5ML	A060400 001
-----------	-----------	-------------

TETRACYCLINE HYDROCHLORIDE

ALPHARMA US PHARMS	125MG/5ML	A060633 001
FERRANTE	125MG/5ML	A060174 001
PROTER	125MG/5ML	A060446 001
PUREPAC PHARM	125MG/5ML	A060291 001

TETRACYCIN

PFI-PHARMECS	125MG/5ML	A060095 001
--------------	-----------	-------------

TETRAMED

IVAX SUB TEVA PHARMS	125MG/5ML	A061468 001
----------------------	-----------	-------------

SUSPENSION/DROPS; OPHTHALMIC

ACHROMYCIN

STORZ	1%	N050268 001
-------	----	-------------

TABLET; ORAL

PANMYCIN

PHARMACIA AND UPJOHN	250MG	A061705 001
	500MG	A061705 002

SUMYCIN

PAR PHARM	50MG	A061147 003
	100MG	A061147 002
	250MG	A061147 001
	500MG	A061147 004

**DISCONTINUED DRUG PRODUCT LIST**

6-346(of 375)

\*\* See List Footnote

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE;ORAL

TETREX

BRISTOL

EQ 100MG HCL  
EQ 250MG HCL  
EQ 250MG HCL  
EQ 250MG HCL  
EQ 500MG HCL  
EQ 500MG HCL  
EQ 500MG HCLA061653 001  
A061653 002  
A061889 002  
N050212 002  
A061653 003  
A061889 001  
N050212 003THALLOUS CHLORIDE TL-201

INJECTABLE;INJECTION

THALLOUS CHLORIDE TL 201

BRACCO

1mCi/ML

N018548 001 Dec 30, 1982

TRACE LIFE

1mCi/ML

A075569 001 Nov 21, 2001

INJECTABLE;INTRAVENOUS

THALLOUS CHLORIDE TL 201

MALLINKRODT NUCLEAR

2mCi/ML

A077698 001 Nov 09, 2006

THEOPHYLLINE

CAPSULE;ORAL

BRONKODYL

SANOFI AVENTIS US

100MG  
200MGA085264 001  
A085264 002

ELIXOPHYLLIN

FOREST LABS

100MG  
200MGA085545 001 Jul 31, 1984  
A083921 001 Jul 31, 1984

SOMOPHYLLIN-T

FISONS

100MG  
200MG  
250MGA087155 001 Feb 25, 1985  
A087155 002 Feb 25, 1985  
A087155 003 Feb 25, 1985

THEOPHYLLINE

KV PHARM

100MG  
200MG  
200MG  
250MGA085263 001  
A085263 002  
A084731 002 Nov 07, 1986  
A084731 001 Nov 07, 1986  
A084731 003 Nov 07, 1986

CAPSULE, EXTENDED RELEASE;ORAL

AEROLATE III

FLEMING PHARMS

65MG

A085075 003 Nov 24, 1986

AEROLATE JR

FLEMING PHARMS

130MG

A085075 002 Nov 24, 1986

AEROLATE SR

FLEMING PHARMS

260MG

A085075 001 Nov 24, 1986

ELIXOPHYLLIN SR

FOREST LABS

125MG  
250MGA086826 001 Jan 29, 1985  
A086826 002 Jan 29, 1985

SLO-BID

SANOFI AVENTIS US

50MG  
75MG  
100MG  
125MG  
200MG  
300MGA088269 001 Jan 31, 1985  
A089539 001 May 10, 1989  
A087892 001 Jan 31, 1985  
A089540 001 May 10, 1989  
A087893 001 Jan 31, 1985  
A087894 001 Jan 31, 1985

SLO-PHYLLIN

SANOFI AVENTIS US

60MG  
125MG  
250MGA085206 001 May 24, 1982  
A085203 001 May 24, 1982  
A085205 001 May 24, 1982

SOMOPHYLLIN-CRT

GRAHAM DM

50MG  
100MG  
200MG  
250MG  
300MGA087763 001 Feb 27, 1985  
A087194 001  
A088382 001 Feb 27, 1985  
A087193 001  
A088383 001 Feb 27, 1985

THEO-DUR

SCHERING

50MG  
75MG  
125MG  
200MGA088022 001 Sep 10, 1985  
A088015 001 Sep 10, 1985  
A088016 001 Sep 10, 1985  
A087995 001 Sep 10, 1985

**DISCONTINUED DRUG PRODUCT LIST**

6-347(of 375)

\*\* See List Footnote

**THEOPHYLLINE**

CAPSULE, EXTENDED RELEASE;ORAL

THEOBID					
WHITBY	260MG			A085983	001 Mar 20, 1985
THEOBID JR.				A087854	001 Mar 20, 1985
WHITBY	130MG			A086569	001 May 27, 1982
THEOCLEAR L.A.-130				A086569	002 May 27, 1982
SCHWARZ PHARMA	130MG			A086480	001 Feb 08, 1985
THEOCLEAR L.A.-260				A086471	001 Feb 08, 1985
SCHWARZ PHARMA	260MG			A088654	001 Feb 12, 1985
THEOPHYL-SR				A088689	001 Feb 12, 1985
ORTHO MCNEIL PHARM	125MG			A089976	001 Jan 04, 1995
250MG				A089977	001 Jan 04, 1995
THEOPHYLLINE				A089932	001 Jan 04, 1995
CENT PHARMS	125MG			A040052	001 Feb 14, 1994
250MG				A040052	002 Feb 14, 1994
HOSPIRA	100MG			A040052	003 Feb 14, 1994
200MG				A040052	004 Feb 14, 1994
INWOOD LABS	300MG			A087462	001 May 11, 1982
100MG				A088255	001 Jun 12, 1986
125MG				A087010	001 Jan 31, 1985
200MG				A087910	001 Jan 31, 1985
300MG					
SANDOZ	260MG				
THEOPHYLLINE-SR					
SCHERER RP	300MG				
THEOVENT					
SCHERING	125MG				
250MG					
ELIXIR;ORAL					
ELIXOMIN					
CENCI	80MG/15ML			A088303	001 Jan 25, 1984
LANOPHYLLIN				A084578	001
LANNETT	80MG/15ML			A084559	001
THEOLIXIR				A086485	001
PANRAY	80MG/15ML			A089223	001 May 27, 1988
THEOPHYL-225				A087679	001 Apr 15, 1982
ORTHO MCNEIL PHARM	112.5MG/15ML			A085952	001
THEOPHYLLINE				A085169	001
ALPHARMA US PHARMS	80MG/15ML			A086720	001
CENCI	80MG/15ML			A085863	001
CHARTWELL RX	80MG/15ML			A084739	001
HALSEY	80MG/15ML			A089626	001 Oct 28, 1988
PHARM ASSOC	80MG/15ML			A086748	001
PRECISION DOSE	80MG/15ML				
ROXANE	80MG/15ML				
TARO	80MG/15ML				
WOCKHARDT	80MG/15ML				
INJECTABLE;INJECTION					
THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER				N019083	001 Nov 07, 1984
B BRAUN	40MG/100ML			N019083	002 Nov 07, 1984
THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER				N019083	003 Nov 07, 1984
B BRAUN	80MG/100ML			N019212	001 Nov 07, 1984
THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER				N019826	004 Aug 14, 1992
B BRAUN	160MG/100ML			N019212	003 Nov 07, 1984
THEOPHYLLINE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER				N019826	005 Aug 14, 1992
B BRAUN	200MG/100ML			N019212	001 Nov 07, 1984
200MG/100ML				N019212	002 Nov 07, 1984
THEOPHYLLINE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER				N019826	004 Aug 14, 1992
B BRAUN	4MG/ML			N019212	003 Nov 07, 1984
400MG/100ML				N019212	002 Nov 07, 1984
400MG/100ML				N019826	005 Aug 14, 1992
THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER				N018649	007 Jul 26, 1982
BAXTER HLTHCARE	4MG/ML			N018649	001 Jul 26, 1982
40MG/100ML				N018649	002 Jul 26, 1982
80MG/100ML				N018649	003 Jul 26, 1982
160MG/100ML				N018649	004 Jul 26, 1982
200MG/100ML				N018649	006 Nov 13, 1985
320MG/100ML					

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-348(of 375)

\*\* See List Footnote

**THEOPHYLLINE****INJECTABLE; INJECTION**

THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER	400MG/100ML	N018649 005 Jul 26, 1982
THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER		
HOSPIRA INC	80MG/100ML	N019211 002 Dec 14, 1984
	200MG/100ML	N019211 004 Dec 14, 1984
	400MG/100ML	N019211 005 Dec 14, 1984

**SOLUTION;ORAL**

AEROLATE		
FLEMING PHARMS	150MG/15ML	A089141 001 Dec 03, 1986
THEOLAIR		
3M	80MG/15ML	A086107 001
THEOPHYLLINE		
ROXANE	80MG/15ML	A087449 001 Sep 15, 1983
SUSPENSION;ORAL		
ELIXICON		
FOREST LABS	100MG/5ML	A085502 001
SYRUP;ORAL		
ACCURBRON		
SANOFI AVENTIS US	150MG/15ML	A088746 001 Nov 22, 1985
AQUAPHYLLIN		
FERNDALE LABS	80MG/15ML	A087917 001 Jan 18, 1983
SLO-PHYLLIN		
SANOFI AVENTIS US	80MG/15ML	A085187 001
THEOCLEAR-80		
CENT PHARMS	80MG/15ML	A087095 001 Mar 01, 1982
THEOPHYLLINE		
ALPHARMA US PHARMS	80MG/15ML	A086001 001
	150MG/15ML	A086545 001

**TABLET;ORAL**

QUIBRON-T		
MONARCH PHARMS	300MG	A088656 001 Aug 22, 1985
SLO-PHYLLIN		
SANOFI AVENTIS US	100MG	A085202 001
	200MG	A085204 001
THEOCLEAR-100		
CENT PHARMS	100MG	A085353 002
THEOCLEAR-200		
CENT PHARMS	200MG	A085353 001
THEOLAIR		
MEDICIS	125MG	A086399 001
	250MG	A086399 002
THEOPHYL-225		
ORTHO MCNEIL PHARM	225MG	A084726 001
TABLET, CHEWABLE;ORAL		
THEOPHYL		
ORTHO MCNEIL PHARM	100MG	A086506 001 Sep 12, 1985
TABLET, EXTENDED RELEASE;ORAL		
DURAPHYL		
FOREST LABS	100MG	A088503 001 Apr 03, 1985
	200MG	A088504 001 Apr 03, 1985
	300MG	A088505 001 Apr 03, 1985
LABID		
WARNER CHILCOTT	250MG	A087225 001
QUIBRON-T/SR		
MONARCH PHARMS	300MG	A087563 001 Jun 21, 1983
SUSTAIRE		
ROERIG	100MG	A085665 001
	300MG	A085665 002
T-PHYL		
PHARM RES ASSOC	200MG	A088253 001 Aug 17, 1983
THEO-DUR		
SCHERING	100MG	A085328 001
	200MG	A086998 001
	300MG	A085328 002
	450MG	A089131 001 Jun 25, 1986
THEOCHRON		
NOSTRUM PHARMS LLC	300MG	A087400 002 Jan 11, 1983

**DISCONTINUED DRUG PRODUCT LIST**

6-349(of 375)

\*\* See List Footnote

**THEOPHYLLINE**

TABLET, EXTENDED RELEASE;ORAL

THEOLAIR-SR

3M	200MG	A088369 001 Jul 16, 1987
	250MG	A086363 002 Jul 16, 1987
	300MG	A088364 001 Jul 16, 1987
	500MG	A089132 001 Jul 16, 1987
THEOPHYLLINE ABLE	300MG	A040548 001 Apr 30, 2004
	400MG	A040543 001 Apr 27, 2004
	450MG	A040546 001 Apr 30, 2004
	600MG	A040539 001 Apr 27, 2004
INWOOD LABS	450MG	A040034 001 Apr 28, 1995
TEVA PHARMS	450MG	A081236 001 Nov 09, 1992
UNI-DUR SCHERING	400MG	A089822 001 Jan 04, 1995
	600MG	A089823 001 Jan 04, 1995

**THEOPHYLLINE SODIUM GLYCINATE**

ELIXIR;ORAL

SYNOPHYLATE

CENT PHARMS	EQ 165MG BASE/15ML	N006333 008
TABLET;ORAL ASBRON		
NOVARTIS	EQ 150MG BASE	A085148 001

**THIABENDAZOLE**

SUSPENSION;ORAL

MINTEZOL

MERCK SHARP DOHME	500MG/5ML	N016097 001
TABLET, CHEWABLE;ORAL MINTEZOL		

MERCK SHARP DOHME	500MG	N016096 001
-------------------	-------	-------------

**THIAMINE HYDROCHLORIDE**

INJECTABLE; INJECTION

BETALIN S

LILLY	100MG/ML	A080853 001
THIAMINE HYDROCHLORIDE		
ABRAXIS PHARM	100MG/ML	A080509 001
AKORN	100MG/ML	A087968 001 Oct 01, 1982
BEL MAR	100MG/ML	A080718 001
	200MG/ML	A080712 001
DELL LABS	100MG/ML	A083775 001
HOSPIRA	100MG/ML	A040079 001 May 03, 1996
LUITPOLD	100MG/ML	A080667 001
PARKE DAVIS	100MG/ML	A080770 001
WATSON LABS	100MG/ML	A080571 001
	100MG/ML	A083534 001
	200MG/ML	A080571 002
	200MG/ML	A083534 002
WEST-WARD PHARMS INT	100MG/ML	A080575 001
WYETH AYERST	100MG/ML	A080553 001

**THIAMYLAL SODIUM**

INJECTABLE; INJECTION

SURITAL

PARKEDALE	1GM/VIAL	N007600 003
	5GM/VIAL	N007600 005
	10GM/VIAL	N007600 009

**THIETHYLPERAZINE MALATE**

INJECTABLE; INJECTION

TORECAN

NOVARTIS	5MG/ML	N012754 002
----------	--------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-350(of 375)

\*\* See List Footnote

THIETHYLPERAZINE MALEATE

SUPPOSITORY;RECTAL

TORECAN

NOVARTIS

10MG

N013247 001

TABLET;ORAL

TORECAN

NOVARTIS

10MG

N012753 001

THIOPENTAL SODIUM

SUSPENSION;RECTAL

PENTOTHAL

ABBOTT

400MG/GM

N011679 001

THIORIDAZINE

SUSPENSION;ORAL

MELLARIL-S

NOVARTIS

EQ 25MG HCL/5ML \*\*

N017923 001

EQ 100MG HCL/5ML \*\*

N017923 002

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE;ORAL

MELLARIL

NOVARTIS

30MG/ML \*\*

N011808 012

100MG/ML \*\*

N011808 018

THIORIDAZINE HYDROCHLORIDE

ACTAVIS MID ATLANTIC

100MG/ML

A088229 001 Aug 23, 1983

ALPHARMA US PHARMS

30MG/ML

A087766 001 Apr 26, 1983

ANI PHARMS INC

30MG/ML

A089602 001 Nov 09, 1987

100MG/ML

A089603 001 Nov 09, 1987

HI TECH PHARMA

30MG/ML

A040125 001 Aug 16, 1996

100MG/ML

A040126 001 Aug 16, 1996

PHARM ASSOC

30MG/ML

A040187 001 Aug 28, 1997

100MG/ML

A040213 001 May 29, 1998

SANDOZ

30MG/ML

A088307 001 Nov 23, 1983

100MG/ML

A088308 001 Nov 23, 1983

WOCKHARDT

30MG/ML

A088258 001 Jul 25, 1983

100MG/ML

A088227 001 Jul 05, 1983

THIORIDAZINE HYDROCHLORIDE INTENSOL

ROXANE

30MG/ML

A088941 001 Dec 16, 1985

100MG/ML

A088942 001 Dec 16, 1985

TABLET;ORAL

MELLARIL

+ NOVARTIS

10MG \*\*

N011808 003

+

15MG \*\*

N011808 016

+

25MG \*\*

N011808 006

+

50MG \*\*

N011808 011

+

100MG \*\*

N011808 009

+

150MG \*\*

N011808 017

+

200MG \*\*

N011808 015

THIORIDAZINE HYDROCHLORIDE

ANI PHARMS INC

10MG

A088270 001 Apr 14, 1983

10MG

A088493 001 May 17, 1985

15MG

A088271 001 Apr 14, 1983

25MG

A088272 001 Apr 14, 1983

50MG

A088194 001 Apr 14, 1983

100MG

A088273 001 Oct 03, 1983

100MG

A088456 001 May 17, 1985

MUTUAL PHARM

10MG

A088375 001 Nov 18, 1983

25MG

A087264 001 Nov 18, 1983

50MG

A088370 001 Nov 18, 1983

100MG

A088379 001 Nov 16, 1983

MYLAN

10MG

A088332 001 Jun 27, 1983

25MG

A088333 001 Jun 27, 1983

50MG

A088334 001 Jun 27, 1983

100MG

A088335 001 Nov 18, 1983

PAR PHARM

10MG

A088351 001 Dec 05, 1983

15MG

A088352 001 Dec 05, 1983

25MG

A088336 001 Dec 05, 1983

50MG

A088322 001 Dec 05, 1983

100MG

A088480 001 Dec 29, 1983

150MG

A089764 001 Feb 09, 1988

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-351(of 375)

\*\* See List Footnote

**THIORIDAZINE HYDROCHLORIDE**

TABLET;ORAL

**THIORIDAZINE HYDROCHLORIDE**

	200MG	A089765 001	Feb 09, 1988
ROXANE	10MG	A088663 001	Mar 15, 1984
	25MG	A088664 001	Mar 15, 1984
	50MG	A088665 001	Mar 15, 1984
	100MG	A089048 001	Feb 26, 1985
SANDOZ	10MG	A088131 001	Aug 30, 1983
	15MG	A088132 001	Aug 30, 1983
	25MG	A088133 001	Aug 30, 1983
	50MG	A088134 001	Aug 30, 1983
	100MG	A088135 001	Nov 20, 1984
	150MG	A088136 001	Sep 17, 1986
	200MG	A088137 001	Sep 17, 1986
SUN PHARM INDUSTRIES	15MG	A088461 001	Nov 18, 1983
	150MG	A088737 001	Sep 26, 1984
	200MG	A088738 001	Oct 16, 1984
SUPERPHARM	10MG	A089103 001	Jul 02, 1985
	25MG	A089104 001	Jul 02, 1985
	50MG	A089105 001	Jul 02, 1985
WATSON LABS	10MG	A088412 001	Sep 12, 1983
	10MG	A088476 001	Nov 08, 1983
	10MG	A088561 001	May 11, 1984
	15MG	A088345 001	Jul 28, 1983
	15MG	A088562 001	May 11, 1984
	25MG	A088296 001	Jul 28, 1983
	25MG	A088478 001	Nov 08, 1983
	25MG	A088755 001	Jul 24, 1984
	50MG	A088323 001	Jul 28, 1983
	50MG	A088479 001	Nov 08, 1983
	50MG	A088563 001	May 11, 1984
	100MG	A088284 001	Aug 25, 1983
	100MG	A088564 001	May 11, 1984
	100MG	A088736 001	Jul 24, 1984
	150MG	A088410 001	Mar 05, 1984
	150MG	A088869 001	Jun 28, 1985
	200MG	A088381 001	Mar 14, 1984
WATSON LABS TEVA	15MG	A088477 001	Nov 08, 1983
	25MG	A088567 001	May 11, 1984
	200MG	A088872 001	Apr 26, 1985
WEST WARD	10MG	A088658 001	Mar 26, 1984
	15MG	A088659 001	Mar 26, 1984
	25MG	A088660 001	Mar 26, 1984
	50MG	A088661 001	Mar 26, 1984

**THIOTEP A**

INJECTABLE;INJECTION

**THIOPLEX**

+ IMMUNEX	15MG/VIAL **	N020058 001	Dec 22, 1994
THIOTEP A			
FRESENIUS KABI USA	15MG/VIAL	A075698 001	Sep 20, 2001
IMMUNEX	15MG/VIAL	N011683 001	
TEVA PARENTERAL	15MG/VIAL **	A075730 001	Apr 20, 2001
	30MG/VIAL **	A075730 002	Apr 20, 2001

**THIOTHIXENE**

CAPSULE;ORAL

**NAVANE**

PFIZER	1MG **	N016584 001	
	2MG **	N016584 002	
	5MG **	N016584 003	
	10MG **	N016584 004	
	20MG **	N016584 005	

**THIOTHIXENE**

AM THERAP	1MG	A071884 001	Aug 12, 1987
	2MG	A071885 001	Aug 12, 1987
	5MG	A071886 001	Aug 12, 1987
	10MG	A071887 001	Aug 12, 1987
	20MG	A072200 001	Dec 17, 1987

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-352(of 375)

\*\* See List Footnote

**THIOTHIXENE**

CAPSULE;ORAL

THIOTHIXENE

SANDOZ

1MG

A071529 002 Jun 24, 1987

2MG

A071529 003 Jun 24, 1987

5MG

A071529 001 Jun 24, 1987

10MG

A071529 004 Jun 24, 1987

WATSON LABS

1MG

A070600 001 Jun 05, 1987

2MG

A070601 001 Jun 05, 1987

2MG

A071626 001 Jun 25, 1987

5MG

A070602 001 Jun 05, 1987

5MG

A071627 001 Jun 25, 1987

10MG

A070603 001 Jun 05, 1987

10MG

A071628 001 Jun 25, 1987

**THIOTHIXENE HYDROCHLORIDE**

CONCENTRATE;ORAL

NAVANE

PFIZER

EQ 5MG BASE/ML

N016758 001

THIOTHIXENE HYDROCHLORIDE

ALPHARMA US PHARMS

EQ 5MG BASE/ML

A070969 001 Oct 16, 1987

PACO

EQ 1MG BASE/ML

A071917 001 Sep 20, 1989

EQ 5MG BASE/ML

A071939 001 Dec 16, 1988

TEVA

EQ 5MG BASE/ML

A071184 001 Jun 22, 1987

TEVA PHARMS

EQ 5MG BASE/ML

A071554 001 Oct 16, 1987

THIOTHIXENE HYDROCHLORIDE INTENSOL

CYCLE PHARMS LTD

EQ 5MG BASE/ML

A073494 001 Jun 30, 1992

INJECTABLE;INJECTION

NAVANE

PFIZER

EQ 2MG BASE/ML

N016904 001

EQ 10MG BASE/VIAL

N016904 002

**THYROGLOBULIN**

TABLET;ORAL

PROLOID

PARKE DAVIS

16MG

N002245 009

32MG

N002245 005

65MG

N002245 002

100MG

N002245 008

130MG

N002245 010

200MG

N002245 007

325MG

N002245 004

THYROGLOBULIN

IMPAK LABS

64.8MG

A080151 001

**THYROTROPIN**

INJECTABLE;INJECTION

THYTROPAR

SANOFI AVENTIS US

10 IU/VIAL

N008682 001

**TIAGABINE HYDROCHLORIDE**

TABLET;ORAL

GABITRIL

CEPHALON

6MG

N020646 006 Nov 29, 2005

8MG

N020646 007 Nov 29, 2005

10MG

N020646 008 Nov 29, 2005

20MG

N020646 004 Sep 30, 1997

**TICARCILLIN DISODIUM**

INJECTABLE;INJECTION

TICAR

GLAXOSMITHKLINE

EQ 1GM BASE/VIAL

N050497 001

EQ 3GM BASE/VIAL

A062690 001 Dec 19, 1986

EQ 3GM BASE/VIAL

N050497 002

EQ 6GM BASE/VIAL

N050497 003

EQ 20GM BASE/VIAL

N050497 004

EQ 30GM BASE/VIAL

N050497 005 Apr 04, 1984

**DISCONTINUED DRUG PRODUCT LIST**

6-353(of 375)

\*\* See List Footnote

**TICLOPIDINE HYDROCHLORIDE**

TABLET;ORAL

TICLID

ROCHE PALO	125MG	N019979 001 Mar 24, 1993
	250MG	N019979 002 Oct 31, 1991
TICLOPIDINE HYDROCHLORIDE		
ACTAVIS ELIZABETH	250MG	A075253 001 Aug 20, 1999
FOSUN PHARMA	250MG	A075318 001 Aug 20, 1999
	250MG	A075326 001 Aug 20, 1999
MYLAN	250MG	A075161 001 Sep 13, 1999
	250MG	A075316 001 Nov 02, 1999
WATSON LABS	250MG	A075309 001 Apr 26, 2000

**TILUDRONATE DISODIUM**

TABLET;ORAL

SKELID

+ SANOFI AVENTIS US	EQ 200MG BASE **	N020707 001 Mar 07, 1997
---------------------	------------------	--------------------------

**TIMOLOL MALEATE**

SOLUTION/DROPS;OPHTHALMIC

TIMOLOL MALEATE

AKORN	EQ 0.25% BASE	A074465 001 Mar 25, 1997
	EQ 0.25% BASE	A074515 001 Mar 25, 1997
APOTEX INC	EQ 0.25% BASE	A075411 001 Sep 08, 2000
	EQ 0.5% BASE	A075412 001 Sep 08, 2000
FOUGERA	EQ 0.25% BASE	A074667 001 Mar 25, 1997
	EQ 0.5% BASE	A074668 001 Mar 25, 1997
TIMOPTIC		
+ ATON	EQ 0.25% BASE **	N018086 001
+	EQ 0.5% BASE **	N018086 002

TABLET;ORAL

BLOCADREN

MERCK	5MG	N018017 001
	10MG	N018017 002
	20MG	N018017 004

TIMOLOL MALEATE

FOSUN PHARMA	5MG	A072550 001 Apr 13, 1989
	10MG	A072551 001 Apr 13, 1989
	20MG	A072552 001 Apr 13, 1989
QUANTUM PHARMICS	5MG	A072466 001 May 19, 1989
	10MG	A072467 001 May 19, 1989
	20MG	A072468 001 May 19, 1989
TEVA	5MG	A072648 001 Jun 16, 1993
	10MG	A072649 001 Jun 16, 1993
	20MG	A072650 001 Jun 16, 1993
USL PHARMA	5MG	A072001 001 Apr 11, 1989
	10MG	A072002 001 Apr 11, 1989
	20MG	A072003 001 Apr 11, 1989
WATSON LABS	5MG	A072269 001 Apr 11, 1989
	5MG	A072917 001 Jul 31, 1991
	10MG	A072270 001 Apr 11, 1989
	10MG	A072918 001 Jul 31, 1991
	20MG	A072271 001 Apr 11, 1989
	20MG	A072919 001 Jul 31, 1991

**TINZAPARIN SODIUM**

INJECTABLE;INJECTION

INNOHEP

LEO PHARMA AS	20,000 IU/ML	N020484 001 Jul 14, 2000
---------------	--------------	--------------------------

**TIOCONAZOLE**

CREAM;TOPICAL

TZ-3

PFIZER	1%	N018682 001 Feb 18, 1983
--------	----	--------------------------

**TIROFIBAN HYDROCHLORIDE**

INJECTABLE;INJECTION

AGGRASTAT

MEDICURE	EQ 12.5MG BASE/50ML (EQ 0.25MG BASE/ML)	N020912 001 May 14, 1998
	EQ 25MG BASE/500ML (EQ 0.05MG BASE/ML)	N020913 001 May 14, 1998

**DISCONTINUED DRUG PRODUCT LIST**

6-354(of 375)

\*\* See List Footnote

TIZANIDINE HYDROCHLORIDE

TABLET;ORAL

## TIZANIDINE HYDROCHLORIDE

ACTAVIS ELIZABETH	EQ 2MG BASE	A076283 001 Jul 12, 2002
	EQ 4MG BASE	A076283 002 Jul 12, 2002
BARR	EQ 2MG BASE	A076371 001 Apr 09, 2003
	EQ 4MG BASE	A076371 002 Apr 09, 2003
IVAX SUB TEVA PHARMS	EQ 2MG BASE	A076321 001 Sep 30, 2004
	EQ 4MG BASE	A076321 002 Sep 30, 2004
MYLAN PHARMS INC	EQ 2MG BASE	A076282 001 Dec 16, 2003
	EQ 4MG BASE	A076282 002 Dec 16, 2003
ZANAFLEX		
+ COVIS PHARMA BV	EQ 2MG BASE **	N020397 002 Feb 04, 2000

TOBRAMYCIN

SOLUTION/DROPS;OPHTHALMIC

## TOBRAMYCIN

ALCON PHARMS LTD	0.3%	A063176 001 May 25, 1994
APOTEX INC	0.3%	A065087 001 Feb 25, 2002

TOBRAMYCIN SULFATE

INJECTABLE;INJECTION

## NEBCIN

LILLY	EQ 10MG BASE/ML	A062008 004
	EQ 10MG BASE/ML	A062707 001 Apr 29, 1987
+	EQ 10MG BASE/ML **	N050477 005
	EQ 40MG BASE/ML	A062008 001
+	EQ 1.2GM BASE/VIAL **	N050519 001

## TOBRAMYCIN SULFATE

APOTHECON	EQ 10MG BASE/ML	A064021 001 May 31, 1994
	EQ 40MG BASE/ML	A064021 002 May 31, 1994
	EQ 40MG BASE/ML	A064026 001 May 31, 1994
HOSPIRA	EQ 10MG BASE/ML	A063080 001 Apr 30, 1991
	EQ 40MG BASE/ML	A063161 001 May 29, 1991
IGI LABS INC	EQ 10MG BASE/ML	A063119 001 Oct 31, 1994
	EQ 40MG BASE/ML	A063120 001 Oct 31, 1994
	EQ 40MG BASE/ML	A063121 001 Oct 31, 1994
WATSON LABS INC	EQ 10MG BASE/ML	A063122 001 Oct 31, 1994
	EQ 40MG BASE/ML	A062945 001 Aug 09, 1989
	EQ 40MG BASE/ML	A062945 002 Aug 09, 1989
WEST-WARD PHARMS INT	EQ 10MG BASE/ML	A063113 001 Apr 26, 1991
	EQ 10MG BASE/ML	A063128 001 Nov 27, 1991
	EQ 40MG BASE/ML	A063118 001 Jul 29, 1991
	EQ 40MG BASE/ML	A063127 001 Nov 27, 1991
TOBRAMYCIN SULFATE (PHARMACY BULK)		
HOSPIRA	EQ 40MG BASE/ML **	A063116 001 May 18, 1992

TOCAINIDE HYDROCHLORIDE

TABLET;ORAL

## TONOCARD

ASTRAZENECA	400MG	N018257 001 Nov 09, 1984
	600MG	N018257 002 Nov 09, 1984

TOLAZAMIDE

TABLET;ORAL

## TOLAZAMIDE

BARR	100MG	A070162 001 Jan 14, 1986
	250MG	A070163 001 Jan 14, 1986
	500MG	A070164 001 Jan 14, 1986
DURAMED PHARMS BARR	100MG	A070165 001 Jan 10, 1986
	250MG	A070166 001 Jan 10, 1986
	500MG	A070167 001 Jan 10, 1986
FOSUN PHARMA	250MG	A070289 001 Mar 13, 1986
	500MG	A070290 001 Mar 13, 1986
G AND W LABS INC	100MG	N018894 001 Nov 02, 1984
	250MG	N018894 002 Nov 02, 1984
	500MG	N018894 003 Nov 02, 1984
INTERPHARM	250MG	A071270 001 Sep 23, 1986
	500MG	A071271 001 Sep 23, 1986
PAR PHARM	100MG	A070159 001 Jan 06, 1986
	250MG	A070160 001 Jan 06, 1986

**DISCONTINUED DRUG PRODUCT LIST**

6-355(of 375)

\*\* See List Footnote

**TOLAZAMIDE**TABLET;ORAL  
TOLAZAMIDE

SANDOZ	500MG	A070161 001	Jan 06, 1986
SUN PHARM INDUSTRIES	100MG	A071633 001	Dec 09, 1987
	100MG	A071357 001	Jul 16, 1987
	250MG	A071358 001	Jul 16, 1987
	500MG	A071359 001	Jul 16, 1987
SUPERPHARM	250MG	A070763 001	Jun 16, 1986
	500MG	A070764 001	Jun 16, 1986
USL PHARMA	100MG	A071355 001	Jan 11, 1988
	250MG	A070168 001	Apr 02, 1986
	500MG	A070169 001	Apr 02, 1986
WATSON LABS	100MG	A070242 001	Aug 01, 1986
	100MG	A070513 001	Jan 09, 1986
	250MG	A070243 001	Aug 01, 1986
	250MG	A070514 001	Jan 09, 1986
	500MG	A070244 001	Aug 01, 1986
	500MG	A070515 001	Jan 09, 1986
TOLINASE			
+ PHARMACIA AND UPJOHN	100MG **	N015500 002	
+	250MG **	N015500 004	
+	500MG **	N015500 005	

**TOLAZOLINE HYDROCHLORIDE**

INJECTABLE;INJECTION

PRISCOLINE

NOVARTIS

25MG/ML

N006403 005 Feb 22, 1985

**TOLBUTAMIDE**

TABLET;ORAL

ORINASE

PHARMACIA AND UPJOHN 250MG \*\*  
500MG \*\*N010670 002  
N010670 001

TOLBUTAMIDE

ALRA	500MG	A086141 001	
ASCOT	500MG	A087541 001	Mar 01, 1983
BARR	500MG	A087121 001	
DAVA PHARMS INC	500MG	A086926 001	
IVAX PHARMS	500MG	A087093 001	
PARKE DAVIS	500MG	A086047 001	
PUREPAC PHARM	500MG	A088950 001	Jun 17, 1985
SANDOZ	500MG	A086574 001	
	500MG	N012678 001	
SUPERPHARM	500MG	A088893 001	Nov 19, 1984
VANGARD	500MG	A087876 001	Apr 20, 1982
WATSON LABS	250MG	A089110 001	May 29, 1987
	500MG	A086109 001	
	500MG	A087318 001	
	500MG	A089111 001	May 29, 1987

**TOLBUTAMIDE SODIUM**

INJECTABLE;INJECTION

ORINASE DIAGNOSTIC

PHARMACIA AND UPJOHN EQ 1GM BASE/VIAL

N012095 001

**TOLCAPONE**

TABLET;ORAL

TASMAR

VALEANT PHARMS LLC 200MG

N020697 002 Jan 29, 1998

**TOLMETIN SODIUM**

CAPSULE;ORAL

TOLECTIN DS

ORTHO MCNEIL JANSSEN EQ 400MG BASE

N018084 001

TOLMETIN SODIUM

ACTAVIS ELIZABETH	EQ 400MG BASE	A073308 001	Jan 24, 1992
IVAX SUB TEVA PHARMS	EQ 400MG BASE	A073392 001	Jan 24, 1992
SANDOZ	EQ 400MG BASE	A073462 001	Apr 30, 1992
SUN PHARM INDUSTRIES	EQ 400MG BASE	A073311 001	Nov 27, 1991
TEVA	EQ 400MG BASE	A073519 001	May 29, 1992

**DISCONTINUED DRUG PRODUCT LIST**

6-356(of 375)

\*\* See List Footnote

TOLMETIN SODIUM

TABLET;ORAL

TOLECTIN

ORTHO MCNEIL JANSSEN	EQ 200MG BASE	N017628 001
TOLECTIN 600		
ORTHO MCNEIL JANSSEN	EQ 600MG BASE	N017628 002 Mar 08, 1989
TOLMETIN SODIUM		
ACTAVIS ELIZABETH	EQ 600MG BASE	A073527 001 Jun 30, 1992
G AND W LABS INC	EQ 600MG BASE	A074399 001 Mar 28, 1996
	EQ 600MG BASE	A074729 001 Feb 27, 1997
SANDOZ	EQ 200MG BASE	A073588 001 Jul 31, 1992
	EQ 600MG BASE	A074002 001 Sep 27, 1993
SUN PHARM INDUSTRIES	EQ 200MG BASE	A073310 001 Nov 27, 1991

TOLVAPTAN

TABLET;ORAL

SAMSCA

+ OTSUKA AMERICA PHARM	60MG **	N022275 003 May 19, 2009
------------------------	---------	--------------------------

TOPIRAMATE

CAPSULE;ORAL

TOPAMAX SPRINKLE

JANSSEN PHARMS	50MG	N020844 003 Oct 26, 1998
TOPIRAMATE		
BARR	15MG	A076448 001 Apr 15, 2009
	25MG	A076448 002 Apr 15, 2009
MYLAN	15MG	A078418 001 Oct 14, 2009
	25MG	A078418 002 Oct 14, 2009
SANDOZ	15MG	A079206 001 Oct 14, 2009
	25MG	A079206 002 Oct 14, 2009

TABLET;ORAL

TOPAMAX

JANSSEN PHARMS	300MG	N020505 003 Dec 24, 1996
	400MG	N020505 006 Dec 24, 1996

TOPIRAMATE		
ACTAVIS TOTOWA	25MG	A078637 001 Feb 27, 2013
	50MG	A078637 002 Feb 27, 2013
	100MG	A078637 003 Feb 27, 2013
	200MG	A078637 004 Feb 27, 2013
BARR	25MG	A076315 001 Mar 27, 2009
	100MG	A076315 002 Mar 27, 2009
	200MG	A076315 003 Mar 27, 2009
MYLAN	25MG	A076314 001 Mar 27, 2009
	50MG	A076314 002 Mar 27, 2009
	100MG	A076314 003 Mar 27, 2009
	200MG	A076314 004 Mar 27, 2009
PLIVA HRVATSKA DOO	25MG	A077905 001 Mar 30, 2009
	50MG	A077905 002 Mar 30, 2009
	100MG	A077905 003 Mar 30, 2009
	200MG	A077905 004 Mar 30, 2009
ROXANE	25MG	A076306 001 Mar 27, 2009
	50MG	A076306 002 Mar 27, 2009
	100MG	A076306 003 Mar 27, 2009
	200MG	A076306 004 Mar 27, 2009
WATSON LABS	25MG	A077643 001 Mar 27, 2009
	50MG	A077643 002 Mar 27, 2009
	100MG	A077643 003 Mar 27, 2009
	200MG	A077643 004 Mar 27, 2009
WOCKHARDT USA	25MG	A090353 001 Sep 01, 2010
	50MG	A090353 002 Sep 01, 2010
	100MG	A090353 003 Sep 01, 2010
	200MG	A090353 004 Sep 01, 2010

TOPIRAMATE		
HIKMA PHARMS	25MG	A091185 001 Nov 25, 2013
	50MG	A091185 002 Nov 25, 2013
	100MG	A091185 003 Nov 25, 2013
	200MG	A091185 004 Nov 25, 2013

**DISCONTINUED DRUG PRODUCT LIST**

6-357(of 375)

\*\* See List Footnote

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

TOPOTECAN HYDROCHLORIDE

FRESENIUS KABI ONCOL	EQ 4MG BASE/VIAL
SUN PHARM IND LTD	EQ 4MG BASE/VIAL

A091376 001	Nov 29, 2010
A202203 001	Aug 29, 2013

SOLUTION; INTRAVENOUS

TOPOTECAN

+ SANDOZ INC	EQ 1MG BASE/ML (EQ 1MG BASE/ML) **
+ +	EQ 3MG BASE/3ML (EQ 1MG BASE/ML) **
+ +	EQ 4MG BASE/4ML (EQ 1MG BASE/ML) **

N200199 001	Feb 25, 2011
N200199 002	Feb 25, 2011
N200199 003	Feb 25, 2011

TORSEMIDE

INJECTABLE; INJECTION

DEMADEX

+ ROCHE	50MG/5ML (10MG/ML) **
+ +	20MG/2ML (10MG/ML) **

N020137 002	Aug 23, 1993
N020137 001	Aug 23, 1993

TORSEMIDE

LUITPOLD	20MG/2ML (10MG/ML)
	50MG/5ML (10MG/ML)
WEST-WARD PHARMS INT	20MG/2ML (10MG/ML)
	50MG/5ML (10MG/ML)

A090656 001	Apr 21, 2010
A090656 002	Apr 21, 2010
A078007 001	Jun 11, 2008
A078007 002	Jun 11, 2008

TABLET; ORAL

TORSEMIDE

SUN PHARM IND	5MG
	10MG
	20MG
	100MG

A078478 001	Feb 26, 2008
A078478 002	Feb 26, 2008
A078478 003	Feb 26, 2008
A078478 004	Feb 26, 2008

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

ACCORD HLTHCARE	50MG
ACTAVIS ELIZABETH	50MG
ASTA	50MG
IVAX SUB TEVA PHARMS	50MG
MYLAN PHARMS INC	50MG
NORTHSTAR HLTHCARE	50MG
SANDOZ	50MG
WATSON LABS	50MG

A202390 001	May 16, 2013
A075960 001	Jun 19, 2002
A075974 001	Jul 12, 2002
A075963 001	Jul 03, 2002
A075980 001	Nov 21, 2002
A078935 001	May 26, 2010
A075968 001	Jun 25, 2002
A075962 001	Jun 24, 2002

ULTRAM

JANSSEN PHARMS	100MG
----------------	-------

N020281 001	Mar 03, 1995
-------------	--------------

TABLET, EXTENDED RELEASE; ORAL

RYZOLT

+ PURDUE PHARMA	100MG **
+ +	200MG **
+ +	300MG **

N021745 001	Dec 30, 2008
N021745 002	Dec 30, 2008
N021745 003	Dec 30, 2008

ULTRAM ER

+ VALEANT PHARMS	100MG
+ +	200MG
+ +	300MG

N021692 001	Sep 08, 2005
N021692 002	Sep 08, 2005
N021692 003	Sep 08, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

RYBIX ODT

SHIONOGI INC	50MG
--------------	------

N021693 001	May 05, 2005
-------------	--------------

TRAMETINIB DIMETHYL SULFOXIDE

TABLET; ORAL

MEKINIST

+ NOVARTIS PHARMS CORP	EQ 1MG
------------------------	--------

N204114 002	May 29, 2013
-------------	--------------

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

CIPLA	1MG
	2MG
	4MG
DR REDDYS LABS LTD	1MG
	2MG
	4MG
EPIC PHARMA LLC	1MG
	2MG
	4MG

A077307 002	Jun 12, 2007
A077307 001	Jun 12, 2007
A077307 003	Jun 12, 2007
A078493 001	Aug 25, 2008
A078493 002	Aug 25, 2008
A078493 003	Aug 25, 2008
A077256 001	Jun 12, 2007
A077256 002	Jun 12, 2007
A077256 003	Jun 12, 2007

**DISCONTINUED DRUG PRODUCT LIST**

6-358(of 375)

\*\* See List Footnote

**TRANDOLAPRIL**

TABLET;ORAL

TRANDOLAPRIL

INVAGEN PHARMS

1MG

A078320 001 Jun 12, 2007

2MG

A078320 002 Jun 12, 2007

4MG

A078320 003 Jun 12, 2007

MYLAN

1MG

A078346 001 Apr 28, 2008

2MG

A078346 002 Apr 28, 2008

4MG

A078346 003 Apr 28, 2008

**TRANEXAMIC ACID**

TABLET;ORAL

CYKLOKAPRON

PHARMACIA AND UPJOHN 500MG

N019280 001 Dec 30, 1986

TRANEXAMIC ACID

AMERIGEN PHARMS LTD 650MG

A203256 001 Jul 25, 2016

**TRAVOPROST**

SOLUTION/DROPS;OPHTHALMIC

IZBA

+ NOVARTIS PHARMS CORP 0.003% \*\*

N204822 001 May 15, 2014

TRAVATAN

+ ALCON PHARMS LTD 0.004% \*\*

N021257 001 Mar 16, 2001

**TRAZODONE HYDROCHLORIDE**

TABLET;ORAL

DESYREL

+ PRAGMA PHARMS LLC 50MG \*\*

N018207 001

+ 100MG \*\*

N018207 002

+ 150MG \*\*

N018207 003 Mar 25, 1985

+ 300MG \*\*

N018207 004 Nov 07, 1988

TRAZODONE HYDROCHLORIDE

AM THERAP 50MG

A071139 001 Oct 29, 1986

100MG

A071140 001 Oct 29, 1986

AUROLIFE PHARMA LLC 50MG

A072484 001 Apr 30, 1990

MYLAN 50MG

A071405 001 Feb 27, 1991

100MG

A071406 001 Feb 27, 1991

MYLAN PHARMS INC 50MG

A090514 001 Jun 02, 2009

100MG

A090514 002 Jun 02, 2009

150MG

A090514 003 Jun 02, 2009

300MG

A090514 004 Jun 02, 2009

QUANTUM PHARMICS 100MG

A070921 001 Dec 01, 1986

100MG

A072483 001 Apr 30, 1990

TEVA 150MG

A074357 001 Apr 30, 1997

USL PHARMA 50MG

A070491 001 Apr 29, 1987

100MG

A070492 001 Apr 29, 1987

WATSON LABS 50MG

A070857 001 Oct 10, 1986

50MG

A071112 001 Nov 17, 1986

100MG

A070858 001 Oct 10, 1986

100MG

A071113 001 Nov 17, 1986

TRIALODINE

QUANTUM PHARMICS 50MG

A070942 001 Dec 01, 1986

TABLET, EXTENDED RELEASE;ORAL

OLEPTRO

+ ANGELINI PHARMA 150MG \*\*

N022411 001 Feb 02, 2010

+ 300MG \*\*

N022411 002 Feb 02, 2010

**TRETINOIN**

CAPSULE;ORAL

VESANOID

+ CHEPLAPHARM 10MG \*\*

N020438 001 Nov 22, 1995

CREAM;TOPICAL

TRETINOIN

ALLERGAN SALES LLC 0.0375%

A090098 001 Mar 22, 2010

0.075%

A202209 001 Oct 11, 2012

SOLUTION;TOPICAL

RETIN-A

+ VALEANT INTL 0.05%

N016921 001

TRETINOIN

TEVA PHARMS 0.05%

A074873 001 Jun 19, 1998

WOCKHARDT 0.05%

A075260 001 Jan 25, 1999

**DISCONTINUED DRUG PRODUCT LIST**

6-359(of 375)

\*\* See List Footnote

TRETINOINSWAB;TOPICAL  
RETIN-A

VALEANT INTL 0.05% N016921 002

TRIAMCINOLONETABLET;ORAL  
ARISTOCORT

ASTELLAS	1MG	N011161 009
	2MG	N011161 004
	4MG	N011161 007
	8MG	N011161 011
	16MG	N011161 010

KENACORT

DELCOR ASSET CORP	1MG	N011283 003
	2MG	N011283 008
	4MG	N011283 006
	8MG	N011283 010

TRIAMCINOLONE

BARR	2MG	A084286 001
	2MG	A084318 001
	4MG	A084267 001
	4MG	A084319 001
	8MG	A084268 001
	8MG	A084320 001
IMPAX LABS	4MG	A084340 001
IVAX SUB TEVA PHARMS	4MG	A083750 001
MYLAN	2MG	A084406 001
PUREPAC PHARM	2MG	A084020 002
	4MG	A084020 003
ROXANE	2MG	A084708 001
	4MG	A084709 001
	8MG	A084707 001
SANDOZ	4MG	A085601 001
TEVA	4MG	A084775 001
WATSON LABS	4MG	A084270 001
	4MG	A085834 001

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED;INHALATION

AZMACORT

ABBVIE 0.1MG/INH N018117 001 Apr 23, 1982

AEROSOL, METERED;NASAL

NASACORT

SANOFI AVENTIS US 0.055MG/INH N019798 001 Jul 11, 1991

CREAM;TOPICAL

ARISTOCORT

ASTELLAS	0.025%	A083017 003
	0.1%	A083016 004
	0.5%	A083015 002

ARISTOCORT A

ASTELLAS	0.025%	A083017 004
	0.025%	A088818 001 Oct 16, 1984
	0.1%	A083016 005
	0.1%	A088819 001 Oct 16, 1984
	0.5%	A083015 003
	0.5%	A088820 001 Oct 16, 1984

FLUTEX

IVAX PHARMS	0.025%	A085539 001
	0.1%	A085539 002
	0.5%	A085539 003

KENALOG

DELCOR ASSET CORP 0.5% A083943 001

KENALOG-H

DELCOR ASSET CORP 0.1% A086240 001

TRIACET

TEVA	0.025%	A084908 001
	0.1%	A084908 002
	0.5%	A084908 003

**DISCONTINUED DRUG PRODUCT LIST**

6-360(of 375)

\*\* See List Footnote

TRIAMCINOLONE ACETONIDE

CREAM;TOPICAL

TRIACORT

SOLVAY 0.1%

A087113 001

TRIAMCINOLONE ACETONIDE

ACTAVIS MID ATLANTIC 0.1%

A087798 001 Jun 04, 1982

ALPHARMA US PHARMS 0.025%

A087797 001 Jun 07, 1982

AMBITX 0.025%

A087932 001 May 09, 1983

MORTON GROVE 0.025%

A088094 001 Sep 01, 1983

0.1%

A088095 001 Sep 01, 1983

0.5%

A088096 001 Sep 01, 1983

PHARMADERM 0.025%

A087990 001 Jul 07, 1983

0.1%

A087991 001 Jul 07, 1983

0.5%

A087992 001 Jul 07, 1983

PHARMAFAIR 0.025%

A087921 001 Aug 10, 1982

0.1%

A087912 001 Aug 10, 1982

0.5%

A087922 001 Aug 10, 1982

TARO 0.025%

A040038 001 Oct 26, 1994

0.025%

A086277 001

0.1%

A086276 001

0.5%

A086275 001

TOPIDERM 0.025%

A089274 001 Feb 21, 1989

0.1%

A089275 001 Feb 21, 1989

0.5%

A089276 001 Feb 21, 1989

TRIATEX

IVAX PHARMS 0.025%

A087430 001 Nov 01, 1988

0.1%

A087429 001 Nov 01, 1988

0.5%

A087428 001 Nov 01, 1988

TRYMEX

SAVAGE LABS 0.025%

A088196 001 Mar 25, 1983

0.1%

A088197 001 Mar 25, 1983

0.5%

A088198 001 Mar 25, 1983

GEL;TOPICAL

ARISTOGEL

ASTELLAS 0.1%

A083380 001

INJECTABLE;INJECTION

TRIAMCINOLONE ACETONIDE

PARNELL 3MG/ML

N019503 001 Oct 16, 1987

SANDOZ INC 10MG/ML

A090166 001 May 27, 2009

40MG/ML

A090164 001 Jun 01, 2009

WATSON LABS 40MG/ML

A085825 001

INJECTABLE;INTRA-ARTICULAR, INTRAMUSCULAR, INTRAVITREAL

TRIVARIS

+ ALLERGAN 8MG/0.1ML (8MG/0.1ML) \*\*

N022220 001 Jun 16, 2008

LOTION;TOPICAL

KENALOG

DELCOR ASSET CORP 0.025% \*\*

A084343 001

+ 0.025% \*\*

N011602 003

0.1% \*\*

A084343 002

+ 0.1% \*\*

N011602 001

TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS 0.025%

A087191 001 Sep 08, 1982

0.1%

A087192 001 Sep 08, 1982

OINTMENT;TOPICAL

ARISTOCORT

ASTELLAS 0.1%

A080750 004

0.5% \*\*

A080745 002

ARISTOCORT A

ASTELLAS 0.1%

A080750 003

0.1%

A088780 001 Oct 01, 1984

0.5% \*\*

A080745 003

0.5%

A088781 001 Oct 05, 1984

FLUTEX

IVAX PHARMS 0.025%

A087375 001 Nov 01, 1988

0.1%

A087377 001 Nov 01, 1988

0.5%

A087376 001 Nov 01, 1988

KENALOG

DELCOR ASSET CORP 0.5% \*\*

A083944 001

+ MYLAN PHARMS INC 0.025%

N011600 003

**DISCONTINUED DRUG PRODUCT LIST**

6-361(of 375)

\*\* See List Footnote

TRIAMCINOLONE ACETONIDE

OINTMENT;TOPICAL

KENALOG

+ TRIAMCINOLONE ACETONIDE ACTAVIS MID ATLANTIC ALPHARMA US PHARMS MORTON GROVE  PHARMADERM  TARO	0.1% 0.1% 0.025% 0.1% 0.5% 0.025% 0.1% 0.025% 0.1% 0.5%	N011600 001 A087799 001 Jun 07, 1982 A089913 001 Dec 23, 1988 A088090 001 Sep 01, 1983 A088091 001 Sep 01, 1983 A088092 001 Sep 01, 1983 A088692 001 Aug 02, 1984 A088690 001 Aug 02, 1984 A040040 001 Sep 30, 1994 A040374 001 Jun 05, 2001 A087902 001 Dec 27, 1982 A040386 001 Jun 05, 2001
TRYMEX SAVAGE LABS	0.025% 0.1%	A088693 001 Aug 02, 1984 A088691 001 Aug 02, 1984

PASTE;DENTAL

KENALOG IN ORABASE

+ DELCOR ASSET CORP ORALONE TARO	0.1% ** 0.1% 0.1%	N012097 001 A071383 001 Jul 06, 1987
SPRAY, METERED;NASAL ALLERNAZE LUPIN ATLANTIS	0.05MG/SPRAY	N020120 001 Feb 04, 2000
NASACORT HFA SANOFI AVENTIS US	0.055MG/SPRAY	N020784 001 Apr 07, 2004
TRIAMCINOLONE ACETONIDE PERRIGO ISRAEL	0.055MG/SPRAY	A078104 001 Jul 30, 2009

TRIAMCINOLONE DIACETATE

INJECTABLE;INJECTION

ARISTOCORT

FOSUN PHARMA + TRIAMCINOLONE DIACETATE AKORN WATSON LABS	25MG/ML 40MG/ML ** 25MG/ML 40MG/ML 40MG/ML 40MG/ML	N011685 003 N012802 001 A085122 001 A086394 001 A084072 001 A085529 001
SYRUP;ORAL ARISTOCORT ASTELLAS	2MG/5ML	N011960 004
KENACORT DELCOR ASSET CORP	EQ 4MG BASE/5ML	N012515 001

TRIAZOLAM

TABLET;ORAL

HALCION

PHARMACIA AND UPJOHN	0.5MG	N017892 002 Nov 15, 1982
TRIAZOLAM WATSON LABS	0.125MG 0.25MG	A074445 001 Oct 20, 1995 A074445 002 Oct 20, 1995

TRICHLORMETHIAZIDE

TABLET;ORAL

METAHYDRIN

SANOFI AVENTIS US NAQUA SCHERING TRICHLOREX LANNETT TRICHLORMAS MAST MM	2MG 4MG  2MG 4MG  4MG 4MG  4MG 4MG  4MG	N012594 001 Jun 16, 1988 N012594 002 Jun 16, 1988  N012265 001 N012265 002  A083436 001 A085630 001  A086259 001
---	---	---

**DISCONTINUED DRUG PRODUCT LIST**

6-362(of 375)

\*\* See List Footnote

TRICHLORMETHIAZIDE

TABLET;ORAL

## TRICHLORMETHIAZIDE

CHARTWELL RX	4MG	A085568 001
IMPAK LABS	4MG	A083967 001
PAR PHARM	2MG	A087007 001
	4MG	A087005 001
SANDOZ	4MG	A086171 001
WATSON LABS	2MG	A083847 001
	2MG	A086458 001
	4MG	A083462 001
	4MG	A083855 001
	4MG	A085962 001

TRICLOFOS SODIUM

SOLUTION;ORAL

## TRICLOS

SANOFI AVENTIS US	1.5GM/15ML	N016830 001
TABLET;ORAL		

## TRICLOS

SANOFI AVENTIS US	750MG	N016809 002
-------------------	-------	-------------

TRIDIHEXETHYL CHLORIDE

INJECTABLE;INJECTION

## PATHILON

LEDERLE	10MG/ML	N009729 001
TABLET;ORAL		

## PATHILON

LEDERLE	25MG	N009489 005
---------	------	-------------

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE;ORAL

## STELAZINE

+ GLAXOSMITHKLINE	EQ 10MG BASE/ML **	N011552 006
-------------------	--------------------	-------------

## TRIFLUOPERAZINE HYDROCHLORIDE

SANDOZ	EQ 10MG BASE/ML	A085787 001 Apr 15, 1982
WOCKHARDT	EQ 10MG BASE/ML	A088143 001 Jul 26, 1983

INJECTABLE;INJECTION

## STELAZINE

+ GLAXOSMITHKLINE	EQ 2MG BASE/ML **	N011552 005
-------------------	-------------------	-------------

TABLET;ORAL

## STELAZINE

+ GLAXOSMITHKLINE	EQ 1MG BASE **	N011552 001
+	EQ 2MG BASE **	N011552 002
+	EQ 5MG BASE **	N011552 003
+	EQ 10MG BASE **	N011552 004

## TRIFLUOPERAZINE HYDROCHLORIDE

DURAMED PHARMS BARR	EQ 1MG BASE	A088967 001 Apr 23, 1985
	EQ 2MG BASE	A088968 001 Apr 23, 1985
	EQ 5MG BASE	A088969 001 Apr 23, 1985
	EQ 10MG BASE	A088970 001 Apr 23, 1985
IVAX PHARMS	EQ 1MG BASE	A087612 001 Nov 19, 1982
	EQ 2MG BASE	A087613 001 Nov 19, 1982
	EQ 5MG BASE	A087328 001 Nov 19, 1982
	EQ 10MG BASE	A087614 001 Nov 19, 1982
SANDOZ	EQ 1MG BASE	A040153 001 Oct 25, 1996
	EQ 2MG BASE	A040153 002 Oct 25, 1996
	EQ 5MG BASE	A040153 003 Oct 25, 1996
	EQ 10MG BASE	A040153 004 Oct 25, 1996
WATSON LABS	EQ 1MG BASE	A085975 001 Jun 23, 1988
	EQ 2MG BASE	A085976 001 Jun 23, 1988
	EQ 5MG BASE	A085973 001 Jun 23, 1988
	EQ 10MG BASE	A088710 001 Jun 23, 1988

TRIFLUPROMAZINE

SUSPENSION;ORAL

## VESPRIN

APOTHECON	EQ 50MG HCL/5ML	N011491 004
-----------	-----------------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-363(of 375)

\*\* See List Footnote

TRIFLUROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

VESPRIN

APOTHECON	3MG/ML	N011325 005
	10MG/ML	N011325 004
	20MG/ML	N011325 001

TABLET; ORAL

VESPRIN

BRISTOL MYERS SQUIBB	10MG	N011123 001
	25MG	N011123 002
	50MG	N011123 003

TRIHEXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

ARTANE

LEDERLE	5MG	N006773 010
	5MG	N012947 001

ELIXIR; ORAL

ARTANE

LEDERLE	2MG/5ML	N006773 009
---------	---------	-------------

TRIHEXYPHENIDYL HYDROCHLORIDE

PHARM VENTURES	2MG/5ML	A089514 001 Apr 07, 1989
----------------	---------	--------------------------

TABLET; ORAL

ARTANE

+ LEDERLE	2MG **	N006773 005
+ +	5MG **	N006773 003

TREMIN

SCHERING	2MG	A080381 001
	5MG	A080381 003

TRIHEXYPHENIDYL HYDROCHLORIDE

HIKMA PHARMS	2MG	A040337 002 Feb 16, 2000
	5MG	A040337 001 Feb 16, 2000

NYLOS	5MG	A085622 001
-------	-----	-------------

VANGARD	2MG	A088035 001 Jul 30, 1982
---------	-----	--------------------------

WATSON LABS	2MG	A040184 001 Feb 06, 1998
	5MG	A085117 001

	5MG	A040184 002 Feb 06, 1998
	5MG	A085105 001

TRILOSTANE

CAPSULE; ORAL

MODRASTANE

BIOENVISION	30MG	N018719 002 Dec 31, 1984
	60MG	N018719 001 Dec 31, 1984

TRIMEPRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 5MG BASE	N011316 004
------------------	-------------	-------------

SYRUP; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE/5ML	N011316 003
------------------	-------------------	-------------

TRIMEPRAZINE TARTRATE

ALPHARMA US PHARMS	EQ 2.5MG BASE/5ML	A085015 001 Feb 18, 1982
MORTON GROVE	EQ 2.5MG BASE/5ML	A088285 001 Apr 11, 1985

TABLET; ORAL

TEMARIL

ALLERGAN HERBERT	EQ 2.5MG BASE	N011316 001
------------------	---------------	-------------

TRIMETHADIONE

CAPSULE; ORAL

TRIDIONE

ABBVIE	300MG	N005856 005
--------	-------	-------------

SOLUTION; ORAL

TRIDIONE

ABBVIE	200MG/5ML	N005856 002
--------	-----------	-------------

**DISCONTINUED DRUG PRODUCT LIST**

6-364(of 375)

\*\* See List Footnote

TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION

ARFONAD

ROCHE

50MG/ML

N008983 001

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TRIMETHOBENZAMIDE HYDROCHLORIDE

HOSPIRA

100MG/ML

A088804 001 Apr 03, 1987

SMITH AND NEPHEW

100MG/ML

A088960 001 Apr 04, 1986

100MG/ML

A089043 001 Apr 04, 1986

SOLOPAK

100MG/ML

A089094 001 Apr 04, 1986

WATSON LABS

100MG/ML

A086577 001 Oct 19, 1982

100MG/ML

A087939 001 Dec 28, 1982

TRIMETHOPRIM

TABLET; ORAL

PROLOPRIM

MONARCH PHARMS

100MG

N017943 001

200MG

N017943 003 Jul 14, 1982

TRIMETHOPRIM

SUN PHARM INDUSTRIES

100MG

A070494 001 Jan 22, 1986

200MG

A070495 001 Sep 24, 1986

TEVA

200MG \*\*

A071259 001 Jun 18, 1987

TRIMPEX

ROCHE

100MG

N017952 001

TRIMPEX 200

ROCHE

200MG

N017952 002 Nov 09, 1982

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL

PRIMSOL

ALLEGIS

EQ 25MG BASE/5ML

N074374 001 Jun 23, 1995

TRIMETREXATE GLUCURONATE

INJECTABLE; INJECTION

NEUTREXIN

MEDIMMUNE ONCOLOGY

EQ 25MG BASE/VIAL

N020326 001 Dec 17, 1993

EQ 200MG BASE/VIAL

N020326 002 Jul 31, 1998

TRIMIPRAMINE MALEATE

CAPSULE; ORAL

TRIMIPRAMINE MALEATE

USL PHARMA

EQ 25MG BASE

A071283 001 Dec 08, 1987

EQ 50MG BASE

A071284 001 Dec 08, 1987

EQ 100MG BASE

A071285 001 Dec 08, 1987

TRIOXSALEN

TABLET; ORAL

TRISORALEN

VALEANT PHARM INTL

5MG

N012697 001

TRIPELENNAMINE CITRATE

ELIXIR; ORAL

PBZ

NOVARTIS

EQ 25MG HCL/5ML

N005914 004

TRIPELENNAMINE HYDROCHLORIDE

TABLET; ORAL

PBZ

NOVARTIS

25MG

A083149 001

50MG

N005914 002

TRIPELENNAMINE HYDROCHLORIDE

ANABOLIC

50MG

A083037 001

BARR

50MG

A080744 001

HEATHER

50MG

A083989 001

IMPAX LABS

50MG

A080785 001

LANNETT

50MG

A083557 001

NYLOS

50MG

A085412 001

PARKE DAVIS

25MG

A083625 001

50MG

A083626 001

WATSON LABS

50MG

A080713 001

50MG

A080790 001

**DISCONTINUED DRUG PRODUCT LIST**

6-365(of 375)

\*\* See List Footnote

TRIPELENNAMINE HYDROCHLORIDE

TABLET;ORAL

TRIPELENNAMINE HYDROCHLORIDE

50MG

A085188 001

TABLET, EXTENDED RELEASE;ORAL  
PBZ-SR

NOVARTIS

50MG

N010533 002

100MG

N010533 001

TRIPLE SULFA (SULFABENZAMIDE;SULFACETAMIDE;SULFHIAZOLE)

CREAM;VAGINAL

GYNE-SULF

G AND W LABS

3.7%;2.86%;3.42%

A088607 001 Jun 09, 1986

SULTRIN

ORTHO MCNEIL PHARM

3.7%;2.86%;3.42%

N005794 001

TRIPLE SULFA

ALPHARMA US PHARMS

3.7%;2.86%;3.42%

A087864 001 Sep 01, 1982

FOUGERA

3.7%;2.86%;3.42%

A086424 001

PERRIGO NEW YORK

3.7%;2.86%;3.42%

A087285 001 Nov 15, 1982

TRYSL

SAVAGE LABS

3.7%;2.86%;3.42%

A087887 001 Jul 23, 1982

VAGILIA

G AND W LABS INC

3.7%;2.86%;3.42%

A088821 001 Nov 09, 1987

TABLET;VAGINAL

SULTRIN

ORTHO MCNEIL PHARM

184MG;143.75MG;172.5MG

N005794 002

TRIPLE SULFA

FOUGERA

184MG;143.75MG;172.5MG

A088463 001 Jan 03, 1985

PHARMADERM

184MG;143.75MG;172.5MG

A088462 001 Jan 03, 1985

TRIPROLIDINE HYDROCHLORIDE

SYRUP;ORAL

ACTIDIL

GLAXOSMITHKLINE

1.25MG/5ML

N011496 002 Jul 01, 1983

MYIDYL

USL PHARMA

1.25MG/5ML

A087963 001 Jan 18, 1983

TRIPROLIDINE HYDROCHLORIDE

ALPHARMA US PHARMS

1.25MG/5ML

A085940 001

HALSEY

1.25MG/5ML

A088735 001 Jan 17, 1985

PHARM ASSOC

1.25MG/5ML

A087514 001 Feb 10, 1982

TABLET;ORAL

ACTIDIL

GLAXOSMITHKLINE

2.5MG

N011110 002 Jul 01, 1983

TRIPROLIDINE HYDROCHLORIDE

VITARINE

2.5MG

A085610 001

WATSON LABS

2.5MG

A085094 001

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

SUSPENSION;ORAL

LANTRISUL

LANNETT

167MG/5ML;167MG/5ML;167MG/5ML

A080123 002

NEOTRIZINE

LILLY

167MG/5ML;167MG/5ML;167MG/5ML

N006317 012

SULFALOID

FOREST PHARMS

167MG/5ML;167MG/5ML;167MG/5ML

A080100 001

SULFOSE

WYETH AYERST

167MG/5ML;167MG/5ML;167MG/5ML

A080013 002

TERFONYL

BRISTOL MYERS SQUIBB

167MG/5ML;167MG/5ML;167MG/5ML

N006904 002

TRIPLE SULFA

ALPHARMA US PHARMS

167MG/5ML;167MG/5ML;167MG/5ML

A080280 001

TRIPLE SULFAS

LEDERLE

167MG/5ML;167MG/5ML;167MG/5ML

N006920 003

TABLET;ORAL

NEOTRIZINE

LILLY

167MG;167MG;167MG

N006317 011

SULFA-TRIPLE #2

IMPAK LABS

167MG;167MG;167MG

A080079 001

SULFALOID

FOREST PHARMS

167MG;167MG;167MG

A080099 001

**DISCONTINUED DRUG PRODUCT LIST**

6-366(of 375)

\*\* See List Footnote

TRISULFAPYRIMIDINES (SULFDIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

TABLET;ORAL SULFOSE			
WYETH AYERST	167MG;167MG;167MG	A080013	001
TERFONYL			
BRISTOL MYERS SQUIBB	167MG;167MG;167MG	N006904	001
TRIPLE SULFA			
PUREPAC PHARM	167MG;167MG;167MG	A080086	001
TRIPLE SULFAS			
LEDERLE	167MG;167MG;167MG	N006920	002
TRIPLE SULFOID			
PAL PAK	167MG;167MG;167MG	A080094	001

TROGLITAZONE

TABLET;ORAL PRELAY			
SANKYO	200MG	N020719	001 Jan 29, 1997
	300MG	N020719	003 Aug 04, 1997
	400MG	N020719	002 Jan 29, 1997
REZULIN			
PFIZER PHARMS	200MG	N020720	001 Jan 29, 1997
	300MG	N020720	003 Aug 04, 1997
	400MG	N020720	002 Jan 29, 1997

TROLAMINE POLYPEPTIDE OLEATE CONDENSATE

SOLUTION/DROPS;OTIC CERUMENEX			
PHARM RES ASSOC	10%	N011340	002

TROLEandomycin

CAPSULE;ORAL TAO			
PFIZER	EQ 250MG BASE	N050336	002
SUSPENSION;ORAL TAO			
PFIZER	EQ 125MG BASE/5ML	N050332	001

TROPICAMIDE

SOLUTION/DROPS;OPHTHALMIC MYDRIACYL			
ALCON	0.5% **	N012111	002
	1% **	N012111	004
MYDRIAFAIR			
PHARMAFAIR	0.5%	A088274	001 Sep 16, 1983
	1%	A088230	001 Sep 16, 1983
TROPICAMIDE			
AKORN	1%	A088447	001 Aug 28, 1985
ALCON PHARMS LTD	1%	A089172	001 Dec 28, 1990
MIZA PHARMS USA	0.5%	A087636	001 Jul 30, 1982
	1%	A087637	001 Aug 09, 1982
WATSON LABS	0.5%	A089171	001 Dec 28, 1990

TROSPiUM CHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL SANCTURA XR			
+ ALLERGAN	60MG **	N022103	001 Aug 03, 2007
TROSPiUM CHLORIDE UPSHER-SMITH LABS	60MG	A091635	001 Apr 29, 2015
TABLET;ORAL SANCTURA			
+ ALLERGAN	20MG **	N021595	001 May 28, 2004

TROVAFLOXACIN MESYLATE

TABLET;ORAL TROVAN			
PFIZER	EQ 100MG BASE	N020759	001 Dec 18, 1997
	EQ 200MG BASE	N020759	002 Dec 18, 1997

**DISCONTINUED DRUG PRODUCT LIST**

6-367(of 375)

\*\* See List Footnote

TUBOCURARINE CHLORIDE

INJECTABLE; INJECTION

TUBOCURARINE CHLORIDE

BRISTOL MYERS SQUIBB	3MG/ML	N005657 001
HOSPIRA	3MG/ML	N006095 001
LILLY	3MG/ML	N006325 001

TYROPOANOATE SODIUM

CAPSULE; ORAL

BILOPAQUE

GE HEALTHCARE	750MG	N013731 001
---------------	-------	-------------

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

+ SUCAMPO PHARMA LLC	0.15% **	N021214 001 Aug 03, 2000
----------------------	----------	--------------------------

URACIL MUSTARD

CAPSULE; ORAL

URACIL MUSTARD

SHIRE	1MG	N012892 001
-------	-----	-------------

UREA

INJECTABLE; INJECTION

STERILE UREA

HOSPIRA	40GM/VIAL	N017698 001
---------	-----------	-------------

UREAPHIL

HOSPIRA	40GM/VIAL	N012154 001
---------	-----------	-------------

UREA C-13

FOR SOLUTION; ORAL

BREATHTEK UBT FOR H-PYLORI

OTSUKA AMERICA	EQ 75MG/POUCH	N020586 002 May 10, 2001
----------------	---------------	--------------------------

HELICOSOL

METABOLIC SOLUTIONS	125MG/VIAL	N021092 001 Dec 17, 1999
---------------------	------------	--------------------------

MERETEK UBT KIT (W/ PRANACTIN)

OTSUKA AMERICA	125MG/VIAL	N020586 001 Sep 17, 1996
----------------	------------	--------------------------

PYLORI-CHEK BREATH TEST

DXS DEVICES	100MG/VIAL	N020900 001 Feb 04, 1999
-------------	------------	--------------------------

UROFOLLITROPIN

INJECTABLE; INTRAMUSCULAR

METRODIN

SERONO	75 IU/AMP	N019415 002 Sep 18, 1986
	150 IU/AMP	N019415 003 Sep 18, 1986

INJECTABLE; SUBCUTANEOUS

FERTINEX

SERONO	75 IU/AMP	N019415 005 Aug 23, 1996
	150 IU/AMP	N019415 004 Aug 23, 1996

UROKINASE

INJECTABLE; INJECTION

KINLYTIC

MICROBIX BIOSYSTEMS	5,000 IU/VIAL	N021846 003
	9,000 IU/VIAL	N021846 002
	250,000 IU/VIAL	N021846 001

URSODIOL

CAPSULE; ORAL

ACTIGALL

ALLERGAN SALES LLC	150MG	N019594 001 Dec 31, 1987
--------------------	-------	--------------------------

URSODIOL

IMPAX LABS INC	300MG	A077895 001 Jul 27, 2006
----------------	-------	--------------------------

TABLET; ORAL

URSODIOL

TEVA PHARMS USA	250MG	A079184 001 May 13, 2009
	500MG	A079184 002 May 13, 2009

**DISCONTINUED DRUG PRODUCT LIST**

6-368(of 375)

\*\* See List Footnote

VALACYCLOVIR HYDROCHLORIDE

TABLET;ORAL

VALACYCLOVIR HYDROCHLORIDE

MYLAN

EQ 500MG BASE

A078070 001 May 24, 2010

EQ 1GM BASE

A078070 002 May 24, 2010

VALDECOXIB

TABLET;ORAL

BEXTRA

GD SEARLE

10MG

N021341 002 Nov 16, 2001

20MG

N021341 003 Nov 16, 2001

VALPROIC ACID

CAPSULE;ORAL

VALPROIC ACID

PAR PHARM

250MG

A070431 001 Feb 28, 1986

SCHERER RP

250MG

A070195 001 Jul 02, 1987

USL PHARMA

250MG

A070631 001 Jun 11, 1987

CAPSULE, DELAYED RELEASE;ORAL

STAVZOR

+ BIONPHARMA INC

125MG \*\*

N022152 001 Jul 29, 2008

+

250MG \*\*

N022152 002 Jul 29, 2008

+

500MG \*\*

N022152 003 Jul 29, 2008

SYRUP;ORAL

VALPROIC ACID

APOTEX INC

250MG/5ML

A077105 001 Jul 29, 2005

VALSARTAN

CAPSULE;ORAL

DIOVAN

NOVARTIS

80MG

N020665 001 Dec 23, 1996

160MG

N020665 002 Dec 23, 1996

VANCOMYCIN HYDROCHLORIDE

FOR SOLUTION;ORAL

VANCOCIN HYDROCHLORIDE

ANI PHARMS INC

EQ 250MG BASE/5ML

A061667 002 Jul 13, 1983

EQ 500MG BASE/6ML

A061667 001

VANCOLED

LEDERLE

EQ 250MG BASE/5ML

A063321 002 Oct 15, 1993

EQ 500MG BASE/6ML

A063321 003 Oct 15, 1993

INJECTABLE;INJECTION

VANCOCIN HYDROCHLORIDE

ANI PHARMS INC

EQ 500MG BASE/VIAL \*\*

A060180 001

EQ 500MG BASE/VIAL

A062476 001 Mar 15, 1984

EQ 500MG BASE/VIAL

A062716 001 Mar 13, 1987

EQ 500MG BASE/VIAL \*\*

A062812 001 Nov 17, 1987

EQ 1GM BASE/VIAL \*\*

A060180 002 Mar 21, 1986

EQ 1GM BASE/VIAL

A062476 002 Mar 21, 1986

EQ 1GM BASE/VIAL

A062716 002 Mar 13, 1987

EQ 1GM BASE/VIAL \*\*

A062812 002 Nov 17, 1987

EQ 10GM BASE/VIAL \*\*

A062812 003 Nov 17, 1987

VANCOLED

WEST-WARD PHARMS INT

EQ 500MG BASE/VIAL \*\*

A062682 001 Jul 22, 1986

EQ 1GM BASE/VIAL \*\*

A062682 002 Mar 30, 1988

EQ 2GM BASE/VIAL \*\*

A062682 003 May 11, 1988

EQ 5GM BASE/VIAL \*\*

A062682 004 May 11, 1988

EQ 10GM BASE/VIAL \*\*

A062682 005 May 11, 1988

VANCOMYCIN HYDROCHLORIDE

TEVA PHARMS USA

EQ 1GM BASE/VIAL

A201251 002 Dec 23, 2015

EQ 5GM BASE/VIAL

A201250 001 Dec 23, 2015

EQ 10GM BASE/VIAL

A201250 002 Dec 23, 2015

EQ 500MG BASE/VIAL

A201251 001 Dec 23, 2015

WEST-WARD PHARMS INT

EQ 500MG BASE/VIAL

A062879 001 Aug 02, 1988

EQ 1GM BASE/VIAL

A062879 002 Aug 02, 1988

VANCOR

PHARMACIA AND UPJOHN

EQ 500MG BASE/VIAL

A062956 001 Aug 01, 1988

EQ 1GM BASE/VIAL

A062956 002 Aug 01, 1988

**DISCONTINUED DRUG PRODUCT LIST**

6-369(of 375)

\*\* See List Footnote

VARDENAFIL HYDROCHLORIDE

TABLET;ORAL

VARDENAFIL HYDROCHLORIDE

TEVA PHARMS	2.5MG	A091347 001	May 03, 2012
	5MG	A091347 002	May 03, 2012
	10MG	A091347 003	May 03, 2012
	20MG	A091347 004	May 03, 2012

VASOPRESSIN TANNATE

INJECTABLE;INJECTION

PITRESSIN TANNATE

+ PARKE DAVIS

5PRESSOR UNITS/ML \*\*

N003402 001

VECURONIUM BROMIDE

INJECTABLE;INJECTION

NORCURON

+ ORGANON USA INC

10MG/VIAL \*\*

N018776 002 Apr 30, 1984

+

20MG/VIAL \*\*

N018776 003 Jan 03, 1992

VECURONIUM BROMIDE

HOSPIRA

4MG/VIAL

A075558 001 Sep 11, 2001

WATSON LABS

10MG/VIAL

A074334 001 Aug 31, 1995

20MG/VIAL

A074334 002 Aug 31, 1995

WEST-WARD PHARMS INT

10MG/VIAL

A075218 001 Aug 23, 1999

20MG/VIAL

A075218 002 Aug 23, 1999

VELAGLUCERASE ALFA

POWDER;IV (INFUSION)

VPRIV

SHIRE HUMAN GENETIC 200 UNITS/VIAL

N022575 002 Feb 26, 2010

VENLAFAKINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE;ORAL

EFFEXOR XR

WYETH PHARMS INC

EQ 100MG BASE

N020699 003 Oct 20, 1997

VENLAFAKINE HYDROCHLORIDE

MYLAN

EQ 37.5MG BASE

A078789 001 Jun 01, 2011

EQ 75MG BASE

A078789 002 Jun 01, 2011

EQ 150MG BASE

A078789 003 Jun 01, 2011

TABLET;ORAL

EFFEXOR

+ WYETH PHARMS INC

EQ 12.5MG BASE \*\*

N020151 001 Dec 28, 1993

+

EQ 25MG BASE \*\*

N020151 002 Dec 28, 1993

+

EQ 37.5MG BASE \*\*

N020151 006 Dec 28, 1993

+

EQ 50MG BASE \*\*

N020151 003 Dec 28, 1993

+

EQ 75MG BASE \*\*

N020151 004 Dec 28, 1993

+

EQ 100MG BASE \*\*

N020151 005 Dec 28, 1993

VENLAFAKINE HYDROCHLORIDE

PLIVA HRVATSKA DOO

EQ 25MG BASE

A078517 001 Jun 13, 2008

EQ 37.5MG BASE

A078517 002 Jun 13, 2008

EQ 50MG BASE

A078517 003 Jun 13, 2008

EQ 75MG BASE

A078517 004 Jun 13, 2008

EQ 100MG BASE

A078517 005 Jun 13, 2008

SANDOZ

EQ 25MG BASE

A077515 001 Jun 13, 2008

EQ 37.5MG BASE

A077515 002 Jun 13, 2008

EQ 50MG BASE

A077515 003 Jun 13, 2008

EQ 75MG BASE

A077515 004 Jun 13, 2008

EQ 100MG BASE

A077515 005 Jun 13, 2008

VERAPAMIL HYDROCHLORIDE

INJECTABLE;INJECTION

CALAN

GD SEARLE LLC

2.5MG/ML

N019038 001 Mar 30, 1984

ISOPTIN

+ MT ADAMS

2.5MG/ML \*\*

N018485 001

VERAPAMIL HYDROCHLORIDE

ABRAXIS PHARM

2.5MG/ML

A070348 001 May 01, 1986

BEDFORD

2.5MG/ML

A072888 001 Jul 28, 1995

HOSPIRA

2.5MG/ML

A070577 001 Feb 02, 1987

2.5MG/ML

A070739 001 May 06, 1987

2.5MG/ML

A070740 001 May 06, 1987

INTL MEDICATION

2.5MG/ML

A070451 001 Dec 16, 1985

LUITPOLD

2.5MG/ML

A070225 001 Nov 12, 1985

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-370(of 375)

\*\* See List Footnote

VERAPAMIL HYDROCHLORIDE

INJECTABLE; INJECTION

VERAPAMIL HYDROCHLORIDE

MARSAM PHARMS LLC	2.5MG/ML 2.5MG/ML 2.5MG/ML	A070617 001 Nov 12, 1985 A072233 001 Feb 26, 1993 A073485 001 Sep 27, 1993
SMITH AND NEPHEW	2.5MG/ML 2.5MG/ML	A070696 001 Jul 31, 1987 A070697 001 Jul 31, 1987
SOLOPAK	2.5MG/ML	A070695 001 Jul 31, 1987
<b>TABLET; ORAL</b>		
CALAN		
GD SEARLE LLC	40MG 160MG	N018817 003 Feb 23, 1988 N018817 004 Feb 23, 1988
ISOPTIN		
MT ADAMS	40MG 80MG 120MG	N018593 003 Nov 23, 1987 N018593 001 Mar 08, 1982 N018593 002 Mar 08, 1982
VERAPAMIL HYDROCHLORIDE		
ACTAVIS ELIZABETH	80MG 120MG	A071019 001 Sep 24, 1986 A070468 001 Sep 24, 1986
FOSUN PHARMA	40MG 80MG 120MG	A073168 001 Jul 31, 1992 A071423 001 May 24, 1988 A071424 001 May 25, 1988
MUTUAL PHARM	80MG 120MG	A070482 001 Sep 24, 1986 A070483 001 Sep 24, 1986
PLIVA	40MG 80MG 120MG	A072751 001 Feb 23, 1996 A072124 001 Jan 26, 1989 A072125 001 Jan 26, 1989
SUN PHARM INDUSTRIES	80MG 120MG	A071489 002 Jan 13, 1988 A071489 001 Jan 13, 1988
WARNER CHILCOTT	80MG 120MG	A070340 001 Sep 24, 1986 A070341 001 Sep 24, 1986
WATSON LABS	40MG 80MG 80MG 120MG 120MG	A072799 001 Apr 28, 1989 A072923 001 Jun 29, 1993 A070855 001 Sep 24, 1986 A071366 001 Oct 01, 1986 A070856 001 Sep 24, 1986
TABLET, EXTENDED RELEASE; ORAL		
CALAN SR		
+ PFIZER	180MG **	N019152 002 Dec 15, 1989
COVERA-HS		
GD SEARLE LLC	180MG 240MG	N020552 001 Feb 26, 1996 N020552 002 Feb 26, 1996
VERAPAMIL HYDROCHLORIDE		
PLIVA	240MG	A072922 001 Mar 01, 1996

VERATRUM VIRIDE ROOT

TABLET; ORAL

VERTAVIS

MEDPOINTE PHARM HLC 130CSR UNIT

N005691 002

VIDARABINE

INJECTABLE; INJECTION

VIRA-A

PARKEDALE EQ 187.4MG BASE/ML

N050523 001

OINTMENT; OPHTHALMIC

VIRA-A

PARKEDALE 3%

N050486 001

VINBLASTINE SULFATE

INJECTABLE; INJECTION

VELBAN

LILLY 10MG/VIAL

N012665 001

VINBLASTINE SULFATE

ABRAXIS PHARM 10MG/VIAL

A089011 001 Nov 18, 1985

HOSPIRA 10MG/VIAL

A089565 001 Aug 18, 1987

**DISCONTINUED DRUG PRODUCT LIST**

6-371(of 375)

\*\* See List Footnote

VINCRISTINE SULFATE

INJECTABLE; INJECTION

ONCOVIN

LILLY	1MG/VIAL 1MG/ML 5MG/VIAL	N014103 001 N014103 003 Mar 07, 1984 N014103 002
VINCASAR PFS TEVA PARENTERAL	1MG/ML	A071426 001 Jul 17, 1987
VINCREX BRISTOL MYERS SQUIBB	5MG/VIAL	A070867 001 Jul 12, 1988
VINCRISTINE SULFATE ABIC	1MG/ML	A070873 001 Feb 19, 1987
ABRAXIS PHARM	1MG/ML	A070411 001 Sep 10, 1986
FRESENIUS KABI USA	1MG/ML 1MG/ML	A076296 001 Dec 20, 2002 A076401 001 Oct 28, 2003
HOSPIRA	1MG/VIAL 2MG/VIAL 5MG/VIAL	A071559 001 Apr 11, 1988 A071560 001 Apr 11, 1988 A071561 001 Apr 11, 1988

VINORELBINE TARTRATE

INJECTABLE; INJECTION

VINORELBINE TARTRATE

EBWE PHARMA	EQ 10MG BASE/ML	A078408 001 Feb 13, 2008
MYLAN LABS LTD	EQ 10MG BASE/ML	A200148 001 Aug 31, 2012

VIOMYCIN SULFATE

INJECTABLE; INJECTION

VIOCIN SULFATE

PFIZER	EQ 1GM BASE/VIAL EQ 5GM BASE/VIAL	A061086 001 A061086 002
--------	--------------------------------------	----------------------------

VITAMIN A

CAPSULE; ORAL

AQUASOL A

ASTRAZENECA	25,000USP UNITS 50,000USP UNITS	A083080 002 A083080 001
VITAMIN A		
BANNER PHARMACAPS	50,000USP UNITS	A083973 001
CHASE CHEM	50,000 IU	A083351 001
EVERYLIFE	50,000 IU	A083134 001
IMPAX LABS	50,000USP UNITS	A080952 001
WEST WARD	50,000USP UNITS	A080985 001

VITAMIN A PALMITATE

CAPSULE; ORAL

AFAXIN

STERLING WINTHROP	EQ 50,000 UNITS BASE	A083187 001
-------------------	----------------------	-------------

ALPHALIN

LILLY	EQ 50,000 UNITS BASE	A080883 001
-------	----------------------	-------------

DEL-VI-A

DEL RAY LABS	EQ 50,000 UNITS BASE	A080830 001
--------------	----------------------	-------------

VI-DOM-A

BAYER PHARMS	EQ 50,000 UNITS BASE	A080972 001
--------------	----------------------	-------------

VITAMIN A

BANNER PHARMACAPS	EQ 50,000 UNITS BASE	A080702 001
BRISTOL MYERS SQUIBB	EQ 50,000 UNITS BASE	A080860 001
CHASE CHEM	EQ 50,000 UNITS BASE	A080746 001
	EQ 50,000 UNITS BASE	A083207 001
ELKINS SINK	EQ 50,000 UNITS BASE	A085479 001
EVERYLIFE	EQ 50,000 UNITS BASE	A080943 001
	EQ 50,000 UNITS BASE	A083114 001
IMPAX LABS	EQ 50,000 UNITS BASE	A080953 001
	EQ 50,000 UNITS BASE	A080955 001
IVAX SUB TEVA PHARMS	EQ 50,000 UNITS BASE	A083035 001
	EQ 50,000 UNITS BASE	A083190 001
MK LABS	EQ 25,000 UNITS BASE	A083457 002
	EQ 50,000 UNITS BASE	A083457 001
WEST WARD	EQ 50,000 UNITS BASE	A080967 001
WHARTON LABS	EQ 50,000 UNITS BASE	A083665 001
VITAMIN A PALMITATE		
ARCUM	EQ 50,000 UNITS BASE	A083311 001

**DISCONTINUED DRUG PRODUCT LIST**

6-372(of 375)

\*\* See List Footnote

VITAMIN A PALMITATE

CAPSULE;ORAL

VITAMIN A PALMITATE

BANNER PHARMACAPS	EQ 50,000 UNITS BASE	A083321 001
	EQ 50,000 UNITS BASE	A083948 001
	EQ 50,000 UNITS BASE	A083981 001
VITAMIN A SOLUBILIZED TEVA	EQ 50,000 UNITS BASE	A080921 001
INJECTABLE;INJECTION		
VITAMIN A PALMITATE BEL MAR	EQ 50,000 UNITS BASE/ML	A080819 001

VORTioxetine Hydrobromide

TABLET;ORAL

TRINTELLIX

+ TAKEDA PHARMS USA EQ 15MG BASE \*\* N204447 003 Sep 30, 2013

Warfarin Potassium

TABLET;ORAL

ATHROMBIN-K

PHARM RES ASSOC	2MG	N011771 007
	5MG	N011771 004
	10MG	N011771 005
	25MG	N011771 006

Warfarin Sodium

INJECTABLE;INJECTION

COUMADIN

BRISTOL MYERS SQUIBB	5MG/VIAL	N009218 024 Feb 07, 1995
	50MG/VIAL	N009218 020
	75MG/VIAL	N009218 012

TABLET;ORAL

ATHROMBIN

PHARM RES ASSOC	5MG	N011771 003
	10MG	N011771 002
	25MG	N011771 001

PANWARFIN

ABBOTT	2MG	N017020 001
	2.5MG	N017020 002
	5MG	N017020 003
	7.5MG	N017020 004
	10MG	N017020 005

WARFARIN SODIUM

FOSUN PHARMA	1MG	A040196 001 Sep 30, 1997
	2MG	A040196 002 Sep 30, 1997
	2.5MG	A040196 003 Sep 30, 1997
	3MG	A040196 008 Jul 26, 2000
	4MG	A040196 004 Sep 30, 1997
	5MG	A040196 005 Sep 30, 1997
	6MG	A040196 009 Jul 26, 2000
	7.5MG	A040196 006 Sep 30, 1997
	10MG	A040196 007 Sep 30, 1997

MYLAN	1MG	A040415 001 Sep 27, 2004
	2MG	A040415 002 Sep 27, 2004
	2.5MG	A040415 003 Sep 29, 2004
	3MG	A040415 004 Sep 27, 2004
	4MG	A040415 005 Sep 27, 2004
	5MG	A040415 006 Sep 27, 2004
	6MG	A040415 007 Sep 27, 2004
	7.5MG	A040415 008 Sep 27, 2004
	10MG	A040415 009 Sep 27, 2004

USL PHARMA	2MG	A088719 001 Jun 27, 1985
	2.5MG	A088720 001 Aug 06, 1985
	5MG	A088721 001 Jul 02, 1985

WATSON LABS	2MG	A086123 001 Aug 17, 1982
	2.5MG	A086120 001 Aug 17, 1982
	5MG	A086119 001 Aug 17, 1982
	7.5MG	A086118 001 Aug 17, 1982
	10MG	A086122 001 Aug 17, 1982

**DISCONTINUED DRUG PRODUCT LIST**

6-373(of 375)

\*\* See List Footnote

XENON XE-127

GAS; INHALATION

XENON XE 127

MALLINCKRODT

5mCi/VIAL

10mCi/VIAL

N018536 001 Oct 01, 1982

N018536 002 Oct 01, 1982

XENON XE-133

GAS; INHALATION

XENON XE 133

GE HEALTHCARE

1 CI/AMP

10mCi/VIAL

20mCi/VIAL

N017256 002

N017687 002

N017687 003

GEN ELECTRIC

5-100 CI/CYLINDER

0.25-5 CI/AMP

N017550 001

N017550 003

XENON XE 133-V.S.S.

GE HEALTHCARE

10mCi/VIAL

N017687 001

INJECTABLE; INJECTION

XENON XE 133

GE HEALTHCARE

1.3-1.7 CI/AMP

LANTHEUS MEDCL

6.3mCi/ML

N017256 001

N017283 001

SOLUTION; INHALATION, INJECTION

XENEISOL

MALLINCKRODT

18-25mCi/AMP

N017262 002

XYLOSE

POWDER; ORAL

XYLO-PFAN

SAVAGE LABS

25GM/BOT

N017605 001

XYLOSE

LYNE

25GM/BOT

N018856 001 Mar 26, 1987

ZALCITABINE

TABLET; ORAL

HIVID

ROCHE

0.375MG

N020199 001 Jun 19, 1992

0.75MG

N020199 002 Jun 19, 1992

ZALEPLON

CAPSULE; ORAL

ZALEPLON

MYLAN

5MG

A077238 001 Jun 06, 2008

10MG

A077238 002 Jun 06, 2008

UPSHER-SMITH LABS

5MG

A078095 001 Jun 06, 2008

5MG

A078706 001 Jun 06, 2008

10MG

A078095 002 Jun 06, 2008

10MG

A078706 002 Jun 06, 2008

ZICONOTIDE ACETATE

INJECTABLE; INTRATHECAL

PRIALT

JAZZ PHARMS INTL

200MCG/2ML (100MCG/ML)

N021060 003 Dec 28, 2004

ZIDOVUDINE

INJECTABLE; INJECTION

ZIDOVUDINE

LIAONING CHENGDA

10MG/ML

A204538 001 Nov 26, 2013

TABLET; ORAL

RETROVIR

VIIV HLTHCARE

200MG

N020518 001 Dec 19, 1995

+

300MG \*\*

N020518 002 Oct 04, 1996

ZIDOVUDINE

AUROBINDO PHARMA

60MG

N022294 001 Jul 23, 2009

HEC PHARM USA INC

300MG

A202058 001 Oct 07, 2011

MATRIX LABS LTD

100MG

N200732 001 Feb 23, 2011

RANBAXY LABS LTD

300MG

A077327 001 Sep 19, 2005

ZILEUTON

TABLET; ORAL

ZYFLO

CHIESI USA INC

300MG

N020471 001 Dec 09, 1996

**DISCONTINUED DRUG PRODUCT LIST**

6-374(of 375)

\*\* See List Footnote

ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

ABRAXIS PHARM

EQ 1MG ZINC/ML

N019229 002 May 05, 1987

ZIPRASIDONE HYDROCHLORIDE

SUSPENSION; ORAL

GEODON

PFIZER INC

EQ 10MG BASE/ML

N021483 001 Mar 29, 2006

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

ZOLEDRONIC ACID

SUN PHARMA GLOBAL

EQ 4MG BASE/5ML

A202746 001 Mar 04, 2013

ZOMETA

+ NOVARTIS

EQ 4MG BASE/VIAL \*\*

N021223 001 Aug 20, 2001

ZOLMITRIPTAN

TABLET; ORAL

ZOLMITRIPTAN

SUN PHARMA GLOBAL

2.5MG

A203476 001 Nov 13, 2014

5MG

A203476 002 Nov 13, 2014

ZOLPIDEM TARTRATE

TABLET; ORAL

ZOLPIDEM TARTRATE

DR REDDYS LABS LTD

5MG

A077985 001 Apr 23, 2007

10MG

A077985 002 Apr 23, 2007

HIKMA

5MG

A078129 001 Apr 30, 2008

10MG

A078129 002 Apr 30, 2008

MYLAN PHARMS INC

5MG

A078016 001 Apr 23, 2007

10MG

A078016 002 Apr 23, 2007

SUN PHARM INDUSTRIES

5MG

A077288 001 Apr 23, 2007

10MG

A077288 002 Apr 23, 2007

SYNTTHON PHARMS

5MG

A077540 001 Apr 23, 2007

10MG

A077540 002 Apr 23, 2007

VIVIMED GLOBAL

5MG

A076062 001 Apr 23, 2007

10MG

A076062 002 Apr 23, 2007

WATSON LABS

5MG

A077773 001 Apr 23, 2007

10MG

A077773 002 Apr 23, 2007

TABLET, EXTENDED RELEASE; ORAL

ZOLPIDEM TARTRATE

SYNTTHON PHARMS

6.25MG

A078483 001 Apr 12, 2011

12.5MG

A078483 002 Jun 06, 2011

TABLET, ORALLY DISINTEGRATING; ORAL

TOVALT ODT

BIOVAIL LABS INTL

5MG

N021412 001 Apr 25, 2007

10MG

N021412 002 Apr 25, 2007

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

ANI PHARMS INC

25MG

A077639 001 Dec 22, 2005

25MG

A077641 003 Dec 22, 2005

50MG

A077639 002 Dec 22, 2005

50MG

A077641 002 Dec 22, 2005

100MG

A077639 003 Dec 22, 2005

100MG

A077641 001 Dec 22, 2005

DR REDDYS LABS LTD

25MG

A077645 002 Sep 29, 2006

50MG

A077645 003 Sep 29, 2006

100MG

A077645 001 Dec 22, 2005

EPIC PHARMA LLC

25MG

A077876 001 Feb 21, 2007

50MG

A077876 002 Feb 21, 2007

100MG

A077876 003 Feb 21, 2007

MYLAN PHARMS INC

25MG

A077647 001 Dec 22, 2005

50MG

A077647 002 Dec 22, 2005

100MG

A077647 003 Dec 22, 2005

ROXANE

25MG

A077648 001 Dec 22, 2005

50MG

A077648 002 Dec 22, 2005

100MG

A077648 003 Dec 22, 2005

SUN PHARM INDUSTRIES

25MG

A077635 001 Dec 22, 2005

50MG

A077635 002 Dec 22, 2005

\*\*Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons\*\*

**DISCONTINUED DRUG PRODUCT LIST**

6-375(of 375)

\*\* See List Footnote

**ZONISAMIDE**CAPSULE;ORAL  
ZONISAMIDE

	100MG	A077635 003 Dec 22, 2005
UPSHER-SMITH LABS	25MG	A077644 001 Dec 22, 2005
	50MG	A077644 002 Dec 22, 2005
	100MG	A077644 003 Dec 22, 2005
WATSON LABS	25MG	A077650 001 Apr 20, 2006
	50MG	A077650 002 Apr 20, 2006
	100MG	A077650 003 Apr 20, 2006



**ORPHAN PRODUCT DESIGNATIONS AND APPROVALS LIST**

The list of Orphan Designations and Approvals is available at:  
<http://www.fda.gov/ForIndustry/DevelopingProductsforRareDiseasesConditions/default.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE IN VIVO BIOAVAILABILITY  
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

ACETAMINOPHEN;ASPIRIN;BUTALBITAL  
CAPSULE OR TABLET; ORAL  
160-165MG;160-165MG;50MG  
325MG;325MG;50MG

ASPIRIN;CAFFEINE;CARISOPRODOL;  
CODEINE PHOSPHATE  
TABLET; ORAL  
160MG;32MG;200MG;16MG

ACETAMINOPHEN;ASPIRIN;BUTALBITAL;  
CAFFEINE  
CAPSULE OR TABLET; ORAL  
160-165MG;160-165MG;50MG;40MG  
325MG;325MG;50MG;40MG

ASPIRIN;CARISOPRODOL  
TABLET; ORAL  
325MG;200MG

ACETAMINOPHEN;BUTALBITAL  
CAPSULE OR TABLET; ORAL  
325MG;50MG

ASPIRIN;CARISOPRODOL;  
CODEINE PHOSPHATE  
TABLET; ORAL  
325MG;200MG;16MG

ACETAMINOPHEN;BUTALBITAL;CAFFEINE  
CAPSULE OR TABLET; ORAL  
325MG;50MG;40MG

ASPIRIN;MEPROBAMATE  
TABLET; ORAL  
325MG;200MG

AMINOPHYLLINE  
TABLET; ORAL  
100MG;200MG

ASPIRIN;METHOCARBAMOL  
TABLET; ORAL  
325MG;400MG

ASPIRIN;BUTALBITAL  
CAPSULE OR TABLET; ORAL  
325MG;50MG  
650MG;50MG

CHLOROTHIAZIDE  
TABLET; ORAL  
250MG

ASPIRIN;BUTALBITAL;CAFFEINE  
CAPSULE OR TABLET; ORAL  
325MG;50MG;40MG  
650MG;50MG;40MG

HYDROXYZINE HYDROCHLORIDE  
TABLET; ORAL  
10MG;25MG;  
50MG;100MG

ASPIRIN;CAFFEINE;CARISOPRODOL  
TABLET; ORAL  
160MG;32MG;200MG

PREDNISONE  
TABLET; ORAL  
1MG;2.5MG;5MG;10MG;  
20MG;25MG;50MG

**APPENDIX A - PRODUCT NAME INDEX**

\*\* A \*\*

A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE  
ABACAVIR SULFATE, ABACAVIR SULFATE  
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE  
ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE  
ABELCET, AMPHOTERICIN B  
ABILITY, ARIPIPRAZOLE  
ABILITY MAINTENA KIT, ARIPIPRAZOLE  
ABILITY MYCITE KIT, ARIPIPRAZOLE  
ABRAXANE, PACLITAXEL  
ABREVA, DOCOSANOL (OTC)  
ABSORICA, ISOTRETINOIN  
ABSTRAL, FENTANYL CITRATE  
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM  
ACANYA, BENZOYL PEROXIDE  
ACARBOSE, ACARBOSE  
ACCOLATE, ZAFIRLUKAST  
ACCUNEB, ALBUTEROL SULFATE  
ACCUPRIL, QUINAPRIL HYDROCHLORIDE  
ACCURETIC, HYDROCHLOROTHIAZIDE  
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE  
ACEPHEN, ACETAMINOPHEN (OTC)  
ACETADOTE, ACETYLCYSTEINE  
ACETAMINOPHEN, ACETAMINOPHEN (OTC)  
ACETAMINOPHEN, ACETAMINOPHEN  
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN  
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)  
ACETAMINOPHEN, CAFFEINE AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN  
ACETASOL HC, ACETIC ACID, GLACIAL  
ACETAZOLAMIDE, ACETAZOLAMIDE  
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM  
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM  
ACETIC ACID, ACETIC ACID, GLACIAL  
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL  
ACETYLCYSTEINE, ACETYLCYSTEINE  
ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE  
ACIPHEX, RABEPRAZOLE SODIUM  
ACIPHEX SPRINKLE, RABEPRAZOLE SODIUM  
ACITRETIN, ACITRETIN  
ACTHREL, CORTICORELIN OVINE TRIFLUTATE  
ACTICLATE, DOXYCYCLINE HYCLATE  
ACTICLATE CAP, DOXYCYCLINE HYCLATE  
ACTIGALL, URSODIOL  
ACTIQ, FENTANYL CITRATE  
ACTIVELLA, ESTRADIOL  
ACTONEL, RISEDRONATE SODIUM  
ACTOPLUS MET, METFORMIN HYDROCHLORIDE  
ACTOPLUS MET XR, METFORMIN HYDROCHLORIDE  
ACTOS, PIOGLITAZONE HYDROCHLORIDE  
ACULAR, KETOROLAC TROMETHAMINE  
ACULAR LS, KETOROLAC TROMETHAMINE  
ACUVAIL, KETOROLAC TROMETHAMINE  
ACYCLOVIR, ACYCLOVIR  
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM  
ACZONE, DAPSONE  
ADAGEN, PEGADEMASE BOVINE  
ADALAT CC, NIFEDIPIINE  
ADAPALENE, ADAPALENE  
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE  
ADASUVE, LOXPINE  
ADCIRCA, Tadalafil  
ADDERALL XR 10, AMPHETAMINE ASPARTATE  
ADDERALL XR 15, AMPHETAMINE ASPARTATE  
ADDERALL XR 20, AMPHETAMINE ASPARTATE  
ADDERALL XR 25, AMPHETAMINE ASPARTATE  
ADDERALL XR 30, AMPHETAMINE ASPARTATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* A \*\*

ADDERALL XR 5, AMPHETAMINE ASPARTATE  
ADDYI, FLIBANSERIN  
ADEFOVIR DIPIVOXIL, ADEFOVIR DIPIVOXIL  
ADEMPAS, RIOCIGUAT  
ADENOCARD, ADENOSINE  
ADENOSINE, ADENOSINE  
ADIPEX-P, PHENTERMINE HYDROCHLORIDE  
ADLYXIN, LIXISENATIDE  
ADMELOG, INSULIN LISPRO  
ADMELOG SOLOSTAR, INSULIN LISPRO  
ADRENAClick, EPINEPHRINE  
ADRENALIN, EPINEPHRINE  
ADREVIEW, IOBENGUANE SULFATE I-123  
ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE  
ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE  
ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE  
ADVAIR HFA, FLUTICASONE PROPIONATE  
ADVIL, IBUPROFEN (OTC)  
ADVIL, IBUPROFEN SODIUM (OTC)  
ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)  
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)  
ADVIL COLD AND SINUS, IBUPROFEN (OTC)  
ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)  
ADVIL LIQUI-GELS, IBUPROFEN (OTC)  
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)  
ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)  
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)  
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)  
ADZENYS ER, AMPHETAMINE  
ADZENYS XR-ODT, AMPHETAMINE  
AEROSPAN HFA, FLUNISOLIDE  
AFEDITAB CR, NIFEDIPINE  
AFINITOR, EVEROLIMUS  
AFINITOR DISPERZ, EVEROLIMUS  
AFIRMELLE, ETHINYLMESTRADIOL  
AFREZZA, INSULIN RECOMBINANT HUMAN  
AFRINOL, PSEUDOEPHEDRINE SULFATE (OTC)  
AGGRASTAT, TIROFIBAN HYDROCHLORIDE  
AGGRENOX, ASPIRIN  
AGRYLIN, ANAGRELIDE HYDROCHLORIDE  
AIRDUO RESPICLICK, FLUTICASONE PROPIONATE  
AK-FLUOR 10%, FLUORESCIN SODIUM  
AK-FLUOR 25%, FLUORESCIN SODIUM  
AKBETA, LEVOBUNOLOL HYDROCHLORIDE  
AKOVAZ, EPHEDRINE SULFATE  
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE  
AKTEN, LIDOCAINE HYDROCHLORIDE  
AKTIPIAK, BENZOYL PEROXIDE  
AKTOB, TOBRAMYCIN  
AKYNZEO, NETUPITANT  
ALA-CORT, HYDROCORTISONE  
ALA-SCALP, HYDROCORTISONE  
ALAVERT, LORATADINE (OTC)  
ALAWAY, KETOTIFEN FUMARATE (OTC)  
ALBENZA, ALBENDAZOLE  
ALBUTEROL SULFATE, ALBUTEROL SULFATE  
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE  
ALCAINE, PROPARACAINA HYDROCHLORIDE  
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE  
ALDACTAZIDE, HYDROCHLOROTHIAZIDE  
ALDACTONE, SPIRONOLACTONE  
ALDARA, IMIQUIMOD  
ALECENSA, ALECTINIB HYDROCHLORIDE  
ALENDRONATE SODIUM, ALENDRONATE SODIUM  
ALEVE, NAPROXEN SODIUM (OTC)

**APPENDIX A - PRODUCT NAME INDEX**

\*\* A \*\*

ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)  
ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)  
ALFENTA, ALFENTANIL HYDROCHLORIDE  
ALFENTANIL, ALFENTANIL HYDROCHLORIDE  
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE  
ALIMTA, PEMETREXED DISODIUM  
ALINIA, NITAZOXANIDE  
ALIQOPA, COPANLISIB DIHYDROCHLORIDE  
ALKERAN, MELPHALAN  
ALLEGRA, FEXOFENADINE HYDROCHLORIDE  
ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)  
ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)  
ALLI, ORLISTAT (OTC)  
ALLOPURINOL, ALLOPURINOL  
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM  
ALLZITAL, ACETAMINOPHEN  
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE  
ALOCRIL, NEDOCROMIL SODIUM  
ALOMIDE, LODOXAMIDE TROMETHAMINE  
ALOPRIM, ALLOPURINOL SODIUM  
ALORA, ESTRADIOL  
ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE  
ALOXI, PALONOSETRON HYDROCHLORIDE  
ALPHAGAN P, BRIMONIDINE TARTRATE  
ALPRAZOLAM, ALPRAZOLAM  
ALPROSTADIL, ALPROSTADIL  
ALREX, LOTEPELDNOL ETABONATE  
ALTABAX, RETAPAMULIN  
ALTACE, RAMIPRIL  
ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE  
ALTAVERA, ETHINYLMESTRADIOL  
ALTOPREV, LOVASTATIN  
ALUNBRIG, BRIGATINIB  
ALVESCO, CICLESONIDE  
ALYACEN 1/35, ETHINYLMESTRADIOL  
ALYACEN 7/7/7, ETHINYLMESTRADIOL  
AMABELZ, ESTRADIOL  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
AMARYL, GLIMEPIRIDE  
AMBIEN, ZOLPIDEM TARTRATE  
AMBIEN CR, ZOLPIDEM TARTRATE  
AMBISOME, AMPHOTERICIN B  
AMCINONIDE, AMCINONIDE  
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE  
AMERGE, NARATRIPTAN HYDROCHLORIDE  
AMICAR, AMINOCAPROIC ACID  
AMIDATE, ETOMIDATE  
AMIFOSTINE, AMIFOSTINE  
AMIKACIN SULFATE, AMIKACIN SULFATE  
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE  
AMILORIDE HYDROCHLORIDE AND HYDROCHLORTHIAZIDE, AMILORIDE HYDROCHLORIDE  
AMINO ACIDS, AMINO ACIDS  
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE  
AMINOCAPROIC, AMINOCAPROIC ACID  
AMINOCAPROIC ACID, AMINOCAPROIC ACID  
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID  
AMINOPHYLLINE, AMINOPHYLLINE  
AMINOSYN 10%, AMINO ACIDS  
AMINOSYN 10% (PH6), AMINO ACIDS  
AMINOSYN 3.5%, AMINO ACIDS  
AMINOSYN 3.5% M, AMINO ACIDS  
AMINOSYN 5%, AMINO ACIDS  
AMINOSYN 7%, AMINO ACIDS

**APPENDIX A - PRODUCT NAME INDEX**

\*\* A \*\*

AMINOSYN 7% (PH6), AMINO ACIDS  
AMINOSYN 7% W/ ELECTROLYTES, AMINO ACIDS  
AMINOSYN 8.5%, AMINO ACIDS  
AMINOSYN 8.5% (PH6), AMINO ACIDS  
AMINOSYN 8.5% W/ ELECTROLYTES, AMINO ACIDS  
AMINOSYN II 10%, AMINO ACIDS  
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS  
AMINOSYN II 10% W/ ELECTROLYTES, AMINO ACIDS  
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS  
AMINOSYN II 7%, AMINO ACIDS  
AMINOSYN II 8.5%, AMINO ACIDS  
AMINOSYN II 8.5% W/ ELECTROLYTES, AMINO ACIDS  
AMINOSYN-HBC 7%, AMINO ACIDS  
AMINOSYN-HF 8%, AMINO ACIDS  
AMINOSYN-PF 10%, AMINO ACIDS  
AMINOSYN-PF 7%, AMINO ACIDS  
AMINOSYN-RF 5.2%, AMINO ACIDS  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
AMITIZA, LUBIPROSTONE  
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE  
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE AND BENAZEPHIL HYDROCHLORIDE, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE  
AMMONIA N 13, AMMONIA N-13  
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE  
AMMONIUM LACTATE, AMMONIUM LACTATE  
AMMONUL, SODIUM BENZOATE  
AMNESTEEM, ISOTRETINOIN  
AMOXAPINE, AMOXAPINE  
AMOXICILLIN, AMOXICILLIN  
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN  
AMOXICILLIN PEDIATRIC, AMOXICILLIN  
AMOXIL, AMOXICILLIN  
AMPHADASE, HYALURONIDASE  
AMPHOTERICIN B, AMPHOTERICIN B  
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM  
AMPICILLIN SODIUM, AMPICILLIN SODIUM  
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE  
AMPYRA, DALFAMPRIDINE  
AMRINONE LACTATE, INAMRINONE LACTATE  
AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE  
AMYVID, FLORBETAPIR F-18  
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT  
ANADROL-50, OXYMETHOLONE  
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE  
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE  
ANAPROX, NAPROXEN SODIUM  
ANAPROX DS, NAPROXEN SODIUM  
ANASTROZOLE, ANASTROZOLE  
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM  
ANCOBON, FLUCYTOSINE  
ANDRODERM, TESTOSTERONE  
ANDROGEL, TESTOSTERONE  
ANDROID 25, METHYLTESTOSTERONE  
ANECTINE, SUCCINYLCHOLINE CHLORIDE  
ANEXSIA 5/325, ACETAMINOPHEN  
ANEXSIA 7.5/325, ACETAMINOPHEN  
ANGELIQ, DROSPIRENONE  
ANGIOMAX, BIVALIRUDIN  
ANORO ELLIPTA, UMECLIDINIUM BROMIDE  
ANTABUSE, DISULFIRAM  
ANTARA (MICRONIZED), FENOFLIBRATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* A \*\*

ANTHELIOS 20, AVOBENZONE (OTC)  
ANTHELIOS 40, AVOBENZONE (OTC)  
ANTHELIOS SX, AVOBENZONE (OTC)  
ANTIZOL, FOMEPIZOLE  
ANUSOL HC, HYDROCORTISONE  
APIDRA, INSULIN GLULISINE RECOMBINANT  
APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT  
APLENZIN, BUPROPION HYDROBROMIDE  
APOKYN, APOMORPHINE HYDROCHLORIDE  
APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE  
APREPITANT, APREPITANT  
APRISO, MESALAMINE  
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE  
APTIOM, ESLICARBAZEPINE ACETATE  
APTIVUS, TIPRANAVIR  
AQUASOL A, VITAMIN A PALMITATE  
ARANELLE, ETHINYLL ESTRADIOL  
ARAVA, LEFLUNOMIDE  
ARCAPTA NEOHALER, INDACATEROL MALEATE  
ARESTIN, MINOCYCLINE HYDROCHLORIDE  
ARGATROBAN, ARGATROBAN  
ARGATROBAN IN 0.9% SODIUM CHLORIDE, ARGATROBAN  
ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN  
ARICEPT, DONEPEZIL HYDROCHLORIDE  
ARICEPT ODT, DONEPEZIL HYDROCHLORIDE  
ARIMIDEX, ANASTROZOLE  
ARIPIPRAZOLE, ARIPIPRAZOLE  
ARISTADA, ARIPIPRAZOLE LAUROXIL  
ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE  
ARIXTRA, FONDAPARINUX SODIUM  
ARMODAFINIL, ARMODAFINIL  
ARMONAIR RESPCLICK, FLUTICASONE PROPIONATE  
ARNUITY ELLIPTA, FLUTICASONE FUROATE  
AROMASIN, EXEMESTANE  
ARRANON, NELARABINE  
ARTHROTEC, DICLOFENAC SODIUM  
ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, ARTICAINE HYDROCHLORIDE  
ARYMO ER, MORPHINE SULFATE  
ASACOL HD, MESALAMINE  
ASCLERA, POLIDOCANOL  
ASCOR, ASCORBIC ACID  
ASHLYNA, ETHINYLL ESTRADIOL  
ASMANEX HFA, MOMETASONE FUROATE  
ASMANEX TWISTHALER, MOMETASONE FUROATE  
ASPIRIN, ASPIRIN (OTC)  
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN  
ASTAGRAF XL, TACROLIMUS  
ASTELIN, AZELASTINE HYDROCHLORIDE  
ASTEPRO, AZELASTINE HYDROCHLORIDE  
ASTRAMORPH PF, MORPHINE SULFATE  
ATACAND, CANDESARTAN CILEXETIL  
ATACAND HCT, CANDESARTAN CILEXETIL  
ATAZANAVIR SULFATE, ATAZANAVIR SULFATE  
ATELVIA, RISEDRONATE SODIUM  
ATENOLOL, ATENOLOL  
ATENOLOL AND CHLORTHALIDONE, ATENOLOL  
ATHENTIA NEXT, LEVONORGESTREL (OTC)  
ATIVAN, LORAZEPAM  
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE  
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM  
ATOVAQUONE, ATOVAQUONE  
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE  
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE  
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE  
ATRALIN, TRETINOIN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* A \*\*

ATRIDOX, DOXYCYCLINE HYCLATE  
ATRIPLA, EFAVIRENZ  
ATROOPEN, ATROPINE  
ATROPINE SULFATE, ATROPINE SULFATE  
ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE  
ATROPINE SULFATE LIFESHIELD ABBOJECT SYRINGE, ATROPINE SULFATE  
ATROVENT, IPRATROPIUM BROMIDE  
ATROVENT HFA, IPRATROPIUM BROMIDE  
AUBAGIO, TERIFLUNOMIDE  
AUGMENTIN '125', AMOXICILLIN  
AUGMENTIN '250', AMOXICILLIN  
AUGMENTIN '875', AMOXICILLIN  
AUGMENTIN XR, AMOXICILLIN  
AUROVELA 1.5/30, ETHINYL ESTRADIOL  
AUROVELA 1/20, ETHINYL ESTRADIOL  
AUROVELA 24 FE, ETHINYL ESTRADIOL  
AUROVELA FE 1.5/30, ETHINYL ESTRADIOL  
AUROVELA FE 1/20, ETHINYL ESTRADIOL  
AURYXIA, FERRIC CITRATE  
AUSTEDO, DEUTETRABENAZINE  
AVI-Q, EPINEPHRINE  
AVAGARD, ALCOHOL (OTC)  
AVAGE, TAZAROTENE  
AVALIDE, HYDROCHLOROTHIAZIDE  
AVANDIA, ROSIGLITAZONE MALEATE  
AVAPRO, IRBESARTAN  
AVC, SULFANILAMIDE  
AVEED, TESTOSTERONE UNDECANOATE  
AVELOX, MOXIFLOXACIN HYDROCHLORIDE  
AVIANE-28, ETHINYL ESTRADIOL  
AVITA, TRETINOIN  
AVODART, DUTASTERIDE  
AVYCAZ, AVIBACTAM SODIUM  
AXERT, ALMOTRIPTAN MALATE  
AXID AR, NIZATIDINE (OTC)  
AXUMIN, FLUCICLOVINE F-18  
AYGESTIN, NORETHINDRONE ACETATE  
AYUNA, ETHINYL ESTRADIOL  
AZACITIDINE, AZACITIDINE  
AZACTAM, AZTREONAM  
AZACTAM IN PLASTIC CONTAINER, AZTREONAM  
AZASAN, AZATHIOPRINE  
AZASITE, AZITHROMYCIN  
AZATHIOPRINE, AZATHIOPRINE  
AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE  
AZELEX, AZELAIC ACID  
AZILECT, RASAGILINE MESYLATE  
AZITHROMYCIN, AZITHROMYCIN  
AZOPT, BRINZOLAMIDE  
AZOR, AMLODIPINE BESYLATE  
AZTREONAM, AZTREONAM  
AZULFIDINE, SULFASALAZINE  
AZULFIDINE EN-TABS, SULFASALAZINE

\*\* B \*\*

BACIIM, BACITRACIN  
BACITRACIN, BACITRACIN  
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC  
BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN  
BACLOFEN, BACLOFEN  
BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* B \*\*

BACTRIM, SULFAMETHOXAZOLE  
BACTRIM DS, SULFAMETHOXAZOLE  
BACTROBAN, MUPIROCIN CALCIUM  
BAL, DIMERCAPROL  
BALANCED SALT, CALCIUM CHLORIDE  
BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM  
BALZIVA-28, ETHINYL ESTRADIOL  
BANZEL, RUFINAMIDE  
BARACLUDE, ENTECAVIR  
BASAGLAR, INSULIN GLARGINE  
BAXDELA, DELAFLOXACIN MEGLUMINE  
BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE  
BEKYREE, DESOGESTREL  
BELBUCA, BUPRENORPHINE HYDROCHLORIDE  
BELEODAQ, BELINOSTAT  
BELSOMRA, SUVOREXANT  
BELVIQ, LORCASERIN HYDROCHLORIDE  
BELVIQ XR, LORCASERIN HYDROCHLORIDE  
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLORTIAZIDE, BENAZEPRIL HYDROCHLORIDE  
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE  
BENDEKA, BENDAMUSTINE HYDROCHLORIDE  
BENICAR, OLMESARTAN MEDOXOMIL  
BENICAR HCT, HYDROCHLORTIAZIDE  
BENTYL, DICYCLOMINE HYDROCHLORIDE  
BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE  
BENZACLIN, BENZOYL PEROXIDE  
BENZAMYCIN, BENZOYL PEROXIDE  
BENZNIDAZOLE, BENZNIDAZOLE  
BENZONATATE, BENZONATATE  
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE  
BEPREVE, BEPOTASTINE BESILATE  
BESIVANCE, BESIFLOXACIN HYDROCHLORIDE  
BETA-VAL, BETAMETHASONE VALERATE  
BETADINE, POVIDONE-IODINE  
BETAGAN, LEVOBUNOLOL HYDROCHLORIDE  
BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE  
BETAPACE, SOTALOL HYDROCHLORIDE  
BETAPACE AF, SOTALOL HYDROCHLORIDE  
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE  
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE  
BETHKIS, TOBRAMYCIN  
BETIMOL, TIMOLOL  
BETOPTIC, BETAXOLOL HYDROCHLORIDE  
BETOPTIC S, BETAXOLOL HYDROCHLORIDE  
BEVESPI AEROSPHERE, FORMOTEROL FUMARATE  
BEVYXXA, BETRIXABAN  
BEXAROTENE, BEXAROTENE  
BEYAZ, DROSPIRENONE  
BIAXIN, CLARITHROMYCIN  
BICALUTAMIDE, BICALUTAMIDE  
BICILLIN C-R, PENICILLIN G BENZATHINE  
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE  
BICILLIN L-A, PENICILLIN G BENZATHINE  
BICNU, CARMUSTINE  
BIDIL, HYDRALAZINE HYDROCHLORIDE  
BILTRICIDE, PRAZIQUANTEL  
BIMATOPROST, BIMATOPROST  
BINOSTO, ALENDRONATE SODIUM  
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)  
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE  
BISOPROLOL FUMARATE AND HYDROCHLORTIAZIDE, BISOPROLOL FUMARATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* B \*\*

BIVALIRUDIN, BIVALIRUDIN  
BIVALIRUDIN IN 0.9% SODIUM CHLORIDE, BIVALIRUDIN  
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE  
BLEPH-10, SULFACETAMIDE SODIUM  
BLEPHAMIDE, PREDNISOLONE ACETATE  
BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE  
BLISOVI 24 FE, ETHINYL ESTRADIOL  
BLISOVI FE 1.5/30, ETHINYL ESTRADIOL  
BLISOVI FE 1/20, ETHINYL ESTRADIOL  
BLOXIVERZ, NEOSTIGMINE METHYLSULFATE  
BONIVA, IBANDRONATE SODIUM  
BONJESTA, DOXYLAMINE SUCCINATE  
BONTRIL PDM, PHENDIMETRAZINE TARTRATE  
BORTEZOMIB, BORTEZOMIB  
BOSULIF, BOSUTINIB MONOHYDRATE  
BRAVELLE, UROFOLLITROPIN  
BREO ELLIPTA, FLUTICASONE FUROATE  
REVIBLOC, ESMOLOL HYDROCHLORIDE  
REVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE  
REVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE  
REVICON 28-DAY, ETHINYL ESTRADIOL  
REVITAL SODIUM, METHOHEXITAL SODIUM  
BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)  
BRIDION, SUGAMMADEX SODIUM  
BRIELLYN, ETHINYL ESTRADIOL  
BRILINTA, TICAGRELOR  
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE  
BRISDELLE, PAROXETINE MESYLATE  
BRIVIACT, BRIVARACETAM  
BROMFED-DM, BROMPHENIRAMINE MALEATE  
BROMFENAC SODIUM, BROMFENAC SODIUM  
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE  
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE,  
BROMSITE, BROMFENAC SODIUM  
BRONCHO SALINE, SODIUM CHLORIDE (OTC)  
BROVANA, ARFORMOTEROL TARTRATE  
BSS, CALCIUM CHLORIDE  
BSS PLUS, CALCIUM CHLORIDE  
BUDESONIDE, BUDESONIDE (OTC)  
BUDESONIDE, BUDESONIDE  
BUMETANIDE, BUMETANIDE  
BUMEX, BUMETANIDE  
BUNAVAIL, BUPRENORPHINE HYDROCHLORIDE  
BUPHENYL, SODIUM PHENYLBUTYRATE  
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE  
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE  
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE  
BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE  
BUPRENEX, BUPRENORPHINE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE  
BUSULFAN, BUSULFAN  
BUSULFEX, BUSULFAN  
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN  
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN  
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN  
BUTALBITAL, ASPIRIN, CAFFEINE AND CODEINE PHOSPHATE, ASPIRIN  
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN  
BUTAPAP, ACETAMINOPHEN  
BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)  
BUTISOL SODIUM, BUTABARBITAL SODIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* B \*\*

BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE  
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE  
BUTRANS, BUPRENORPHINE  
BYDUREON, EXENATIDE SYNTHETIC  
BYDUREON BCISE, EXENATIDE  
BYDUREON PEN, EXENATIDE SYNTHETIC  
BYETTA, EXENATIDE SYNTHETIC  
BYSTOLIC, NEBIVOLOL HYDROCHLORIDE  
BYVALSON, NEBIVOLOL HYDROCHLORIDE

\*\* C \*\*

CABERGOLINE, CABERGOLINE  
CABOMETYX, CABOZANTINIB S-MALATE  
CADUET, AMLODIPINE BESYLATE  
CAF'CIT, CAFFEINE CITRATE  
CAFERGOT, CAFFEINE  
CAFFEINE CITRATE, CAFFEINE CITRATE  
CALAN, VERAPAMIL HYDROCHLORIDE  
CALAN SR, VERAPAMIL HYDROCHLORIDE  
CALCIPOTRIENE, CALCIPOTRIENE  
CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CALCITONIN-SALMON, CALCITONIN SALMON  
CALCITRIOL, CALCITRIOL  
CALCIUM ACETATE, CALCIUM ACETATE  
CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE  
CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM  
CALCIUM GLUCONATE, CALCIUM GLUCONATE  
CALDOLOR, IBUPROFEN  
CALQUENCE, ACALABRUTINIB  
CAMBIA, DICLOFENAC POTASSIUM  
CAMILA, NORETHINDRONE  
CAMPTOSAR, IRINOTECAN HYDROCHLORIDE  
CANASA, MESALAMINE  
CANCIDAS, CASPOFUNGIN ACETATE  
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL  
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL  
CAPASTAT SULFATE, CAPREOMYCIN SULFATE  
CAPECITABINE, CAPECITABINE  
CAPEX, FLUOCINOLONE ACETONIDE  
CAPITAL SOLEIL 15, AVOBENZONE (OTC)  
CAPRELSA, VANDETANIB  
CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE  
CAPTOPRIL, CAPTOPRIL  
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL  
CARAC, FLUOROURACIL  
CARAFATE, SUCRALFATE  
CARBAGLU, CARGLUMIC ACID  
CARBAMAZEPINE, CARBAMAZEPINE  
CARBATROL, CARBAMAZEPINE  
CARBIDOPA, CARBIDOPA  
CARBIDOPA AND LEVODOPA, CARBIDOPA  
CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA  
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE  
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE  
CARBOPLATIN, CARBOPLATIN  
CARDENE, NICARDIPINE HYDROCHLORIDE  
CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82  
CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT  
CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
CARDIZEM, DILTIAZEM HYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* C \*\*

CARDIZEM CD, DILTIAZEM HYDROCHLORIDE  
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE  
CARDURA, DOXAZOSIN MESYLATE  
CARDURA XL, DOXAZOSIN MESYLATE  
CARISOPRODOL, CARISOPRODOL  
CARISOPRODOL AND ASPIRIN, ASPIRIN  
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN  
CARNEXIV, CARBAMAZEPINE  
CARNITOR, LEVOCARNITINE  
CARNITOR SF, LEVOCARNITINE  
CAROSPIR, SPIRONOLACTONE  
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE  
CARTIA XT, DILTIAZEM HYDROCHLORIDE  
CARVEDILOL, CARVEDILOL  
CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE  
CASODEX, BICALUTAMIDE  
CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE  
CASPORYN HC, HYDROCORTISONE  
CATAPRES, CLONIDINE HYDROCHLORIDE  
CATAPRES-TTS-1, CLONIDINE  
CATAPRES-TTS-2, CLONIDINE  
CATAPRES-TTS-3, CLONIDINE  
CAVERJECT, ALPROSTADIL  
CAVERJECT IMPULSE, ALPROSTADIL  
CAYSTON, AZTREONAM  
CEFACLOR, CEFACLOR  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM  
CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM  
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM  
CEFDINIR, CEFDINIR  
CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE  
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE  
CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE  
CEFIXIME, CEFIXIME  
CEFOTAN, CEFOTETAN DISODIUM  
CEFOTAXIME, CEFOTAXIME SODIUM  
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM  
CEFOTETAN, CEFOTETAN DISODIUM  
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, CEFOTETAN DISODIUM  
CEFOXITIN, CEFOXITIN SODIUM  
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM  
CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM  
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL  
CEFPROZIL, CEFPROZIL  
CEFTAZIDIME, CEFTAZIDIME  
CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME  
CEFTIN, CEFUROXIME AXETIL  
CEFTRIAXONE, CEFTRIAXONE SODIUM  
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM  
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM  
CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM  
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM  
CEFUROXIME AXETIL, CEFUROXIME AXETIL  
CEFUROXIME SODIUM, CEFUROXIME SODIUM  
CELEBREX, CELECOXIB  
CELECOXIB, CELECOXIB  
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE  
CELEXA, CITALOPRAM HYDROBROMIDE  
CELLCEPT, MYCOPHENOLATE MOFETIL  
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE  
CELONTIN, METHSUXIMIDE  
CENTANY, MUPIROCIN  
CEPHALEXIN, CEPHALEXIN  
CERDELGA, ELIGLUSTAT TARTRATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* C \*\*

CEREBYX, FOSPHENYTOIN SODIUM  
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT  
CEREZYME, IMIGLUCERASE  
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE  
CERVIDIL, DINOPROSTONE  
CESAMET, NABILONE  
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE  
CETROTIDE, CETRORELIK  
CETYLEV, ACETYL CYSTEINE  
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE  
CHANTIX, VARENICLINE TARTRATE  
CHEMET, SUCCIMER  
CHENODIOL, CHENODIOL  
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)  
CHILDREN'S ADVIL, IBUPROFEN (OTC)  
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)  
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)  
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)  
CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
CHILDREN'S ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CLARITIN, LORATADINE (OTC)  
CHILDREN'S ELIXSURE, IBUPROFEN (OTC)  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)  
CHILDREN'S MOTRIN, IBUPROFEN (OTC)  
CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)  
CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN  
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)  
CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE  
CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)  
CHLDIAZEPoxide AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE  
CHLDIAZEPoxide HYDROCHLORIDE, CHLDIAZEPoxide HYDROCHLORIDE  
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE  
CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE  
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE  
CHLORTHIAZIDE, CHLORTHIAZIDE  
CHLORTHIAZIDE SODIUM, CHLORTHIAZIDE SODIUM  
CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)  
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE  
CHLORPROPAMIDE, CHLORPROPAMIDE  
CHLORTHALIDONE, CHLORTHALIDONE  
CHLORZOXAZONE, CHLORZOXAZONE  
CHOLAC, LACTULOSE  
CHOLBAM, CHOLIC ACID  
CHOLEDYL SA, OXTRIPHYLLINE  
CHOLESTYRAMINE, CHOLESTYRAMINE  
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE  
CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT  
CHOLINE C-11, CHOLINE C-11

**APPENDIX A - PRODUCT NAME INDEX****\*\* C \*\***

CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC  
CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE  
CIALIS, TADALAFIL  
CICLOPIROX, CICLOPIROX  
CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)  
CIDOFOVIR, CIDOFOVIR  
CILOSTAZOL, CILOSTAZOL  
CILOXAN, CIPROFLOXACIN HYDROCHLORIDE  
CIMETIDINE, CIMETIDINE (OTC)  
CIMETIDINE, CIMETIDINE  
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE  
CINVANTI, APREPITANT  
CIPRO, CIPROFLOXACIN  
CIPRO, CIPROFLOXACIN HYDROCHLORIDE  
CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE  
CIPRODEX, CIPROFLOXACIN  
CIPROFLOXACIN, CIPROFLOXACIN  
CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN  
CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT  
CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT  
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE  
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE  
CISPLATIN, CISPLATIN  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
CITANESE FORTE DENTAL, EPINEPHRINE BITARTRATE  
CLADRIBINE, CLADRIBINE  
CLAFORAN, CEFOTAXIME SODIUM  
CLARAVIS, ISOTRETINOIN  
CLARINEX, DESLORATADINE  
CLARINEX D 24 HOUR, DESLORATADINE  
CLARINEX-D 12 HOUR, DESLORATADINE  
CLARITHROMYCIN, CLARITHROMYCIN  
CLARITIN, LORATADINE (OTC)  
CLARITIN HIVES RELIEF, LORATADINE (OTC)  
CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)  
CLARITIN REDITABS, LORATADINE (OTC)  
CLARITIN-D, LORATADINE (OTC)  
CLARITIN-D 24 HOUR, LORATADINE (OTC)  
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)  
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE  
CLENPIQ, CITRIC ACID  
CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE  
CLEOCIN, CLINDAMYCIN PHOSPHATE  
CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE  
CLEOCIN T, CLINDAMYCIN PHOSPHATE  
CLEVIPREX, CLEVIDIPIINE  
CLIMARA, ESTRADIOL  
CLIMARA PRO, ESTRADIOL  
CLINDA-DERM, CLINDAMYCIN PHOSPHATE  
CLINDAGEL, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
CLINDAMYCIN HYDROCLHORIDE, CLINDAMYCIN HYDROCHLORIDE  
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE  
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE  
CLINDEsse, CLINDAMYCIN PHOSPHATE  
CLINDEts, CLINDAMYCIN PHOSPHATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* C \*\*

CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLOBETASOL PROPIONATE (EMOLlient), CLOBETASOL PROPIONATE  
CLOBEX, CLOBETASOL PROPIONATE  
CLODERM, CLOCORTOLONE PIVALATE  
CLOFARABINE, CLOFARABINE  
CLOLAR, CLOFARABINE  
CLOMID, CLOMIPHENE CITRATE  
CLOMIPHENE CITRATE, CLOMIPHENE CITRATE  
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE  
CLONAZEPAM, CLONAZEPAM  
CLONIDINE, CLONIDINE  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
CLONIDINE HYDROCHLORIDE , CLONIDINE HYDROCHLORIDE  
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE  
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM  
CLOROTEKAL, CHLOROPROCaine HYDROCHLORIDE  
CLORPRES, CHLORTHALIDONE  
CLOTrimazole, CLOTrimazole (OTC)  
CLOTrimazole, CLOTrimazole  
CLOTrimazole AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CLOZAPINE, CLOZAPINE  
CLOZARIL, CLOZAPINE  
COARTEM, ARTEMETHER  
CODEINE SULFATE, CODEINE SULFATE  
COGENTIN, BENZTROPINE MESYLATE  
COL-PROBENECID, COLCHICINE  
COLAZAL, BALSALAZIDE DISODIUM  
COLCRYS, COLCHICINE  
COLESTID, COLESTIPOL HYDROCHLORIDE  
COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE  
COLGATE TOTAL, SODIUM FLUORIDE (OTC)  
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM  
COLOCORT, HYDROCORTISONe  
COLPREP KIT, MAGNESIUM SULFATE  
COLY-MYCIN M, COLISTIMETHATE SODIUM  
COLY-MYCIN S, COLISTIN SULFATE  
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350  
COMBIGAN, BRIMONIDINE TARTRATE  
COMBIPATCH, ESTRADIOL  
COMBIVENT RESPIMAT, ALBUTEROL SULFATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* C \*\*

COMBIVIR, LAMIVUDINE  
COMETRIQ, CABOZANTINIB S-MALATE  
COMMIT, NICOTINE POLACRILEX (OTC)  
COMPLERA, EMTRICITABINE  
COMPROM, PROCHLORPERAZINE  
COMTAN, ENTACAPONE  
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE  
CONDYLOX, PODOFILOX  
CONRAY, IOTHALAMATE MEGLUMINE  
CONRAY 30, IOTHALAMATE MEGLUMINE  
CONRAY 43, IOTHALAMATE MEGLUMINE  
CONSTILAC, LACTULOSE  
CONTRAVE, BUPROPION HYDROCHLORIDE  
CONZIP, TRAMADOL HYDROCHLORIDE  
COPAXONE, GLATIRAMER ACETATE  
COPEGUS, RIBAVIRIN  
CORDRAN, FLURANDRENOLIDE  
CORDRAN SP, FLURANDRENOLIDE  
COREG, CARVEDILOL  
COREG CR, CARVEDILOL PHOSPHATE  
CORGARD, NADOLOL  
CORLANOR, IVABRADINE HYDROCHLORIDE  
CORLOPAM, FENOLDOPAM MESYLATE  
CORMAX, CLOBETASOL PROPIONATE  
CORPHEDRA, EPHEDRINE SULFATE  
CORTEF, HYDROCORTISONE  
CORTENEMA, HYDROCORTISONE  
CORTIFOAM, HYDROCORTISONE ACETATE  
CORTISONE ACETATE, CORTISONE ACETATE  
CORTISPORIN, BACITRACIN ZINC  
CORTISPORIN, HYDROCORTISONE  
CORTISPORIN, HYDROCORTISONE ACETATE  
CORTROSYN, COSYNTROPIN  
CORVERT, IBUTILIDE FUMARATE  
CORZIDE, BENDROFLUMETHIAZIDE  
COSMEGEN, DACTINOMYCIN  
COSOPT, DORZOLAMIDE HYDROCHLORIDE  
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE  
COSYNTROPIN, COSYNTROPIN  
COTELLIC, COBIMETINIB FUMARATE  
COTEMPLA XR-ODT, METHYLPHENIDATE  
COUMADIN, WARFARIN SODIUM  
COZAAR, LOSARTAN POTASSIUM  
CREON, PANCRELIPASE (AMYLASE)  
CRESEMBIA, ISAVUCONAZONIUM SULFATE  
CRESTOR, ROSUVASTATIN CALCIUM  
CRINONE, PROGESTERONE  
CRIXIVAN, INDINAVIR SULFATE  
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)  
CROMOLYN SODIUM, CROMOLYN SODIUM  
CROTAN, CROTAMITON  
CRYSELLE, ETHINYLY ESTRADIOL  
CUBICIN, DAPTOMYCIN  
CUBICIN RF, DAPTOMYCIN  
CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE  
CUPRIMINE, PENICILLAMINE  
CUROSURF, PORACTANT ALFA  
CUTIVATE, FLUTICASONE PROPIONATE  
CUVPOSA, GLYCOPYRROLATE  
CYANOCOBALAMIN, CYANOCOBALAMIN  
CYANOKIT, HYDROXOCOBALAMIN  
CYCLAFEM 0.5/35, ETHINYLY ESTRADIOL  
CYCLAFEM 1/35, ETHINYLY ESTRADIOL  
CYCLAFEM 7/7/7, ETHINYLY ESTRADIOL  
CYCLESSA, DESOGESTREL

**APPENDIX A - PRODUCT NAME INDEX**

\*\* C \*\*

CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE  
CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE  
CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE  
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE  
CYCLOSET, BROMOCRIPTINE MESYLATE  
CYCLOSPORINE, CYCLOSPORINE  
CYKLOKAPRON, TRANEXAMIC ACID  
CYMBALTA, DULOXETINE HYDROCHLORIDE  
CYONANZ, ETHINYLMESTRADIOL  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
CYSTADANE, BETAINE HYDROCHLORIDE  
CYSTAGON, CYSTEAMINE BITARTRATE  
CYSTARAN, CYSTEAMINE HYDROCHLORIDE  
CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE  
CYSTOGRAFIN, DIATRIZOATE MEGLUMINE  
CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE  
CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE  
CYTARABINE, CYTARABINE  
CYTOMEL, LIOTHYRONINE SODIUM  
CYTOTEC, MISOPROSTOL  
CYTOVENE, GANCICLOVIR SODIUM

\*\* D \*\*

D.H.E. 45, DIHYDROERGOTAMINE MESYLATE  
DACARBAZINE, DACARBAZINE  
DACOGEN, DECITABINE  
DACTINOMYCIN, DACTINOMYCIN  
DAKLINZA, DACLATASVIR DIHYDROCHLORIDE  
DALFAMPRIDINE, DALFAMPRIDINE  
DALIRESP, ROFLUMILAST  
DALVANCE, DALBAVANCIN HYDROCHLORIDE  
DANAZOL, DANAZOL  
DANTRIUM, DANTROLENE SODIUM  
DANTROLENE SODIUM, DANTROLENE SODIUM  
DAPSONE, DAPSONE  
DAPTOMYCIN, DAPTOMYCIN  
DARAPRIM, PYRIMETHAMINE  
DARIFENACIN, DARIFENACIN HYDROBROMIDE  
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE  
DARUNAVIR ETHANOLATE, DARUNAVIR ETHANOLATE  
DASATINIB, DASATINIB  
DASETTA 1/35, ETHINYLMESTRADIOL  
DASETTA 7/7/7, ETHINYLMESTRADIOL  
DATSCAN, IOFLUPANE I-123  
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE  
DAYPRO, OXaprozin  
DAYSEE, ETHINYLMESTRADIOL  
DAYTRANA, METHYLPHENIDATE  
DDAVP, DESMOPRESSIN ACETATE  
DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE  
DECITABINE, DECITABINE  
DEFERASIROX, DEFERASIROX  
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE  
DEFINITY, PERFLUTREN  
DEFITELIO, DEFIBROTIDE SODIUM  
DELAESTRYL, TESTOSTERONE ENANTHATE  
DELESTROGEN, ESTRADIOL VALERATE  
DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* D \*\*

DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)  
DELZICOL, MESALAMINE  
DEMADEX, TORSEMIDE  
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE  
DEMEROL, MEPERIDINE HYDROCHLORIDE  
DEM SER, METYROSINE  
DENAVIR, PENCICLOVIR  
DEPACON, VALPROATE SODIUM  
DEPAKENE, VALPROIC ACID  
DEPAKOTE, DIVALPROEX SODIUM  
DEPAKOTE ER, DIVALPROEX SODIUM  
DEPEN, PENICILLAMINE  
DEPO-ESTRADIOL, ESTRADIOL CYPIONATE  
DEPO-MEDROL, METHYL PREDNISOLONE ACETATE  
DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE  
DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE  
DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE  
DEPOCYT, CYTARABINE  
DERMA-SMOOTH/F/S, FLUOCINOLONE ACETONIDE  
DERMABET, BETAMETHASONE VALERATE  
DERMATOP, PREDNICARBATE  
DERMATOP E EMOLlient, PREDNICARBATE  
DERMOTIC, FLUOCINOLONE ACETONIDE  
DESCOVY, EMTRICITABINE  
DESFERAL, DEFEROXAMINE MESYLATE  
DESIPIRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE  
DESLORATADINE, DESLORATADINE  
DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE  
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE  
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE  
DESOGEN, DESOGESTREL  
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL  
DESONATE, DESONIDE  
DESONIDE, DESONIDE  
DESOWEN, DESONIDE  
DESOXIMETASONE, DESOXIMETASONE  
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE  
DESVENLAFAxINE, DESVENLAFAxINE  
DESVENLAFAxINE, DESVENLAFAxINE FUMARATE  
DESVENLAFAxINE SUCCINATE, DESVENLAFAxINE SUCCINATE  
DETROL, TOLTERODINE TARTRATE  
DETROL LA, TOLTERODINE TARTRATE  
DEXAMETHASONE, DEXAMETHASONE  
DEXAMETHASONE INTENSOL, DEXAMETHASONE  
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE  
DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE  
DEXASPORIN, DEXAMETHASONE  
DEXBROMPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE SULFATE, DEXBROMPHENIRAMINE MALEATE (OTC)  
DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE  
DEXEDRINE, DEXTROAMPHETAMINE SULFATE  
DEXFERRUM, IRON DEXTRAN  
DEXILANT, DEXLANSOPRAZOLE  
DEXLANSOPRAZOLE, DEXLANSOPRAZOLE  
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE  
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE  
DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE  
DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)  
DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* D \*\*

DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 25%, DEXTROSE  
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE  
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,  
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,  
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,  
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,  
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,  
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,  
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER,  
DEXTROSE 50%, DEXTROSE  
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 60% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE  
DIABETA, GLYBURIDE  
DIABINESE, CHLORPROPAMIDE  
DIAMOX, ACETAZOLAMIDE  
DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

**APPENDIX A - PRODUCT NAME INDEX****\*\* D \*\***

DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DIASTAT, DIAZEPAM  
DIASTAT ACUDIAL, DIAZEPAM  
DIAZEPAM, DIAZEPAM  
DIAZEPAM INTENSOL, DIAZEPAM  
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE  
DICLEGIS, DOXYLAMINE SUCCINATE  
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DICLOFENAC SODIUM , DICLOFENAC SODIUM  
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM  
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM  
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE  
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE  
DIDANOSINE, DIDANOSINE  
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE  
DIFFERIN, ADAPALENE (OTC)  
DIFFERIN, ADAPALENE  
DIFICID, FIDAXOMICIN  
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE  
DIFLUCAN, FLUCONAZOLE  
DIFLUNISAL, DIFLUNISAL  
DIGOXIN, DIGOXIN  
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE  
DILANTIN, PHENYTOIN  
DILANTIN, PHENYTOIN SODIUM  
DILANTIN-125, PHENYTOIN  
DILATRATE-SR, ISOSORBIDE DINITRATE  
DILAUDID, HYDROMORPHONE HYDROCHLORIDE  
DILAUDID-HP, HYDROMORPHONE HYDROCHLORIDE  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DILTZAC, DILTIAZEM HYDROCHLORIDE  
DIMENHYDRINATE, DIMENHYDRINATE  
DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE  
DIOVAN, VALSARTAN  
DIOVAN HCT, HYDROCHLOROTHIAZIDE  
DIPENTUM, OLSALAZINE SODIUM  
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE  
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE  
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE  
DIPRIVAN, PROPOFOL  
DIPROLENE, BETAMETHASONE DIPROPIONATE  
DIPROLENE AF, BETAMETHASONE DIPROPIONATE  
DIPYRIDAMOLE, DIPYRIDAMOLE  
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE  
DISULFIRAM, DISULFIRAM  
DISULFIRAM , DISULFIRAM  
DITROPAN XL, OXYBUTYNIN CHLORIDE  
DIURIL, CHLOROTHIAZIDE  
DIURIL, CHLOROTHIAZIDE SODIUM  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
DIVIGEL, ESTRADIOL  
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE  
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE  
DOCETAXEL, DOCETAXEL  
DOFETILIDE, DOFETILIDE  
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* D \*\*

DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE  
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE  
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE  
DOXAPRAM, DOXAPRAM HYDROCHLORIDE  
DORAL, QUAZEPAM  
DORYX, DOXYCYCLINE HYCLATE  
DORYX MPC, DOXYCYCLINE HYCLATE  
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE  
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE  
DOTAREM, GADOTERATE MEGLUMINE  
DOVONEX, CALCIPOTRIENE  
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE  
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE  
DOXEPIН HYDROCHLORIDE, DOXEPIН HYDROCHLORIDE  
DOXERCALCIFEROL, DOXERCALCIFEROL  
DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE  
DOXY 100, DOXYCYCLINE HYCLATE  
DOXY 200, DOXYCYCLINE HYCLATE  
DOXYCYCLINE, DOXYCYCLINE  
DOXYCYCLINE, DOXYCYCLINE HYCLATE  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)  
DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE  
DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT  
DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE  
DRISDOL, ERGOCALCIFEROL  
DRONABINOL, DRONABINOL  
DROPERIDOL, DROPERIDOL  
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE  
DROSPIRENONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONE  
DROXIA, HYDROXYUREA  
DTPA, TECHNETIUM TC-99M PENTETATE KIT  
DUAC, BENZOYL PEROXIDE  
DUAVEE, BAZEDOXIFENE ACETATE  
DUETACT, GLIMEPIRIDE  
DUEXIS, FAMOTIDINE  
DULERA, FORMOTEROL FUMARATE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
DUODOTE, ATROPINE  
DUOPA, CARBIDOPA  
DURACLON, CLONIDINE HYDROCHLORIDE  
DURAGESIC-100, FENTANYL  
DURAGESIC-12, FENTANYL  
DURAGESIC-25, FENTANYL  
DURAGESIC-50, FENTANYL  
DURAGESIC-75, FENTANYL  
DURAMORPH PF, MORPHINE SULFATE  
DURAPREP, IODINE POVACRYLEX (OTC)  
DUREZOL, DIFLUPREDNATE  
DURLAZA, ASPIRIN  
DUTASTERIDE, DUTASTERIDE  
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE  
DUTOPROL, HYDROCHLOROTHIAZIDE  
DUVOID, BETHANECHOL CHLORIDE  
DUZALLO, ALLOPURINOL  
DYANAVEL XR, AMPHETAMINE  
DYAZIDE, HYDROCHLOROTHIAZIDE  
DYLOJECT, DICLOFENAC SODIUM  
DYMISTA, AZELASTINE HYDROCHLORIDE  
DYNA-HEX, CHLORHEXIDINE GLUCONATE (OTC)  
DYNACIN, MINOCYCLINE HYDROCHLORIDE  
DYRENIUM, TRIAMTERENE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* E \*\*

E-Z SCRUB 201, POVIDONE-IODINE (OTC)  
E-Z SCRUB 241, POVIDONE-IODINE (OTC)  
E-Z-HD, BARIUM SULFATE  
E-Z-PAQUE, BARIUM SULFATE  
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE  
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE  
EC-NAPROSYN, NAPROXEN  
ECONAZOLE NITRATE, ECONAZOLE NITRATE  
ECOZA, ECONAZOLE NITRATE  
EDARBI, AZILSARTAN KAMEDOXOMIL  
EDARBYCLOL, AZILSARTAN KAMEDOXOMIL  
EDECRIN, ETHACRYNATE SODIUM  
EDECRIN, ETHACRYNIC ACID  
EDEX, ALPROSTADIL  
EDLUAR, ZOLPIDEM TARTRATE  
EDURANT, RILPIVIRINE HYDROCHLORIDE  
EFAVIRENZ, EFAVIRENZ  
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE  
EFFIENT, PRASUGREL HYDROCHLORIDE  
EFUDEX, FLUOROURACIL  
EGRIFTA, TESAMORELIN ACETATE  
ELDEPRYL, SELEGILINE HYDROCHLORIDE  
ELELYSO, TALIGLUCERASE ALFA  
ELESTAT, EPINASTINE HYDROCHLORIDE  
ELESTRIN, ESTRADIOL  
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE  
ELIDEL, PIMECROLIMUS  
ELIFEMME, ETHINYLMESTRADIOL  
ELIGARD, LEUPROLIDE ACETATE  
ELIMITREX, PERMETHRIN  
ELINEST, ETHINYLMESTRADIOL  
ELIPHOS, CALCIUM ACETATE  
ELIQUIS, APIXABAN  
ELIXOPHYLLIN, THEOPHYLLINE  
ELLA, ULIPRISTAL ACETATE  
ELLENCE, EPIRUBICIN HYDROCHLORIDE  
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE  
ELMIRON, PENTOSAN POLYSULFATE SODIUM  
ELOCON, MOMETASONE FUROATE  
ELOXATIN, OXALIPLATIN  
EMADINE, EMEDASTINE DIFUMARATE  
EMBEDA, MORPHINE SULFATE  
EMBELINE, CLOBETASOL PROPIONATE  
EMBELINE E, CLOBETASOL PROPIONATE  
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM  
EMEND, APREPITANT  
EMEND, FOSAPREPITANT DIMEGLUMINE  
EMFLAZA, DEFLAZACORT  
EMLA, LIDOCAINE  
EMOQUETTE, DESOGESTREL  
EMSAM, SELEGILINE  
EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE  
EMTRIVA, EMTRICITABINE  
EMVERM, MEBENDAZOLE  
ENABLEX, DARIFENACIN HYDROBROMIDE  
ENALAPRIL MALEATE, ENALAPRIL MALEATE  
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE  
ENALAPRILAT, ENALAPRILAT  
ENDARI, L-GLUTAMINE  
ENDOMETRIN, PROGESTERONE  
ENDOSOL EXTRA, CALCIUM CHLORIDE  
ENLON, EDROPHONIUM CHLORIDE  
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM  
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM  
ENPRESSE-28, ETHINYLMESTRADIOL

**APPENDIX A - PRODUCT NAME INDEX**

\*\* E \*\*

ENSKYCE, DESOGESTREL  
ENSTILAR, BETAMETHASONE DIPROPIONATE  
ENTACAPONE, ENTACAPONE  
ENTECAVIR, ENTECAVIR  
ENTEREG, ALVIMOPAN  
ENTOCORT EC, BUDESONIDE  
ENTRESTO, SACUBITRIL  
ENULOSE, LACTULOSE  
ENVARSUS XR, TACROLIMUS  
EOVIST, GADOXETATE DISODIUM  
EPANED, ENALAPRIL MALEATE  
EPANED KIT, ENALAPRIL MALEATE  
EPANOVA, OMEGA-3-CARBOXYLIC ACIDS  
EPCLUSA, SOFOSBUVIR  
EPHEDRINE SULFATE, EPHEDRINE SULFATE  
EPIDUO, ADAPALENE  
EPIDUO FORTE, ADAPALENE  
EPIFOAM, HYDROCORTISONE ACETATE  
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE  
EPINEPHRINE, EPINEPHRINE  
EPIPEN, EPINEPHRINE  
EPIPEN JR., EPINEPHRINE  
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE  
EPITOL, CARBAMAZEPINE  
EPIVIR, LAMIVUDINE  
EPIVIR-HBV, LAMIVUDINE  
EPLERENONE, EPLERENONE  
EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM  
EPROSARTAN MESYLATE, EPROSARTAN MESYLATE  
EPTIFIBATIDE, EPTIFIBATIDE  
EPZICOM, ABACAVIR SULFATE  
EQUETRO, CARBAMAZEPINE  
ERAXIS, ANIDULAFUNGIN  
ERGOCALCIFEROL, ERGOCALCIFEROL  
ERGOLOID MESYLATES, ERGOLOID MESYLATES  
ERGOMAR, ERGOTAMINE TARTRATE  
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE  
ERIVEDGE, VISMODEGIB  
ERRIN, NORETHINDRONE  
ERTACZO, SERTACONAZOLE NITRATE  
ERY-TAB, ERYTHROMYCIN  
ERYC, ERYTHROMYCIN  
ERYGEL, ERYTHROMYCIN  
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE  
ERYTHROCID, ERYTHROMYCIN LACTOBIONATE  
ERYTHROCID STEARATE, ERYTHROMYCIN STEARATE  
ERYTHROMYCIN, ERYTHROMYCIN  
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE  
ESBRIET, PIRFENIDONE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
ESKATA, HYDROGEN PEROXIDE  
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE  
ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE  
ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE  
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)  
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM  
ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM  
ESOMEPRAZOLE STRONTIUM, ESOMEPRAZOLE STRONTIUM  
ESTARYLLA, ETHINYL ESTRADIOL  
ESTAZOLAM, ESTAZOLAM  
ESTRACE, ESTRADIOL  
ESTRADIOL, ESTRADIOL  
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL  
ESTRADIOL AND NORGESTIMATE, ESTRADIOL

**APPENDIX A - PRODUCT NAME INDEX**

\*\* E \*\*

ESTRADIOL VALERATE, ESTRADIOL VALERATE  
ESTRASORB, ESTRADIOL HEMIHYDRATE  
ESTRING, ESTRADIOL  
ESTROGEL, ESTRADIOL  
ESTROPIPATE, ESTROPIPATE  
ESTROSTEP FE, ETHINYL ESTRADIOL  
ESZOPICLONE, ESZOPICLONE  
ETHACRYNATE SODIUM, ETHACRYNATE SODIUM  
ETHACRYNIC ACID, ETHACRYNIC ACID  
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE  
ETHAMOLIN, ETHANOLAMINE OLEATE  
ETHOSUXIMIDE, ETHOSUXIMIDE  
ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
ETHYOL, AMIFOSTINE  
ETIDRONATE DISODIUM, ETIDRONATE DISODIUM  
ETODOLAC, ETODOLAC  
ETOMIDATE, ETOMIDATE  
ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE  
ETOPOSIDE, ETOPOSIDE  
EUCRISA, CRISABOROLE  
EURAX, CROTAMITON  
EVAMIST, ESTRADIOL  
EVEKEO, AMPHETAMINE SULFATE  
EVISTA, RALOXIFENE HYDROCHLORIDE  
EVOCLIN, CLINDAMYCIN PHOSPHATE  
EVOMELA, MELPHALAN HYDROCHLORIDE  
EVOTAZ, ATAZANAVIR SULFATE  
EVOXAC, CEVIMELINE HYDROCHLORIDE  
EVZIO, NALOXONE HYDROCHLORIDE  
EXALGO, HYDROMORPHONE HYDROCHLORIDE  
EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)  
EXELDERM, SULCONAZOLE NITRATE  
EXELON, RIVASTIGMINE  
EXELON, RIVASTIGMINE TARTRATE  
EXEMESTANE, EXEMESTANE  
EXFORGE, AMLODIPINE BESYLATE  
EXFORGE HCT, AMLODIPINE BESYLATE  
EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)  
EXJADE, DEFERASIROX  
EXONDYS 51, ETEPLIRSEN  
EXPAREL, BUPIVACAINE  
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM  
EXTINA, KETOCONAZOLE  
EXTRANEAL, ICODEXTRIN  
EZETIMIBE, EZETIMIBE  
EZETIMIBE AND ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM  
EZETIMIBE AND SIMVASTATIN, EZETIMIBE

\*\* F \*\*

FABIOR, TAZAROTENE  
FACTIVE, GEMIFLOXACIN MESYLATE  
FALLBACK SOLO, LEVONORGESTREL (OTC)  
FALMINA, ETHINYL ESTRADIOL  
FAMCICLOVIR, FAMCICLOVIR  
FAMOTIDINE, FAMOTIDINE (OTC)  
FAMOTIDINE, FAMOTIDINE  
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE  
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE  
FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)  
FANAPT, ILOPERIDONE  
FARESTON, TOREMIFENE CITRATE  
FARXIGA, DAPAGLIFLOZIN PROPANEDIOL  
FARYDAK, PANOBINOSTAT LACTATE  
FASLODEX, FULVESTRANT  
FAYOSIM, ETHINYL ESTRADIOL

**APPENDIX A - PRODUCT NAME INDEX**

\*\* F \*\*

FAZACLO ODT, CLOZAPINE  
FELBAMATE, FELBAMATE  
FELBATOL, FELBAMATE  
FELDENE, PIROXICAM  
FELODIPINE, FELODIPINE  
FEMARA, LETROZOLE  
FEMCON FE, ETHINYL ESTRADIOL  
FEMHRT, ETHINYL ESTRADIOL  
FEMRING, ESTRADIOL ACETATE  
FENOFIBRATE, FENOFIBRATE  
FENOFIBRATE (MICRONIZED), FENOFIBRATE  
FENOFIBRIC ACID, CHOLINE FENOFIBRATE  
FENOGLIDE, FENOFIBRATE  
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE  
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM  
FENTANYL CITRATE, FENTANYL CITRATE  
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE  
FENTANYL-100, FENTANYL  
FENTANYL-12, FENTANYL  
FENTANYL-25, FENTANYL  
FENTANYL-37, FENTANYL  
FENTANYL-50, FENTANYL  
FENTANYL-62, FENTANYL  
FENTANYL-75, FENTANYL  
FENTANYL-87, FENTANYL  
FENTORA, FENTANYL CITRATE  
FERAHHEME, FERUMOXYTOL  
FERRIPROX, DEFERIPRONE  
FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX  
FESOTERODINE FUMARATE, FESOTERODINE FUMARATE  
FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE  
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE  
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE  
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
FIASP, INSULIN ASPART  
FIASP FLEXTOUCH, INSULIN ASPART  
FIBRICOR, FENOFIBRIC ACID  
FINACEA, AZELAIC ACID  
FINACEA, AZELAIC ACID  
FINASTERIDE, FINASTERIDE  
FIORICET W/ CODEINE, ACETAMINOPHEN  
FIORINAL, ASPIRIN  
FIORINAL W/CODEINE, ASPIRIN  
FIRAZYR, ICATIBANT ACETATE  
FIRMAGON, DEGARELIX ACETATE  
FLAGYL, METRONIDAZOLE  
FLAGYL ER, METRONIDAZOLE  
FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE  
FLAREX, FLUOROMETHOLONE ACETATE  
FLAVORED COlestid, COlestipol HYDROCHLORIDE  
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE  
FLECAINIDE ACETATE, FLECAINIDE ACETATE  
FLECTOR, DICLOFENAC EPOLAMINE  
FLOLAN, EPOPROSTENOL SODIUM  
FLOLIPID, SIMVASTATIN  
FLOMAX, TAMSULOSIN HYDROCHLORIDE  
FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)  
FLONASE SENSI MIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)  
FLOVENT DISKUS 100, FLUTICASONE PROPIONATE  
FLOVENT DISKUS 250, FLUTICASONE PROPIONATE  
FLOVENT DISKUS 50, FLUTICASONE PROPIONATE  
FLOVENT HFA, FLUTICASONE PROPIONATE  
FLOWTUSS, GUAIFENESIN  
FLOXURIDINE, FLOXURIDINE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* F \*\*

FLUCONAZOLE, FLUCONAZOLE  
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUCYTOSINE, FLUCYTOSINE  
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE  
FLUMADINE, RIMANTADINE HYDROCHLORIDE  
FLUMAZENIL, FLUMAZENIL  
FLUNISOLIDE, FLUNISOLIDE  
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE  
FLUOCINONIDE, FLUOCINONIDE  
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE  
FLUORESCITE, FLUORESCIN SODIUM  
FLUOROPLEX, FLUOROURACIL  
FLUOROURACIL, FLUOROURACIL  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
FLUOXYMESTERONE, FLUOXYMESTERONE  
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE  
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE  
FLURANDRENOLIDE, FLURANDRENOLIDE  
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE  
FLURBIPROFEN, FLURBIPROFEN  
FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM  
FLUTAMIDE, FLUTAMIDE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
FLUVASTATIN SODIUM, FLUVASTATIN SODIUM  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
FML, FLUOROMETHOLONE  
FML FORTE, FLUOROMETHOLONE  
FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE  
FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE  
FOLIC ACID, FOLIC ACID  
FOLLISTIM AQ, FOLLITROPIN ALFA/BETA  
FOLOTYN, PRALATREXATE  
FOMEPIZOLE, FOMEPIZOLE  
FONDAPARINUX SODIUM, FONDAPARINUX SODIUM  
FORANE, ISOFLURANE  
FORFIVO XL, BUPROPION HYDROCHLORIDE  
FORTAMET, METFORMIN HYDROCHLORIDE  
FORTAZ, CEFTAZIDIME  
FORTAZ IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM  
FORTEO, TERIPARATIDE RECOMBINANT HUMAN  
FORTESTA, TESTOSTERONE  
FOSAMAX, ALENDRONATE SODIUM  
FOSAMAX PLUS D, ALENDRONATE SODIUM  
FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM  
FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE  
FOSCAVIR, FOSCARNET SODIUM  
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM  
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM  
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM  
FOSRENOL, LANTHANUM CARBONATE  
FRAGMIN, DALTEPARIN SODIUM  
FREAMINE HBC 6.9%, AMINO ACIDS  
FREAMINE III 10%, AMINO ACIDS  
FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS  
FREAMINE III 8.5%, AMINO ACIDS  
FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS  
FROVA, FROVATRIPTAN SUCCINATE  
FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE  
FULYZAQ, CROFELEMER  
FURADANTIN, NITROFURANTOIN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* F \*\*

FUROSEMIDE, FUROSEMIDE  
FUSILEV, LEVOLEUCOVORIN CALCIUM  
FUZEON, ENFUVIRTIDE  
FYAVOLV, ETHINYL ESTRADIOL  
FYCOMPA, PERAMPANEL

\*\* G \*\*

GABAPENTIN, GABAPENTIN  
GABITRIL, TIAGABINE HYDROCHLORIDE  
GABLOFEN, BACLOFEN  
GADAVIST, GADOBUTROL  
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE  
GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67  
GALZIN, ZINC ACETATE  
GANCICLOVIR, GANCICLOVIR  
GANCICLOVIR, GANCICLOVIR SODIUM  
GANCICLOVIR SODIUM, GANCICLOVIR SODIUM  
GANIRELIX ACETATE, GANIRELIX ACETATE  
GASTROCROM, CROMOLYN SODIUM  
GASTROGRAFIN, DIATRIZOATE MEGLUMINE  
GATIFLOXACIN, GATIFLOXACIN  
GATTEX KIT, TEDUGLUTIDE RECOMBINANT  
GAVISCON, ALUMINUM HYDROXIDE (OTC)  
GELNIQUE, OXYBUTYNIN CHLORIDE  
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE  
GEMFIBROZIL, GEMFIBROZIL  
GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE  
GEMZAR, GEMCITABINE HYDROCHLORIDE  
GEN-XENE, CLORAZEPATE DIPOTASSIUM  
GENERLAC, LACTULOSE  
GENGRAF, CYCLOSPORINE  
GENOPTIC, GENTAMICIN SULFATE  
GENOTROPIN, SOMATROPIN RECOMBINANT  
GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT  
GENTAK, GENTAMICIN SULFATE  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE  
GENVOYA, COBICISTAT  
GEODON, ZIPRASIDONE HYDROCHLORIDE  
GEODON, ZIPRASIDONE MESYLATE  
GIAPREZA, ANGIOTENSIN II  
GIAZO, BALSALAZIDE DISODIUM  
GILDAGIA, ETHINYL ESTRADIOL  
GILDESS 1.5/30, ETHINYL ESTRADIOL  
GILDESS 1/20, ETHINYL ESTRADIOL  
GILDESS 24 FE, ETHINYL ESTRADIOL  
GILDESS FE 1.5/30, ETHINYL ESTRADIOL  
GILDESS FE 1/20, ETHINYL ESTRADIOL  
GILENYA, FINGOLIMOD  
GILOTrif, AFATINIB DIMALEATE  
GLATIRAMER ACETATE, GLATIRAMER ACETATE  
GLATOPA, GLATIRAMER ACETATE  
GLEEVEC, IMATINIB MESYLATE  
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE  
GLEOSTINE, LOMUSTINE  
GLIADEL, CARMUSTINE  
GLIMEPIRIDE, GLIMEPIRIDE  
GLIPIZIDE, GLIPIZIDE  
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE  
GLOFIL-125, IOTHALAMATE SODIUM I-125  
GLUCAGEN, GLUCAGON HYDROCHLORIDE RECOMBINANT  
GLUCAGON, GLUCAGON HYDROCHLORIDE  
GLUCAGON, GLUCAGON RECOMBINANT  
GLUCAMIDE, CHLORPROPAMIDE  
GLUCOPHAGE, METFORMIN HYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* G \*\*

GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE  
 GLUCOTROL, GLIPIZIDE  
 GLUCOTROL XL, GLIPIZIDE  
 GLUMETZA, METFORMIN HYDROCHLORIDE  
 GLYBURIDE, GLYBURIDE  
 GLYBURIDE (MICRONIZED), GLYBURIDE  
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE  
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE  
 GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)  
 GLYCOLAX, POLYETHYLENE GLYCOL 3350  
 GLCOPYRROLATE, GLCOPYRROLATE  
 GLYDO, LIDOCAINE HYDROCHLORIDE  
 GLYNASE, GLYBURIDE  
 GLYSET, MIGLITOL  
 GLYXAMBI, EMPAGLIFLOZIN  
 GOCOVRI, AMANTADINE HYDROCHLORIDE  
 GOLYTELY, POLYETHYLENE GLYCOL 3350  
 GONAL-F, FOLLITROPIN ALFA/BETA  
 GONAL-F RFF, FOLLITROPIN ALFA/BETA  
 GONAL-F RFF REDI-JECT, FOLLITROPIN ALFA/BETA  
 GONITRO, NITROGLYCERIN  
 GOPRELTO, COCAINE HYDROCHLORIDE  
 GRALISE, GABAPENTIN  
 GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE  
 GRANisetron HYDROCHLORIDE PRESERVATIVE FREE, GRANisetron HYDROCHLORIDE  
 GRIS-PEG, GRiseofulvin, ULTRAMICROSIZE  
 GRiseofulvin, GRiseofulvin, MICROSIZE  
 GRiseofulvin, ULTRAMICROSIZE, GRiseofulvin, ULTRAMICROSIZE  
 GRiseofulvin, ULTRAMICROSIZE, GRiseofulvin, ULTRAMICROSIZE  
 GUAIFENESIN, GUAIFENESIN (OTC)  
 GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)  
 GUAIFENESIN AND PSEUDOEPHENDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)  
 GUANABENZ ACETATE, GUANABENZ ACETATE  
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE  
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE  
 GYNAZOLE-1, BUTOCONAZOLE NITRATE

\*\* H \*\*

H.P. ACTHAR GEL, CORTICOTROPIN  
 HABITROL, NICOTINE (OTC)  
 HAILEY FE 1/20, ETHINYLEDIESTRADIOL  
 HALAVEN, ERIBULIN MESYLATE  
 HALCION, TRIAZOLAM  
 HALDOL, HALOPERIDOL DECANOATE  
 HALDOL, HALOPERIDOL LACTATE  
 HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE  
 HALOG, HALCINONIDE  
 HALOPERIDOL, HALOPERIDOL  
 HALOPERIDOL, HALOPERIDOL LACTATE  
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE  
 HARVONI, LEDIPASVIR  
 HEATHER, NORETHINDRONE  
 HECTOROL, DOXERCALCIEROL  
 HEMABATE, CARBOPROST TROMETHAMINE  
 HEMANGEOL, PROPRANOLOL HYDROCHLORIDE  
 HEPARIN SODIUM, HEPARIN SODIUM  
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* H \*\*

HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM  
HEPATAMINE 8%, AMINO ACIDS  
HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT  
HEPSERA, ADEFOVIR DIPIVOXIL  
HER STYLE, LEVONORGESTREL (OTC)  
HETLIOZ, TASIMELTEON  
HEXALEN, ALTRETAMINE  
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)  
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)  
HICON, SODIUM IODIDE I-131  
HIPREX, METHENAMINE HIPPURATE  
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE  
HORIZANT, GABAPENTIN ENACARBIL  
HUMALOG, INSULIN LISPRO RECOMBINANT  
HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT  
HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT  
HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT  
HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT  
HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT  
HUMATROPE, SOMATROPIN RECOMBINANT  
HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)  
HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)  
HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)  
HUMULIN R, INSULIN HUMAN  
HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)  
HUMULIN R KWIKPEN, INSULIN HUMAN  
HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)  
HYCAMTIN, TOPOTECAN HYDROCHLORIDE  
HYCOFENIX, GUAIFENESIN  
HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE  
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE  
HYDREA, HYDROXYUREA  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE  
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE  
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE  
HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE, HYDROCODONE BITARTRATE  
HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,  
HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,  
HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX  
HYDROCODONE POLISTIREX AND CHLORPHENIRAMNE POLISTIREX, CHLORPHENIRAMINE POLISTIREX  
HYDROCORTISONE, HYDROCORTISONE  
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL  
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE  
HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE  
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE  
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE  
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN  
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE  
HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE  
HYDROXYUREA, HYDROXYUREA  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE  
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN  
HYSINGLA, HYDROCODONE BITARTRATE  
HYZAAR, HYDROCHLOROTHIAZIDE

\*\* I \*\*

IBANDRONATE SODIUM, IBANDRONATE SODIUM  
IBRANCE, PALBOCICLIB  
IBU-TAB, IBUPROFEN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* I \*\*

IBU-TAB 200, IBUPROFEN (OTC)  
IBUPROFEN, IBUPROFEN (OTC)  
IBUPROFEN, IBUPROFEN  
IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)  
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)  
IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)  
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)  
IBUPROFEN LYSINE, IBUPROFEN LYSINE  
IBUPROFEN SODIUM, IBUPROFEN SODIUM (OTC)  
IBUPROHM, IBUPROFEN (OTC)  
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)  
IBUTILIDE FUMARATE, IBUTILIDE FUMARATE  
IC-GREEN, INDOCYANINE GREEN  
ICLUSIG, PONATINIB HYDROCHLORIDE  
IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE  
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE  
IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE  
IDHIFA, ENASIDENIB MESYLATE  
IDKIT:HP, CITRIC ACID  
IFEX, IFOSFAMIDE  
IFOSFAMIDE, IFOSFAMIDE  
ILEVRO, NEPAFENAC  
ILOPERIDONE, ILOPERIDONE  
ILUVIEN, FLUOCINOLONE ACETONIDE  
IMATINIB MESYLATE, IMATINIB MESYLATE  
IMBRUVICA, IBRUTINIB  
IMIPENEM AND CILASTATIN, CILASTATIN SODIUM  
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE  
IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE  
IMIQUIMOD, IMIQUIMOD  
IMITREX, SUMATRIPTAN  
IMITREX, SUMATRIPTAN SUCCINATE  
IMITREX STATDOSE, SUMATRIPTAN SUCCINATE  
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)  
IMODIUM MULTI-SYMPMOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)  
IMPAVIDO, MILTEFOSINE  
IMPOYZ, CLOBETASOL PROPIONATE  
IMURAN, AZATHIOPRINE  
INAPSINE, DROPERIDOL  
INCASSIA, NORETHINDRONE  
INCRELEX, MECASERMIN RECOMBINANT  
INCRUSE ELLIPTA, UMECLIDINIUM BROMIDE  
INDAPAMIDE, INDAPAMIDE  
INDERAL LA, PROPRANOLOL HYDROCHLORIDE  
INDICLOR, INDIUM IN-111 CHLORIDE  
INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE  
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE  
INDOCIN, INDOMETHACIN  
INDOCIN, INDOMETHACIN SODIUM  
INDOCYANINE GREEN, INDOCYANINE GREEN  
INDOMETHACIN, INDOMETHACIN  
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM  
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)  
INFASURF PRESERVATIVE FREE, CALFACTANT  
INFED, IRON DEXTRAN  
INFUMORPH, MORPHINE SULFATE  
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE  
INFUVITE PEDIATRIC, ASCORBIC ACID  
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID  
INGREZZA, VALBENAZINE TOSYLATE  
INJECTAFER, FERRIC CARBOXYMALTOSE  
INLYTA, AXITINIB  
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE  
INOMAX, NITRIC OXIDE  
INSPRA, EPLERENONE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* I \*\*

INTEGRILIN, EPTIFIBATIDE  
INTELENCE, ETRAVIRINE  
INTERMEZZO, ZOLPIDEM TARTRATE  
INTRALIPID 10%, SOYBEAN OIL  
INTRALIPID 20%, SOYBEAN OIL  
INTRALIPID 30%, SOYBEAN OIL  
INTRAROSA, PRASTERONE  
INTROVALE, ETHINYLMESTRADIOL  
INTUNIV, GUANFACINE HYDROCHLORIDE  
INVANZ, ERТАPENEM SODIUM  
INVEGA, PALIPERIDONE  
INVEGA SUSTENNA, PALIPERIDONE PALMITATE  
INVEGA TRINZA, PALIPERIDONE PALMITATE  
INVIRASE, SAQUINAVIR MESYLATE  
INVOKAMET, CANAGLIFLOZIN  
INVOKAMET XR, CANAGLIFLOZIN  
INVOKANA, CANAGLIFLOZIN  
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
IONSYS, FENTANYL HYDROCHLORIDE  
IOPIDINE, APRACLONIDINE HYDROCHLORIDE  
IOSAT, POTASSIUM IODIDE (OTC)  
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE  
IPRIVASK, DESIRUDIN RECOMBINANT  
IRBESARTAN, IRBESARTAN  
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
IRESSA, GEFITINIB  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
ISENTRESS, RALTEGRAVIR POTASSIUM  
ISENTRESS HD, RALTEGRAVIR POTASSIUM  
ISIBLOOM, DESOGESTREL  
ISOCaine HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE  
ISOFLURANE, ISOFLURANE  
ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
ISONIAZID, ISONIAZID  
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE  
ISOPTO ATROPINE, ATROPINE SULFATE  
ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE  
ISORDIL, ISOSORBIDE DINITRATE  
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE  
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE  
ISOSULFAN BLUE, ISOSULFAN BLUE  
ISOTRETINOIN, ISOTRETINOIN  
ISOVUE-200, IOPAMIDOL  
ISOVUE-250, IOPAMIDOL  
ISOVUE-300, IOPAMIDOL  
ISOVUE-370, IOPAMIDOL  
ISOVUE-M 200, IOPAMIDOL  
ISOVUE-M 300, IOPAMIDOL  
ISRADIPINE, ISRADIPINE  
ISTALOL, TIMOLOL MALEATE  
ISTODAX, ROMIDEPSIN  
ISUPREL, ISOPROTERENOL HYDROCHLORIDE  
ITRACONAZOLE, ITRACONAZOLE  
IVERMECTIN, IVERMECTIN  
IVY BLOCK, BENTOQUATAM (OTC)  
IXEMPRA KIT, IXABEPILONE

\*\* J \*\*

JADENU, DEFERASIROX  
JADENU SPRINKLE, DEFERASIROX  
JAIMESS, ETHINYLMESTRADIOL  
JAKAFI, RUXOLITINIB PHOSPHATE  
JALYN, DUTASTERIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* J \*\*

JANTOVEN, WARFARIN SODIUM  
JANUMET, METFORMIN HYDROCHLORIDE  
JANUMET XR, METFORMIN HYDROCHLORIDE  
JANUVIA, SITAGLIPTIN PHOSPHATE  
JARDIANCE, EMPAGLIFLOZIN  
JEANATOPE, ALBUMIN IODINATED I-125 SERUM  
JENCYCLLA, NORETHINDRONE  
JENTADUETO, LINAGLIPTIN  
JENTADUETO XR, LINAGLIPTIN  
JEVTANA KIT, CABAZITAXEL  
JUBLIA, EFINACONAZOLE  
JULUCA, DOLUTEGRAVIR SODIUM  
JUNEL 1.5/30, ETHINYLMESTRADIOL  
JUNEL 1/20, ETHINYLMESTRADIOL  
JUNEL FE 1.5/30, ETHINYLMESTRADIOL  
JUNEL FE 1/20, ETHINYLMESTRADIOL  
JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)  
JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)  
JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)  
JUXTAPID, LOMITAPIDE MESYLATE

\*\* K \*\*

K-TAB, POTASSIUM CHLORIDE  
KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS  
KADIAN, MORPHINE SULFATE  
KAITLIB FE, ETHINYLMESTRADIOL  
KALETRA, LOPINAVIR  
KALEXATE, SODIUM POLYSTYRENE SULFONATE  
KALLIGA, DESOGESTREL  
KALYDECO, IVACAFTOR  
KANAMYCIN SULFATE, KANAMYCIN SULFATE  
KAPVAY, CLONIDINE HYDROCHLORIDE  
KARBINAL ER, CARBINOXAMINE MALEATE  
KARIVA, DESOGESTREL  
KAZANO, ALOGLIPTIN BENZOATE  
KEFLEX, CEPHALEXIN  
KEFZOL, CEFAZOLIN SODIUM  
KELNOR, ETHINYLMESTRADIOL  
KENALOG, TRIAMCINOLONE ACETONIDE  
KENALOG-10, TRIAMCINOLONE ACETONIDE  
KENALOG-40, TRIAMCINOLONE ACETONIDE  
KENGREAL, CANGRELOR  
KEPPRA, LEVETIRACETAM  
KEPPRA XR, LEVETIRACETAM  
KERYDIN, TAVABOROLE  
KETALAR, KETAMINE HYDROCHLORIDE  
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE  
KETOCONAZOLE, KETOCONAZOLE  
KETOPROFEN, KETOPROFEN  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)  
KETOZOLE, KETOCONAZOLE  
KEVEYIS, DICHLORPHENAMIDE  
KHEDEZLA, DESVENLAFAKINE  
KIMIDESS, DESOGESTREL  
KINEVAC, SINCALIDE  
KIONEX, SODIUM POLYSTYRENE SULFONATE  
KISQALI, RIBOCICLIB SUCCINATE  
KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE  
KITABIS PAK, TOBRAMYCIN  
KLARON, SULFACETAMIDE SODIUM  
KLONOPIN, CLONAZEPAM  
KLOR-CON, POTASSIUM CHLORIDE  
KLOR-CON M10, POTASSIUM CHLORIDE  
KLOR-CON M15, POTASSIUM CHLORIDE

**APPENDIX A - PRODUCT NAME INDEX****\*\* K \*\***

KLOR-CON M20, POTASSIUM CHLORIDE  
KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE  
KORLYM, MIFEPRISTONE  
KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE  
KURVELO, ETHINYLMESTRADIOL  
KUVAN, SAPROPTERIN DIHYDROCHLORIDE  
KYBELLA, DEOXYCHOLIC ACID  
KYLEENA, LEVONORGESTREL  
KYNAMRO, MIPOMERSEN SODIUM  
KYPROLIS, CARFILZOMIB

**\*\* L \*\***

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
LACRISERT, HYDROXYPROPYL CELLULOSE  
LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
LACTULOSE, LACTULOSE  
LAMICTAL, LAMOTRIGINE  
LAMICTAL CD, LAMOTRIGINE  
LAMICTAL ODT, LAMOTRIGINE  
LAMICTAL XR, LAMOTRIGINE  
LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)  
LAMISIL, TERBINAFINE HYDROCHLORIDE  
LAMISIL AT, TERBINAFINE (OTC)  
LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)  
LAMIVUDINE, LAMIVUDINE  
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE  
LAMOTRIGINE, LAMOTRIGINE  
LANIAZID, ISONIAZID  
LANORINAL, ASPIRIN  
LANOXIN, DIGOXIN  
LANOXIN PEDIATRIC, DIGOXIN  
LANSOPRAZOLE, LANSOPRAZOLE (OTC)  
LANSOPRAZOLE, LANSOPRAZOLE  
LANSOPRAZOLE, LANSOPRAZOLE  
LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN  
LANTHANUM CARBONATE, LANTHANUM CARBONATE  
LANTUS, INSULIN GLARGINE RECOMBINANT  
LANTUS SOLOSTAR, INSULIN GLARGINE RECOMBINANT  
LARIN 1.5/30, ETHINYLMESTRADIOL  
LARIN 1/20, ETHINYLMESTRADIOL  
LARIN 24 FE, ETHINYLMESTRADIOL  
LARIN FE 1.5/30, ETHINYLMESTRADIOL  
LARIN FE 1/20, ETHINYLMESTRADIOL  
LAROTID, AMOXICILLIN  
LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE  
LASIX, FUROSEMIDE  
LASTACRAFT, ALCAFTADINE  
LATANOPROST, LATANOPROST  
LATISSE, BIMATOPROST  
LATUDA, LURASIDONE HYDROCHLORIDE  
LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350  
LAZANDA, FENTANYL CITRATE  
LEFLUNOMIDE, LEFLUNOMIDE  
LENVIMA, LENVATINIB MESYLATE  
LERIBANE, ETHINYLMESTRADIOL  
LESCOL XL, FLUVASTATIN SODIUM  
LESSINA-28, ETHINYLMESTRADIOL  
LETAIRIS, AMBRISENTAN  
LETROZOLE, LETROZOLE  
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM  
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM  
LEUKERAN, CHLORAMBUCIL  
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE  
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* L \*\*

LEVAQUIN, LEVOFLOXACIN  
LEVEMIR, INSULIN DETEMIR RECOMBINANT  
LEVEMIR FLEXTOUCH, INSULIN DETEMIR RECOMBINANT  
LEVETIRACETAM, LEVETIRACETAM  
LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM  
LEVITRA, VARDENAFIL HYDROCHLORIDE  
LEVO-T, LEVOTHYROXINE SODIUM \*\*  
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE  
LEVOCARNITINE, LEVOCARNITINE  
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE  
LEVOFLOXACIN, LEVOFLOXACIN  
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN  
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM  
LEVONEST, ETHINYLMESTRADIOL  
LEVONORGESTREL, LEVONORGESTREL (OTC)  
LEVONORGESTREL, LEVONORGESTREL  
LEVONORGESTREL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL  
LEVONORGESTREL AND ETHINYLMESTRADIOL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL  
LEVOPHED, NOREPINEPHRINE BITARTRATE  
LEVORA 0.15/30-28, ETHINYLMESTRADIOL  
LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE  
LEVOHYROXINE SODIUM, LEVOHYROXINE SODIUM  
LEVOHYROXINE SODIUM, LEVOHYROXINE SODIUM \*\*  
LEVOXYL, LEVOHYROXINE SODIUM \*\*  
LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE  
LEXAPRO, ESCITALOPRAM OXALATE  
LEXISCAN, REGADENOSON  
LEXIVA, FOSAMPRENAVIR CALCIUM  
LIALDA, MESALAMINE  
LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE  
LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE  
LIDEX, FLUOCINONIDE  
LIDOCAINE, LIDOCAINE  
LIDOCAINE AND PRILOCaine, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE  
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE  
LIDODERM, LIDOCAINE  
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE  
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE  
LILETTA, LEVONORGESTREL  
LINCOCIN, LINCOMYCIN HYDROCHLORIDE  
LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE  
LINDANE, LINDANE  
LINEZOLID, LINEZOLID  
LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID  
LINZESS, LINACLOTIDE  
LIORESAL, BACLOFEN  
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM  
LIPIODOL, ETHIODIZED OIL  
LIPITOR, ATORVASTATIN CALCIUM  
LIPOFEN, FENOFIBRATE  
LIQUID E-Z-PAQUE, BARIUM SULFATE  
LISINOPRIL, LISINOPRIL  
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LITHIUM CARBONATE, LITHIUM CARBONATE  
LITHIUM CITRATE, LITHIUM CITRATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* L \*\*

LITHOBID, LITHIUM CARBONATE  
LITHOSTAT, ACETOHYDROXAMIC ACID  
LIVALO, PITAVASTATIN CALCIUM  
LO LOESTRIN FE, ETHINYL ESTRADIOL  
LO MINASTRIN FE, ETHINYL ESTRADIOL  
LO SIMPESSE, ETHINYL ESTRADIOL  
LOCOID, HYDROCORTISONE BUTYRATE  
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE  
LODOSYN, CARBIDOPA  
LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL  
LOESTRIN 21 1/20, ETHINYL ESTRADIOL  
LOESTRIN 24 FE, ETHINYL ESTRADIOL  
LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL  
LOESTRIN FE 1/20, ETHINYL ESTRADIOL  
LOGILIA, ULIPRISTAL ACETATE  
LOMAIRA, PHENTERMINE HYDROCHLORIDE  
LOMOTIL, ATROPINE SULFATE  
LONHALA MAGNAIR KIT, GLYCOPYRROLATE  
LONSURF, TIPIRACIL HYDROCHLORIDE  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE  
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)  
LOPID, GEMFIBROZIL  
LOPINAVIR AND RITONAVIR, LOPINAVIR  
LOPRESSOR, METOPROLOL TARTRATE  
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE  
LOPROX, CICLOPIROX  
LOPURIN, ALLOPURINOL  
LORATADINE, LORATADINE (OTC)  
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)  
LORATADINE REDIDOSE, LORATADINE (OTC)  
LORAZEPAM, LORAZEPAM  
LORAZEPAM INTENSOL, LORAZEPAM  
LORAZEPAM PRESERVATIVE FREE, LORAZEPAM  
LORYNA, DROSPIRENONE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LOSEASONIQUE, ETHINYL ESTRADIOL  
LOTEMAX, LOTEPREDNOL ETABONATE  
LOTENSIN, BENAZEPRIL HYDROCHLORIDE  
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE  
LOTREL, AMLODIPINE BESYLATE  
LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)  
LOTRISONE, BETAMETHASONE DIPROPIONATE  
LOTRONEX, ALOSETRON HYDROCHLORIDE  
LOVASTATIN, LOVASTATIN  
LOVAZA, OMEGA-3-ACID ETHYL ESTERS  
LOVENOX, ENOXAPARIN SODIUM  
LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM  
LOW-OGESTREL-28, ETHINYL ESTRADIOL  
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE  
LTA II KIT, LIDOCAINE HYDROCHLORIDE  
LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES  
LUMIFY, BRIMONIDINE TARTRATE (OTC)  
LUMIGAN, BIMATOPROST  
LUNESTA, ESZOPICLONE  
LUPANETA PACK, LEUPROLIDE ACETATE  
LUPRON DEPOT, LEUPROLIDE ACETATE  
LUPRON DEPOT-PED, LEUPROLIDE ACETATE  
LUVOX, FLUVOXAMINE MALEATE  
LUXIQ, BETAMETHASONE VALERATE  
LUZU, LULICONAZOLE  
LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT  
LYNPARZA, OLAPARIB  
LYRICA, PREGABALIN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* L \*\*

LYRICA CR, PREGABALIN  
LYSODREN, MITOTANE  
LYSTEDA, TRANEXAMIC ACID

\*\* M \*\*

M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
M.V.I. ADULT, ASCORBIC ACID  
M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID  
M.V.I. PEDIATRIC, ASCORBIC ACID  
MACRILEN, MACIMORELIN ACETATE  
MACROBID, NITROFURANTOIN  
MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE  
MACUGEN, PEGAPANTIB SODIUM  
MAFENIDE ACETATE, MAFENIDE ACETATE  
MAGNESIUM SULFATE, MAGNESIUM SULFATE  
MAGNESIUM SULFATE, MAGNESIUM SULFATE  
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE  
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE  
MAGNEVIST, GADOPENTETATE DIMEGLUMINE  
MAKENA, HYDROXYPROGESTERONE CAPROATE  
MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE  
MALARONE, ATOVAQUONE  
MALARONE PEDIATRIC, ATOVAQUONE  
MALATHION, MALATHION  
MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE  
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 25%, MANNITOL  
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL  
MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE  
MARCAINE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE  
MARINOL, DRONABINOL  
MARLISSA, ETHINYLMESTRADIOL  
MARPLAN, ISOCARBOXAZID  
MARQIBO KIT, VINCRISTINE SULFATE  
MATULANE, PROCARBAZINE HYDROCHLORIDE  
MAVIK, TRANDOLAPRIL  
MAVYRET, GLECAPREVIR  
MAXALT, RIZATRIPTAN BENZOATE  
MAXALT-MLT, RIZATRIPTAN BENZOATE  
MAXIDEX, DEXAMETHASONE  
MAXIPIME, CEFEPIME HYDROCHLORIDE  
MAXITROL, DEXAMETHASONE  
MAXZIDE, HYDROCHLOROTHIAZIDE  
MAXZIDE-25, HYDROCHLOROTHIAZIDE  
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE  
MDP-BRACCO, TECHNETIUM TC-99M MEDRONATE KIT  
MECAMYLAMINE HYDROCHLORIDE, MECAMYLAMINE HYDROCHLORIDE  
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE  
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM  
MEDROL, METHYLPREDNISOLONE  
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE  
MEFENAMIC ACID, MEFENAMIC ACID  
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE  
MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM  
MEGACE ES, MEGESTROL ACETATE  
MEGATOPE, ALBUMIN IODINATED I-131 SERUM  
MEGESTROL ACETATE, MEGESTROL ACETATE  
MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE  
MELAMISA, DROSPIRENONONE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* M \*\*

MELOXICAM, MELOXICAM  
MELPHALAN, MELPHALAN  
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
MEMBRANEBLUE, TRYPLAN BLUE  
MEN'S ROGAINE, MINOXIDIL (OTC)  
MENEST, ESTROGENS, ESTERIFIED  
MENOPUR, MENOTROPINS (FSH  
MENOSTAR, ESTRADIOL  
MENTAX, BUTENAFINE HYDROCHLORIDE  
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE  
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE  
MEPHYTON, PHYTONADIONE  
MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE  
MEPROBAMATE, MEPROBAMATE  
MEPRON, ATOVAQUONE  
MERCAPTOPURINE, MERCAPTOPURINE  
MEROPENEM, MEROPENEM  
MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM  
MERREM, MEROPENEM  
MESALAMINE, MESALAMINE  
MESNA, MESNA  
MESNEX, MESNA  
MESTINON, PYRIDOSTIGMINE BROMIDE  
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE  
METADATE ER, METHYLPHENIDATE HYDROCHLORIDE  
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE  
METARAMINOL BITARTRATE, METARAMINOL BITARTRATE  
METASTRON, STRONTIUM CHLORIDE SR-89  
METAXALONE, METAXALONE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE  
METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE  
METHADOSE, METHADONE HYDROCHLORIDE  
METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE  
METHAZOLAMIDE, METHAZOLAMIDE  
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE  
METHERGINE, METHYLERGONOVINE MALEATE  
METHIMAZOLE, METHIMAZOLE  
METHOCARBAMOL, METHOCARBAMOL  
METHOCARBAMOL AND ASPIRIN, ASPIRIN  
METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM  
METHOTREXATE SODIUM, METHOTREXATE SODIUM  
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM  
METHOXSALEN, METHOXSALEN  
METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE  
METHYCLOTHIAZIDE, METHYCLOTHIAZIDE  
METHYLDOPA, METHYLDOPA  
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE  
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE  
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE  
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE  
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
METHYLPREDNISOLONE, METHYLPREDNISOLONE  
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE  
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE  
METHYLTESTOSTERONE, METHYLTESTOSTERONE  
METIPRANOLOL, METIPRANOLOL HYDROCHLORIDE  
METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE  
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
METOLAZONE, METOLAZONE  
METOPIRONE, METYRAPONE  
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* M \*\*

METOPROLOL TARTRATE, METOPROLOL TARTRATE  
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
METOZOLOV ODT, METOCLOPRAMIDE HYDROCHLORIDE  
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE  
METROCREAM, METRONIDAZOLE  
METROGEL, METRONIDAZOLE  
METROGEL-VAGINAL, METRONIDAZOLE  
METROLOTION, METRONIDAZOLE  
METRONIDAZOLE, METRONIDAZOLE  
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE  
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE  
MIACALCIN, CALCITONIN SALMON  
MIBELAS 24 FE, ETHINYL ESTRADIOL  
MICARDIS, TELMISARTAN  
MICARDIS HCT, HYDROCHLOROTHIAZIDE  
MICONAZOLE 3, MICONAZOLE NITRATE (OTC)  
MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
MICONAZOLE 7, MICONAZOLE NITRATE (OTC)  
MICONAZOLE 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)  
MICONAZOLE NITRATE, MICONAZOLE NITRATE  
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
MICORT-HC, HYDROCORTISONE ACETATE  
MICRO-K, POTASSIUM CHLORIDE  
MICRO-K 10, POTASSIUM CHLORIDE  
MICROGESTIN 1.5/30, ETHINYL ESTRADIOL  
MICROGESTIN 1/20, ETHINYL ESTRADIOL  
MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL  
MICROGESTIN FE 1/20, ETHINYL ESTRADIOL  
MICRONOR, NORETHINDRONE  
MICROZIDE, HYDROCHLOROTHIAZIDE  
MIDAMOR, AMILORIDE HYDROCHLORIDE  
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE  
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE  
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE  
MIDOL LIQUID GELS, IBUPROFEN (OTC)  
MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE  
MIFEPREX, MIFEPRISTONE  
MIGERGOT, CAFFEINE  
MIGLITOL, MIGLITOL  
MIGRANAL, DIHYDROERGOTAMINE MESYLATE  
MILI, ETHINYL ESTRADIOL  
MILNACIPRAN HYDROCHLORIDE, MILNACIPRAN HYDROCHLORIDE  
MILRINONE LACTATE, MILRINONE LACTATE  
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE  
MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE  
MINASTRIN 24 FE, ETHINYL ESTRADIOL  
MINIPRESS, PRAZOSIN HYDROCHLORIDE  
MINIRIN, DESMOPRESSIN ACETATE  
MINITRAN, NITROGLYCERIN  
MINIVELLE, ESTRADIOL  
MINOCIN, MINOCYCLINE HYDROCHLORIDE  
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE  
MINOLIRA, MINOCYCLINE HYDROCHLORIDE  
MINOXIDIL, MINOXIDIL (OTC)  
MINOXIDIL, MINOXIDIL  
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)  
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)  
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)  
MIOCHOL-E, ACETYLCHOLINE CHLORIDE  
MIOSTAT, CARBACHOL  
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)  
MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE  
MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* M \*\*

MIRENA, LEVONORGESTREL  
MIRTAZAPINE, MIRTAZAPINE  
MIRVASO, BRIMONIDINE TARTRATE  
MISOPROSTOL, MISOPROSTOL  
MITIGARE, COLCHICINE  
MITOMYCIN, MITOMYCIN  
MITOSOL, MITOMYCIN  
MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE  
MIVACRON, MIVACURUM CHLORIDE  
MOBIC, MELOXICAM  
MODAFINIL, MODAFINIL  
MODICON 28, ETHINYLMESTRADIOL  
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE  
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE  
MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE  
MOMETASONE FUROATE, MOMETASONE FUROATE  
MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
MONISTAT 3, MICONAZOLE NITRATE (OTC)  
MONISTAT 3, MICONAZOLE NITRATE  
MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)  
MONISTAT 7, MICONAZOLE NITRATE (OTC)  
MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
MONO-LINYAH, ETHINYLMESTRADIOL  
MONODOX, DOXYCYCLINE  
MONOKET, ISOSORBIDE MONONITRATE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
MONUROL, FOSFOMYCIN TROMETHAMINE  
MORPHABOND ER, MORPHINE SULFATE  
MORPHINE SULFATE, MORPHINE SULFATE  
MOTOFEN, ATROPINE SULFATE  
MOTRIN IB, IBUPROFEN (OTC)  
MOVANTIK, NALOXEGOL OXALATE  
MOVIPREP, ASCORBIC ACID  
MOXATAG, AMOXICILLIN  
MOXEZA, MOXIFLOXACIN HYDROCHLORIDE  
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE  
MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN  
MOZOBIL, PLERIXAFOR  
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM  
MS CONTIN, MORPHINE SULFATE  
MUCINEX, GUAIFENESIN (OTC)  
MUCINEX D, GUAIFENESIN (OTC)  
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)  
MULTAQ, DRONEDARONE HYDROCHLORIDE  
MULTIHANCE, GADOBENATE DIMEGLUMINE  
MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE  
MUPIROCIN, MUPIROCIN  
MUPIROCIN, MUPIROCIN CALCIUM  
MUSE, ALPROSTADIL  
MUSTARGEN, MECHLORETHAMINE HYDROCHLORIDE  
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE  
MYCAMINE, MICAFUNGIN SODIUM  
MYCELEX-7, CLOTrimazole (OTC)  
MYCELEX-7 COMBINATION PACK, CLOTrimazole (OTC)  
MYCOBUTIN, RIFABUTIN  
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL  
MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE  
MYCOPHENOLIC ACID, MYCOPHENOLIC ACID  
MYDAYIS, AMPHETAMINE ASPARTATE  
MYDRIACYL, TROPICAMIDE  
MYFORTIC, MYCOPHENOLIC ACID  
MYKACET, NYSTATIN  
MYLERAN, BUSULFAN  
MYORISAN, ISOTRETINOIN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* M \*\*

MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT

MYRBETRIQ, MIRABEGRON

MYSOLINE, PRIMIDONE

MYZILRA, ETHINYL ESTRADIOL

\*\* N \*\*

NABUMETONE, NABUMETONE

NADOLOL, NADOLOL

NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE

NAFCILLIN SODIUM, NAFCILLIN SODIUM

NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE

NAFTIN, NAFTIFINE HYDROCHLORIDE

NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE

NALFON, FENOPROFEN CALCIUM

NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM

NALOXONE, NALOXONE HYDROCHLORIDE

NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE

NAMENDA, MEMANTINE HYDROCHLORIDE

NAMENDA XR, MEMANTINE HYDROCHLORIDE

NAMZARIC, DONEPEZIL HYDROCHLORIDE

NANDROLONE DECANOATE, NANDROLONE DECANOATE

NAPHAZOLINE HYDROCHLORIDE, NAPHAZOLINE HYDROCHLORIDE

NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)

NAPHCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

NAPRELAN, NAPROXEN SODIUM

NAPROSYN, NAPROXEN

NAPROXEN, NAPROXEN

NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM

NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

NAPROXEN SODIUM, NAPROXEN SODIUM

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)

NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE

NARCAN, NALOXONE HYDROCHLORIDE

NARDIL, PHENELZINE SULFATE

NAROPIN, ROPIVACAINE HYDROCHLORIDE

NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)

NASCOBAL, CYANOCOBALAMIN

NASONEX, MOMETASONE FUROATE

NATACYN, NATAMYCIN

NATAZIA, DIENOGENEST

NATEGLINIDE, NATEGLINIDE

NATESTO, TESTOSTERONE

NATRECOR, NESIRITIDE RECOMBINANT

NATROBA, SPINOSAD

NAVELBINE, VINORELBINE TARTRATE

NEBUPENT, PENTAMIDINE ISETHIONATE

NEDOCROMIL SODIUM, NEDOCROMIL SODIUM

NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE

NEMBUTAL SODIUM, PENTOBARBITAL SODIUM

NEO-SYNALAR, FLUOCINOLONE ACETONIDE

NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC

NEOMYCIN SULFATE, NEOMYCIN SULFATE

NEOPAP, ACETAMINOPHEN (OTC)

NEOPROFEN, IBUPROFEN LYSINE

NEORAL, CYCLOSPORINE

NEOSPORIN, BACITRACIN ZINC

NEOSPORIN, GRAMICIDIN

NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* N \*\*

NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE  
NEPHRAMINE 5.4%, AMINO ACIDS  
NERLYNX, NERATINIB MALEATE  
NESACAINE, CHLOROPROCAINE HYDROCHLORIDE  
NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE  
NESINA, ALOGLIPTIN BENZOATE  
NETSPOT, GALLIUM DOTATATE GA-68  
NEUPRO, ROTIGOTINE  
NEURACEQ, FLORBETABEN F-18  
NEUROLITE, TECHNETIUM TC-99M BICISATE KIT  
NEURONTIN, GABAPENTIN  
NEVANAC, NEPAFENAC  
NEVIRAPINE, NEVIRAPINE  
NEXAVAR, SORAFENIB TOSYLATE  
NEXESTA FE, ETHINYLMESTRADIOL  
NEXIUM, ESOMEPRAZOLE MAGNESIUM  
NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)  
NEXIUM IV, ESOMEPRAZOLE SODIUM  
NEXPLANON, ETONOGESTREL  
NEXTERONE, AMIODARONE HYDROCHLORIDE  
NIACIN, NIACIN  
NIACOR, NIACIN  
NIASPAN, NIACIN  
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE  
NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE  
NICODERM CQ, NICOTINE (OTC)  
NICORETTE, NICOTINE POLACRILEX (OTC)  
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)  
NICOTINE, NICOTINE (OTC)  
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)  
NICOTROL, NICOTINE  
NIFEDIPINE, NIFEDIPINE  
NIKKI, DROSPIRENONE  
NILANDRON, NILUTAMIDE  
NILUTAMIDE, NILUTAMIDE  
NIMBEX, CISATRACURIUM BESYLATE  
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE  
NIMODIPINE, NIMODIPINE  
NINLARO, IXAZOMIB CITRATE  
NIPENT, PENTOSTATIN  
NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE  
NISOLDIPINE, NISOLDIPINE  
NITHIODOTE, SODIUM NITRITE  
NITRO-DUR, NITROGLYCERIN  
NITROFURANTOIN, NITROFURANTOIN  
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE  
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN  
NITROGLYCERIN, NITROGLYCERIN  
NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN  
NITROLINGUAL PUMPSPRAY, NITROGLYCERIN  
NITROMIST, NITROGLYCERIN  
NITROPRESS, SODIUM NITROPRUSSIDE  
NITROSTAT, NITROGLYCERIN  
NITYR, NITISINONE  
NIX, PERMETHRIN (OTC)  
NIZATIDINE, NIZATIDINE  
NIZORAL, KETOCONAZOLE  
NIZORAL A-D, KETOCONAZOLE (OTC)  
NOCTIVA, DESMOPRESSIN ACETATE  
NOR-QD, NORETHINDRONE  
NORCO, ACETAMINOPHEN  
NORDITROPIN FLEXPRO, SOMATROPIN RECOMBINANT  
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE  
NORETHINDRONE, NORETHINDRONE  
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* N \*\*

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE,  
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL  
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL  
NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL  
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL  
NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL  
NORINYL 1+50 28-DAY, MESTRANOL  
NORITATE, METRONIDAZOLE  
NORMOCARB HF 25, MAGNESIUM CHLORIDE  
NORMOCARB HF 35, MAGNESIUM CHLORIDE  
NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
NORPACE, DISOPYRAMIDE PHOSPHATE  
NORPACE CR, DISOPYRAMIDE PHOSPHATE  
NORPRAMIN, DESIPRAMINE HYDROCHLORIDE  
NORTHERA, DROXIDOPA  
NORTREL 0.5/35-28, ETHINYL ESTRADIOL  
NORTREL 1/35-21, ETHINYL ESTRADIOL  
NORTREL 1/35-28, ETHINYL ESTRADIOL  
NORTREL 7/7/7, ETHINYL ESTRADIOL  
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE  
NORVASC, AMLODIPINE BESYLATE  
NORVIR, RITONAVIR  
NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)  
NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)  
NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)  
NOVOLOG, INSULIN ASPART RECOMBINANT  
NOVOLOG FLEXPEN, INSULIN ASPART RECOMBINANT  
NOVOLOG FLEXTOUCH, INSULIN ASPART RECOMBINANT  
NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT  
NOVOLOG MIX 70/30 FLEXPEN, INSULIN ASPART PROTAMINE RECOMBINANT  
NOVOLOG PENFILL, INSULIN ASPART RECOMBINANT  
NOXAFILE, POSACONAZOLE  
NUCYNTA, TAPENTADOL HYDROCHLORIDE  
NUCYNTA ER, TAPENTADOL HYDROCHLORIDE  
NUEDEXTA, DEXTROMETHORPHAN HYDROBROMIDE  
NULYTELY, POLYETHYLENE GLYCOL 3350  
NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350  
NUPLAZID, PIMAVANSERIN TARTRATE  
NUTRESTORE, L-GLUTAMINE  
NUTRILIPID 10%, SOYBEAN OIL  
NUTRILIPID 20%, SOYBEAN OIL  
NUTROPIN AQ NUSPIN, SOMATROPIN RECOMBINANT  
NUTROPIN AQ PEN, SOMATROPIN RECOMBINANT  
NUVARING, ETHINYL ESTRADIOL  
NUVESSA, METRONIDAZOLE  
NUVIGIL, ARMODAFINIL  
NYLIA 1/35, ETHINYL ESTRADIOL  
NYLIA 7/7/7, ETHINYL ESTRADIOL  
NYMALIZE, NIMODIPINE  
NYSTATIN, NYSTATIN  
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN  
NYSTOP, NYSTATIN

\*\* O \*\*

OCALIVA, OBETICHOLIC ACID  
OCTOCAINE, EPINEPHRINE  
OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT  
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE  
OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE  
OCUFEN, FLURBIPROFEN SODIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* O \*\*

OCUFLOX, OFLOXACIN  
ODEFSEY, EMTRICITABINE  
ODOMZO, SONIDEGIB PHOSPHATE  
OFEV, NINTEDANIB ESYLATE  
OFIRMEV, ACETAMINOPHEN  
OFLOXACIN, OFLOXACIN  
OGEN 5, ESTROPIPATE  
OGESTREL 0.5/50-28, ETHINYL ESTRADIOL  
OLANZAPINE, OLANZAPINE  
OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL  
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE  
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE  
OLUX, CLOBETASOL PROPIONATE  
OLUX E, CLOBETASOL PROPIONATE  
OLYSIO, SIMEPREVIR SODIUM  
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS  
OMEPRAZOLE, OMEPRAZOLE (OTC)  
OMEPRAZOLE, OMEPRAZOLE  
OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN  
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)  
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE  
OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)  
OMIDRIA, KETOROLAC TROMETHAMINE  
OMNARIS, CICLESONIDE  
OMNIPAQUE 140, IOHESOL  
OMNIPAQUE 180, IOHESOL  
OMNIPAQUE 240, IOHESOL  
OMNIPAQUE 300, IOHESOL  
OMNIPAQUE 350, IOHESOL  
OMNIPRED, PREDNISOLONE ACETATE  
OMNISCAN, GADODIAMIDE  
OMNITROPE, SOMATROPIN RECOMBINANT  
ONDANSETRON, ONDANSETRON  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONEXTON, BENZOYL PEROXIDE  
ONFI, CLOBAZAM  
ONGLYZA, SAXAGLIPTIN HYDROCHLORIDE  
ONIVYDE, IRINOTECAN HYDROCHLORIDE  
ONMEL, ITRACONAZOLE  
ONZETRA XSAIL, SUMATRIPTAN SUCCINATE  
OPANA, OXYMORPHONE HYDROCHLORIDE  
OPCICON ONE-STEP, LEVONORGESTREL (OTC)  
OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)  
OPSUMIT, MACITENTAN  
OPTIMARK, GADOVERSETAMIDE  
OPTIMARK IN PLASTIC CONTAINER, GADOVERSETAMIDE  
OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE  
OPTIRAY 240, IOVERSOL  
OPTIRAY 300, IOVERSOL  
OPTIRAY 320, IOVERSOL  
OPTIRAY 350, IOVERSOL  
OPTISON, ALBUMIN HUMAN  
ORABLOC, ARTICAINE HYDROCHLORIDE  
ORACEA, DOXYCYCLINE  
ORALTAG, IOHESOL  
ORAP, PIMOZIDE  
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE  
ORAQIX, LIDOCAINE  
ORAVERSE, PHENTOLAMINE MESYLATE  
ORAVIG, MICONAZOLE  
ORBACTIV, ORITAVANCIN DIPHOSPHATE  
ORENITRAM, TREPROSTINIL DIOLAMINE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* O \*\*

ORFADIN, NITISINONE  
ORKAMBI, IVACAFTOR  
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE  
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN  
ORSYTHIA, ETHINYLMESTRADIOL  
ORTHO CYCLEN-28, ETHINYLMESTRADIOL  
ORTHO TRI-CYCLEN, ETHINYLMESTRADIOL  
ORTHO TRI-CYCLEN LO, ETHINYLMESTRADIOL  
ORTHO-NOVUM 1/35-28, ETHINYLMESTRADIOL  
ORTHO-NOVUM 7/7/7-28, ETHINYLMESTRADIOL  
ORVATEN, MIDODRINE HYDROCHLORIDE  
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE  
OSENI, ALOGLIPTIN BENZOATE  
OSMITROL 10% IN WATER, MANNITOL  
OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 15% IN WATER, MANNITOL  
OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 20% IN WATER, MANNITOL  
OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 5% IN WATER, MANNITOL  
OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS  
OSPHENA, OSPEMIFENE  
OTEZLA, APREMILAST  
OTICAIR, HYDROCORTISONE  
OTIPRIO, CIPROFLOXACIN  
OTOVEL, CIPROFLOXACIN HYDROCHLORIDE  
OTREXUP, METHOTREXATE  
OVIDE, MALATHION  
OVIDREL, CHORIOGONADOTROPIN ALFA  
OXACILLIN SODIUM, OXACILLIN SODIUM  
OXALIPLATIN, OXALIPLATIN  
OXANDRIN, OXANDROLONE  
OXANDROLONE, OXANDROLONE  
OXaprozin, OXaprozin  
OXAYDO, OXYCODONE HYDROCHLORIDE  
OXAZEPAM, OXAZEPAM  
OXCARBAZEPINE, OXCARBAZEPINE  
OXICONAZOLE NITRATE, OXICONAZOLE NITRATE  
OXISTAT, OXICONAZOLE NITRATE  
OXSORALEN-ULTRA, METHOXALEN  
OXTELLAR XR, OXCARBAZEPINE  
OXYBUTYNIN, OXYBUTYNIN  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE  
OXYCET, ACETAMINOPHEN  
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
OXYCODONE AND ASPIRIN, ASPIRIN  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN  
OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN  
OXYCONTIN, OXYCODONE HYDROCHLORIDE  
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE  
OXYTOCIN, OXYTOCIN  
OXYTROL, OXYBUTYNIN  
OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)  
OZEMPIC, SEMAGLUTIDE  
OZURDEX, DEXAMETHASONE

\*\* P \*\*

PACERONE, AMIODARONE HYDROCHLORIDE  
PACITAXEL, PACLITAXEL  
PACLITAXEL, PACLITAXEL  
PALIPERIDONE, PALIPERIDONE  
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE  
PAMELOR, NORTRIPTYLINE HYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* P \*\*

PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM  
PANCREAZE, PANCRELIPASE (AMYLASE)  
PANCURONIUM BROMIDE, PANCURONIUM BROMIDE  
PANDEL, HYDROCORTISONE PROBUTATE  
PANRETIN, ALITRETNINOIN  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM  
PARAGARD T 380A, COPPER  
PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE  
PARICALCITOL, PARICALCITOL  
PARLODEL, BROMOCRIPTINE MESYLATE  
PARNATE, TRANYLCYPROMINE SULFATE  
PAROEX, CHLORHEXIDINE GLUCONATE  
PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE  
PAROXETINE, PAROXETINE HYDROCHLORIDE  
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE  
PAROXETINE MESYLATE, PAROXETINE MESYLATE  
PARSABIV, ETELCALCETIDE  
PASER, AMINOSALICYLIC ACID  
PATADAY, OLOPATADINE HYDROCHLORIDE  
PATANASE, OLOPATADINE HYDROCHLORIDE  
PATANOL, OLOPATADINE HYDROCHLORIDE  
PAXIL, PAROXETINE HYDROCHLORIDE  
PAXIL CR, PAROXETINE HYDROCHLORIDE  
PAZEO, OLOPATADINE HYDROCHLORIDE  
PCE, ERYTHRHYMYCIN  
PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE  
PEDIATRIC ADVIL, IBUPROFEN (OTC)  
PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350  
PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL  
PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL  
PEG-3350, SODIUM CHLORIDE, SODIUM BICARBONATE, POTASSIUM CHLORIDE AND BISACODYL, BISACODYL  
PEGANONE, ETHOTOIN  
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM  
PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM  
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE  
PENICILLIN G SODIUM, PENICILLIN G SODIUM  
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM  
PENICILLIN-VK, PENICILLIN V POTASSIUM  
PENLAC, CICLOPIROX  
PENNsaid, DICLOFENAC SODIUM  
PENTAM, PENTAMIDINE ISETHIONATE  
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE  
PENTASA, MESALAMINE  
PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM  
PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM  
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM  
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE  
PENTOSTATIN, PENTOSTATIN  
PENTOXIFYLLINE, PENTOXIFYLLINE  
PENTOXIL, PENTOXIFYLLINE  
PEPCID, FAMOTIDINE  
PEPCID AC, FAMOTIDINE (OTC)  
PEPCID AC, FAMOTIDINE (OTC)  
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)  
PERCOSET, ACETAMINOPHEN  
PERCODAN, ASPIRIN  
PERFOROMIST, FORMOTEROL FUMARATE  
PERIDEX, CHLORHEXIDINE GLUCONATE  
PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS  
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE  
PERIOCHIP, CHLORHEXIDINE GLUCONATE  
PERIOGARD, CHLORHEXIDINE GLUCONATE  
PERMAPEN, PENICILLIN G BENZATHINE  
PERMETHRIN, PERMETHRIN (OTC)  
PERMETHRIN, PERMETHRIN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* P \*\*

PERPHENAZINE, PERPHENAZINE  
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE  
PERSANTINE, DIPYRIDAMOLE  
PERTZYE, PANCRELIPASE (AMYLASE)  
PEXEVA, PAROXETINE MESYLATE  
PFIZERPEN, PENICILLIN G POTASSIUM  
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE  
PHENELZINE SULFATE, PHENELZINE SULFATE  
PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE  
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE  
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE  
PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE  
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE  
PHENYTEK, PHENYTOIN SODIUM  
PHENYTOIN, PHENYTOIN  
PHENYTOIN SODIUM, PHENYTOIN SODIUM  
PHILITH, ETHINYL ESTRADIOL  
PHOSLO GELCAPS, CALCIUM ACETATE  
PHOSLYRA, CALCIUM ACETATE  
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE  
PHOTFRIN, PORFIMER SODIUM  
PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM  
PHOTREXA VISCOSUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM  
PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
PHYTONADIONE, PHYTONADIONE  
PICATO, INGENOL MEButATE  
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE  
PIMOZIDE, PIMOZIDE  
PIMTREA, DESOGESTREL  
PINDOLOL, PINDOLOL  
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE  
PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE  
PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
PIPERACILLIN, PIPERACILLIN SODIUM  
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM  
PIRMELLA 1/35, ETHINYL ESTRADIOL  
PIRMELLA 7/7/7, ETHINYL ESTRADIOL  
PIROXICAM, PIROXICAM  
PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM  
PITOCIN, OXYTOCIN  
PLAN B ONE-STEP, LEVONORGESTREL (OTC)  
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE  
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
PLAVIX, CLOPIDOGREL BISULFATE  
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PLIAGLIS, LIDOCAINE  
PODOFILOX, PODOFILOX  
POLOCAINE, MEPIVACAINE HYDROCHLORIDE  
POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350  
POLYMYCIN B SULFATE, POLYMYXIN B SULFATE  
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE  
POLYTRIM, POLYMYXIN B SULFATE  
POMALYST, POMALIDOMIDE  
PONSTEL, MEFENAMIC ACID  
PORTIA-28, ETHINYL ESTRADIOL  
POTASSIUM ACETATE, POTASSIUM ACETATE  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER,

## **APPENDIX A - PRODUCT NAME INDEX**

\*\*\* P \*\*\*

**APPENDIX A - PRODUCT NAME INDEX**

\*\* P \*\*

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CITRATE, POTASSIUM CITRATE  
POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)  
POVIDONE IODINE, POVIDONE-IODINE (OTC)  
PRADAXA, DABIGATRAN ETEXILATE MESYLATE  
PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE  
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE  
PRAMOSONE, HYDROCORTISONE ACETATE  
PRANDIN, REPAGLINIDE  
PRASUGREL, PRASUGREL HYDROCHLORIDE  
PRAVACHOL, PRAVASTATIN SODIUM  
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM  
PRAZIQUANTEL, PRAZIQUANTEL  
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE  
PRE-OP, HEXACHLOROPHENE  
PRE-OP II, HEXACHLOROPHENE  
PRE-PEN, BENZYL PENICILLOYL POLYLYSINE  
PRECEDEX, DEXMEDETomidine HYDROCHLORIDE  
PRECOSE, ACARBOSE  
PRED FORTE, PREDNISOLONE ACETATE  
PRED MILD, PREDNISOLONE ACETATE  
PRED-G, GENTAMICIN SULFATE  
PREDNICARBATE, PREDNICARBATE  
PREDNISOLONE, PREDNISOLONE  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE  
PREDNISONE, PREDNISONE  
PREDNISONE INTENSOL, PREDNISONE  
PREGNYL, GONADOTROPIN, CHORIONIC  
PRELONE, PREDNISOLONE  
PREMARIN, ESTROGENS, CONJUGATED  
PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS

**APPENDIX A - PRODUCT NAME INDEX**

\*\* P \*\*

PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS  
PREMPHASE 14/14, ESTROGENS, CONJUGATED  
PREMPRO, ESTROGENS, CONJUGATED  
PREPIDIL, DINOPROSTONE  
PREPOPIK, CITRIC ACID  
PRESTALIA, AMLODIPINE BESYLATE  
PREVACID, LANSOPRAZOLE  
PREVACID 24 HR, LANSOPRAZOLE (OTC)  
PREVALITE, CHOLESTYRAMINE  
PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)  
PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
PREVIFEM, ETHINYLY ESTRADIOL  
PREVPAC, AMOXICILLIN  
PREVYMIS, LETERMUVIR  
PREXXARTAN, VALSARTAN  
PREZCOBIX, COBICISTAT  
PREZISTA, DARUNAVIR ETHANOLATE  
PRIALT, ZICONOTIDE ACETATE  
PRIFTIN, RIFAPENTINE  
PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE  
PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE  
PRILOSEC, OMEPRAZOLE MAGNESIUM  
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)  
PRIMAQUINE, PRIMAQUINE PHOSPHATE  
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE  
PRIMAXIN, CILASTATIN SODIUM  
PRIMIDONE, PRIMIDONE  
PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE  
PRINIVIL, LISINOPRIL  
PRISMASOL B22GK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISTIQ, DESVENLAFAZINE SUCCINATE  
PROAIR HFA, ALBUTEROL SULFATE  
PROAIR RESPICLICK, ALBUTEROL SULFATE  
PROBALAN, PROBENECID  
PROBENECID, PROBENECID  
PROBENECID AND COLCHICINE, COLCHICINE  
PROBUPHINE, BUPRENORPHINE HYDROCHLORIDE  
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE  
PROCALAMINE, AMINO ACIDS  
PROCARDIA, NIFEDIPINE  
PROCARDIA XL, NIFEDIPINE  
PROCHLORPERAZINE, PROCHLORPERAZINE  
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE  
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE  
PROCOMP, PROCHLORPERAZINE MALEATE  
PROCTOFOAM HC, HYDROCORTISONE ACETATE  
PROCYSB, CYSTEAMINE BITARTRATE  
PROFEN, IBUPROFEN (OTC)  
PROFERDEX, IRON DEXTRAN  
PROGESTERONE, PROGESTERONE  
PROGLYCEM, DIAZOXIDE  
PROGRAF, TACROLIMUS  
PROHANCE, GADOTERIDOL  
PROHANCE MULTIPACK, GADOTERIDOL  
PROLENZA, BROMFENAC SODIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* P \*\*

PROMACTA, ELTROMBOPAG OLAMINE  
PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE PHOSPHATE  
PROMETH VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE  
PROMETH VC W/ CODEINE, CODEINE PHOSPHATE  
PROMETH W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE  
PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE  
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN  
PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE  
PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE  
PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE  
PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE  
PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE  
PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE  
PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE  
PROMETRIUM, PROGESTERONE  
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE  
PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE  
PROPARACAIN HYDROCHLORIDE, PROPARACAIN HYDROCHLORIDE  
PROPECIA, FINASTERIDE  
PROPOFOL, PROPOFOL  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE  
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
PROPYLTIOURACIL, PROPYLTIOURACIL  
PROSCAR, FINASTERIDE  
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS  
PROSTIN E2, DINOPROSTONE  
PROSTIN VR PEDIATRIC, ALPROSTADIL  
PROTAMINE SULFATE, PROTAMINE SULFATE  
PROTONIX, PANTOPRAZOLE SODIUM  
PROTONIX IV, PANTOPRAZOLE SODIUM  
PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE  
PROTOPIC, TACROLIMUS  
PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE  
PROVAYBLUE, METHYLENE BLUE  
PROVENTIL-HFA, ALBUTEROL SULFATE  
PROVERA, MEDROXYPROGESTERONE ACETATE  
PROVIGIL, MODAFINIL  
PROVOCHOLINE, METHACHOLINE CHLORIDE  
PROZAC, FLUOXETINE HYDROCHLORIDE  
PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE  
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)  
PULMICORT FLEXHALER, BUDESONIDE  
PULMICORT RESPULES, BUDESONIDE  
PULMOLITE, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT  
PUR-WASH, PURIFIED WATER (OTC)  
PURINETHOL, MERCAPTOPURINE  
PURIXAN, MERCAPTOPURINE  
PYLERA, BISMUTH SUBCITRATE POTASSIUM  
PYRAZINAMIDE, PYRAZINAMIDE  
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE  
PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE  
PYTEST, UREA, C-14  
PYTEST KIT, UREA, C-14

\*\* Q \*\*

QBRELIS, LISINOPRIL  
QNASL, BECLOMETHASONE DIPROPIONATE  
QOLIANA, BRIMONIDINE TARTRATE  
QSYMIA, PHENTERMINE HYDROCHLORIDE  
QTERN, DAPAGLIFLOZIN PROPANEDIOL  
QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM  
QUALAQWIN, QUININE SULFATE  
QUARTETTE, ETHINYLMESTRADIOL

**APPENDIX A - PRODUCT NAME INDEX**

\*\* Q \*\*

QUASENSE, ETHINYL ESTRADIOL  
QUDEXY XR, TOPIRAMATE  
QUELICIN, SUCCINYLCHOLINE CHLORIDE  
QUELICIN PRESERVATIVE FREE, SUCCINYLCHOLINE CHLORIDE  
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE  
QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE  
QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE  
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE  
QUINAPRIL HYDROCHLORIDE AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE  
QUINARETIC, HYDROCHLORTHIAZIDE  
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE  
QUINIDINE SULFATE, QUINIDINE SULFATE  
QUININE SULFATE, QUININE SULFATE  
QUTENZA, CAPSAICIN  
QVAR 40, BECLOMETHASONE DIPROPIONATE  
QVAR 80, BECLOMETHASONE DIPROPIONATE  
QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

\*\* R \*\*

R-GENE 10, ARGinine HYDROCHLORIDE  
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM  
RADICAVA, EDARAVONE  
RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)  
RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE  
RAMELTEON, RAMELTEON  
RAMIPRIL, RAMIPRIL  
RANEXA, RANOLAZINE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
RAPAFLO, SILODOSIN  
RAPAMUNE, SIROLIMUS  
RAPIVAB, PERAMIVIR  
RASAGILINE MESYLATE, RASAGILINE MESYLATE  
RASUVO, METHOTREXATE  
RAVICTI, GLYCEROL PHENYLBUTYRATE  
RAYALDEE, CALCIFEDIOL  
RAYOS, PREDNISONE  
RAZADYNE, GALANTAMINE HYDROBROMIDE  
RAZADYNE ER, GALANTAMINE HYDROBROMIDE  
READI-CAT 2, BARIUM SULFATE  
READI-CAT 2 SMOOTHIES, BARIUM SULFATE  
REBETOL, RIBAVIRIN  
RECLAST, ZOLEDRONIC ACID  
RECTIV, NITROGLYCERIN  
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE  
REGONOL, PYRIDOSTIGMINE BROMIDE  
RELENZA, ZANAMIVIR  
RELISTOR, METHYLNALTREXONE BROMIDE  
RELPAX, ELETRIPTAN HYDROBROMIDE  
REMERON, MIRTAZAPINE  
REMERON SOLTAB, MIRTAZAPINE  
REMODULIN, TREPROSTINIL  
RENACIDIN, CITRIC ACID  
RENAGEL, SEVELAMER HYDROCHLORIDE  
RENOGRAFIN-76, DIATRIZOATE MEGLUMINE  
RENOVA, TRETINOIN  
RENELVA, SEVELAMER CARBONATE  
REPAGLINIDE, REPAGLINIDE  
REPAGLINIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
REPREXAIN, HYDROCODONE BITARTRATE  
REQUIP, ROPINIROLE HYDROCHLORIDE  
REQUIP XL, ROPINIROLE HYDROCHLORIDE  
RESCRIPTOR, DELAVIRDINE MESYLATE  
RESECTISOL IN PLASTIC CONTAINER, MANNITOL  
RESTASIS, CYCLOSPORINE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* R \*\*

RESTASIS MULTIDOSE, CYCLOSPORINE  
RESTORIL, TEMAZEPAM  
RETIN-A, TRETINOIN  
RETIN-A MICRO, TRETINOIN  
RETIN-A-MICRO, TRETINOIN  
RETISERT, FLUOCINOLONE ACETONIDE  
RETROVIR, ZIDOVUDINE  
REVATIO, SILDENAFIL CITRATE  
REVLIMID, LENALIDOMIDE  
REVONTO, DANTROLENE SODIUM  
REXULTI, BREXPIPRAZOLE  
REYATAZ, ATAZANAVIR SULFATE  
REZIRA, HYDROCODONE BITARTRATE  
RHINOCORT ALLERGY, BUDESONIDE (OTC)  
RHOFADE, OXYMETAZOLINE HYDROCHLORIDE  
RHOPRESSA, NETARSUDIL DIMESYLATE  
RIBASPHERE, RIBAVIRIN  
RIBAVARIN, RIBAVIRIN  
RIBAVIRIN, RIBAVIRIN  
RIDAURA, AURANOFIN  
RIFABUTIN, RIFABUTIN  
RIFADIN, RIFAMPIN  
RIFAMATE, ISONIAZID  
RIFAMPIN, RIFAMPIN  
RIFATER, ISONIAZID  
RILUTEK, RILUZOLE  
RILUZOLE, RILUZOLE  
RIMACTANE, RIFAMPIN  
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE  
RIMSO-50, DIMETHYL SULFOXIDE  
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
RIOMET, METFORMIN HYDROCHLORIDE  
RISEDRONATE SODIUM, RISETDRONATE SODIUM  
RISPERDAL, RISPERIDONE  
RISPERDAL CONSTA, RISPERIDONE  
RISPERIDONE, RISPERIDONE  
RITALIN, METHYLPHENIDATE HYDROCHLORIDE  
RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE  
RITALIN-SR, METHYLPHENIDATE HYDROCHLORIDE  
RITONAVIR, RITONAVIR  
RIVASTIGMINE, RIVASTIGMINE  
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE  
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
ROBAXIN, METHOCARBAMOL  
ROBAXIN-750, METHOCARBAMOL  
ROBINUL, GLYCOPYRROLATE  
ROBINUL FORTE, GLYCOPYRROLATE  
ROCALTROL, CALCITRIOL  
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE  
ROGAINE (FOR MEN), MINOXIDIL (OTC)  
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)  
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)  
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE  
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE  
ROSIGLITAZONE MALEATE, ROSIGLITAZONE MALEATE  
ROSIGLITAZONE MALEATE AND GLIMEPIRIDE, GLIMEPIRIDE  
ROSIGLITAZONE MALEATE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM  
ROWASA, MESALAMINE  
ROWEEPRA, LEVETIRACETAM  
ROXICET, ACETAMINOPHEN  
ROXICODONE, OXYCODONE HYDROCHLORIDE  
ROXYBOND, OXYCODONE HYDROCHLORIDE  
ROZEREM, RAMELTEON  
RUBRACA, RUCAPARIB CAMSYLATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* R \*\*

RUBY-FILL, RUBIDIUM CHLORIDE RB-82  
RUFINAMIDE, RUFINAMIDE  
RYANODEX, DANTROLENE SODIUM  
RYDAPT, MIDOSTAURIN  
RYTARY, CARBIDOPA  
RYTHMOL, PROPAFENONE HYDROCHLORIDE  
RYTHMOL SR, PROPAFENONE HYDROCHLORIDE  
RYZODEG 70/30, INSULIN ASPART

\*\* S \*\*

SABRIL, VIGABATRIN  
SAFYRAL, DROSPIRENONE  
SAIZEN, SOMATROPIN RECOMBINANT  
SALAGEN, PILOCARPINE HYDROCHLORIDE  
SALONPAS, MENTHOL (OTC)  
SAMSCA, TOLVAPTAN  
SANCUSO, GRANisetron  
SANDIMMUNE, CYCLOSPORINE  
SANDOSTATIN, OCTREOTIDE ACETATE  
SANDOSTATIN LAR, OCTREOTIDE ACETATE  
SAPHRIS, ASENAPINE MALEATE  
SARAFEM, FLUOXETINE HYDROCHLORIDE  
SAVAYSA, EDOXABAN TOSYLATE  
SAVELLA, MILNACIPRAN HYDROCHLORIDE  
SAXENDA, LIRAGLUTIDE RECOMBINANT  
SCANDONEST L, LEVONORDEFRIN  
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE  
SCANLUX-300, IOPAMIDOL  
SCANLUX-370, IOPAMIDOL  
SCLEROSOL, TALC  
SCOPOLAMINE, SCOPOLAMINE  
SEASONALE, ETHINYL ESTRADIOL  
SEASONIQUE, ETHINYL ESTRADIOL  
SECONAL SODIUM, SECOBARBITAL SODIUM  
SECTRAL, ACEBUTOLOL HYDROCHLORIDE  
SEEGBRI, GLYCOPYRROLATE  
SEGLUROMET, ERTUGLIFLOZIN  
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE  
SELENIUM SULFIDE, SELENIUM SULFIDE  
SELFEMRA, FLUOXETINE HYDROCHLORIDE  
SELSUN, SELENIUM SULFIDE  
SELZENTRY, MARAVIROC  
SEMPREX-D, ACRIVASTINE  
SENSIPAR, CINACALCET HYDROCHLORIDE  
SENSORCAINE, BUPIVACAINE HYDROCHLORIDE  
SEPTOCaine, ARTICaine HYDROCHLORIDE  
SEPTRA, SULFAMETHOXAZOLE  
SEPTRA DS, SULFAMETHOXAZOLE  
SEREVENT, SALMETEROL XINAFOATE  
SERNIVO, BETAMETHASONE DIPROPIONATE  
SEROMYCIN, CYCLOSERINE  
SEROQUEL, QUETIAPINE FUMARATE  
SEROQUEL XR, QUETIAPINE FUMARATE  
SEROSTIM, SOMATROPIN RECOMBINANT  
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE  
SETLAKIN, ETHINYL ESTRADIOL  
SEVELAMER CARBONATE, SEVELAMER CARBONATE  
SEVOFLURANE, SEVOFLURANE  
SFROWASA, MESALAMINE  
SIGNIFOR, PASIREOTIDE DIASPARTATE  
SIGNIFOR LAR, PASIREOTIDE PAMOATE  
SIKLOS, HYDROXYUREA  
SILDENAFIL CITRATE, SILDENAFIL CITRATE  
SILENOR, DOXEPEP HYDROCHLORIDE  
SILODOSIN, SILODOSIN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* S \*\*

SILVADENE, SILVER SULFADIAZINE  
SIMBRINZA, BRIMONIDINE TARTRATE  
SIMPESSE, ETHINYLI ESTRADIOL  
SIMVASTATIN, SIMVASTATIN  
SINE-AID IB, IBUPROFEN (OTC)  
SINEMET, CARBIDOPA  
SINEMET CR, CARBIDOPA  
SINGULAIR, MONTELUKAST SODIUM  
SINOGRAFIN, DIATRIZOATE MEGLUMINE  
SINUVA, MOMETASONE FUROATE  
SIROLIMUS, SIROLIMUS  
SIRTURO, BEDAQUILINE FUMARATE  
SITAVIG, ACYCLOVIR  
SIVEXTRO, TEDIZOLID PHOSPHATE  
SKELAXIN, METAXALONE  
SKLICE, IVERMECTIN  
SKYLA, LEVONORGESTREL  
SMOFLIPID 20%, FISH OIL  
SODIUM ACETATE, SODIUM ACETATE  
SODIUM BICARBONATE, SODIUM BICARBONATE  
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18  
SODIUM IODIDE I 123, SODIUM IODIDE I-123  
SODIUM IODIDE I 131, SODIUM IODIDE I-131  
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE  
SODIUM NITRITE, SODIUM NITRITE  
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE  
SODIUM OXYBATE, SODIUM OXYBATE  
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE  
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE  
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE  
SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE  
SODIUM THIOSULFATE, SODIUM THIOSULFATE  
SOJOURN, SEVOFLURANE  
SOLARAZE, DICLOFENAC SODIUM  
SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE  
SOLIQUA 100/33, INSULIN GLARGINE  
SOLODYN, MINOCYCLINE HYDROCHLORIDE  
SOLOSEC, SECNIDAZOLE  
SOLTAMOX, TAMOXIFEN CITRATE  
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE  
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE  
SOMA, CARISOPRODOL  
SOMATULINE DEPOT, LANREOTIDE ACETATE  
SOMAVERT, PEGVISOMANT  
SONATA, ZALEPLON  
SOOLANTRA, IVERMECTIN  
SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL  
SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL  
SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL  
SORIATANE, ACITRETIN  
SORILUX, CALCIPOTRIENE  
SORINE, SOTALOL HYDROCHLORIDE  
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE  
SOTRADECOL, SODIUM TETRADECYL SULFATE  
SOTYLIZE, SOTALOL HYDROCHLORIDE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* S \*\*

SOVALDI, SOFOSBUVIR  
SPECTAZOLE, ECONAZOLE NITRATE  
SPINRAZA, NUSINERSEN SODIUM  
SPIRIVA, TIOTROPIUM BROMIDE  
SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE  
SPIRONOLACTONE, SPIRONOLACTONE  
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
SPORANOX, ITRACONAZOLE  
SPRINTEC, ETHINYLMESTRADIOL  
SPRITAM, LEVETIRACETAM  
SPRIX, KETOROLAC TROMETHAMINE  
SPRYCEL, DASATINIB  
SPS, SODIUM POLYSTYRENE SULFONATE  
SSD, SILVER SULFADIAZINE  
SSD AF, SILVER SULFADIAZINE  
STALEVO 100, CARBIDOPA  
STALEVO 125, CARBIDOPA  
STALEVO 150, CARBIDOPA  
STALEVO 200, CARBIDOPA  
STALEVO 50, CARBIDOPA  
STALEVO 75, CARBIDOPA  
STARLIX, NATEGLINIDE  
STAVUDINE, STAVUDINE  
STAXYN, VARDENAFIL HYDROCHLORIDE  
STEGLATRO, ERTUGLIFLOZIN  
STELUJAN, ERTUGLIFLOZIN  
STENDRA, AVANAFIL  
STERILE WATER, STERILE WATER FOR IRRIGATION  
STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION  
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION  
STERITALC, TALC  
STIE-CORT, HYDROCORTISONE  
STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE  
STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE  
STIVARGA, REGORAFENIB  
STRATTERA, ATOMOXETINE HYDROCHLORIDE  
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE  
STRIANT, TESTOSTERONE  
STRIBILD, COBICISTAT  
STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE  
STROMECTOL, IVERMECTIN  
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89  
SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE  
SUBLOCADE, BUPRENORPHINE  
SUBOXONE, BUPRENORPHINE HYDROCHLORIDE  
SUBSYS, FENTANYL  
SUCRAID, SACROSIDASE  
SUCRALFATE, SUCRALFATE  
SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)  
SUDAFED 24 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)  
SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE  
SUFENTANIL CITRATE, SUFENTANIL CITRATE  
SULAR, NISOLDIPINE  
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM  
SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE  
SULFADIAZINE, SULFADIAZINE  
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE  
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE  
SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE  
SULFAMYLYN, MAFENIDE ACETATE  
SULFASALAZINE, SULFASALAZINE  
SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE  
SULINDAC, SULINDAC  
SUMATRIPTAN, SUMATRIPTAN

**APPENDIX A - PRODUCT NAME INDEX**

\*\* S \*\*

SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE  
SUMAVEL DOSEPRO, SUMATRIPTAN SUCCINATE  
SUPPRELIN LA, HISTRELIN ACETATE  
SUPRANE, DESFLURANE  
SUPRAX, CEFIXIME  
SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE  
SURMONTIL, TRIMIPRAMINE MALEATE  
SURVANTA, BERACTANT  
SUSTIVA, EFAVIRENZ  
SUSTOL, GRANISETRON  
SUTENT, SUNITINIB MALATE  
SYEDA, DROSPIRENONE  
SYMBICORT, BUDESONIDE  
SYMBYAX, FLUOXETINE HYDROCHLORIDE  
SYMJEPI, EPINEPHRINE  
SYMLIN, PRAMLINE ACETATE  
SYMPROIC, NALDEMEDINE TOSYLATE  
SYNALAR, FLUOCINOLONE ACETONIDE  
SYNALGOS-DC, ASPIRIN  
SYNAREL, NAFARELIN ACETATE  
SYNDROS, DRONABINOL  
SYNERA, LIDOCAINE  
SYNERCID, DALFOPRISTIN  
SYNJARDY, EMPAGLIFLOZIN  
SYNJARDY XR, EMPAGLIFLOZIN  
SYNRIBO, OMACETAXINE MEPESUCCINATE  
SYNTHROID, LEVOTHYROXINE SODIUM \*\*  
SYPRINE, TRIENTINE HYDROCHLORIDE

\*\* T \*\*

TAB-PROFEN, IBUPROFEN (OTC)  
TACLONEX, BETAMETHASONE DIPROPIONATE  
TACROLIMUS, TACROLIMUS  
TAFINLAR, DABRAFENIB MESYLATE  
TAGAMET HB, CIMETIDINE (OTC)  
TAGITOL V, BARIUM SULFATE  
TAGRISSO, OSIMERTINIB MESYLATE  
TALC, TALC  
TALWIN, PENTAZOCINE LACTATE  
TAMBOCOR, FLECAINIDE ACETATE  
TAMIFLU, OSELTAMIVIR PHOSPHATE  
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE  
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE  
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE  
TAPAZOLE, METHIMAZOLE  
TARCEVA, ERLOTINIB HYDROCHLORIDE  
TARGRETIN, BEXAROTENE  
TARKA, TRANDOLAPRIL  
TASIGNA, NILOTINIB HYDROCHLORIDE MONOHYDRATE  
TASMAR, TOLCAPONE  
TAVIST-1, CLEMASTINE FUMARATE (OTC)  
TAXOL, PACLITAXEL  
TAXOTERE, DOCETAXEL  
TAYTULLA, ETHINYLMESTRADIOL  
TAZAROTENE, TAZAROTENE  
TAZICEF, CEFTAZIDIME  
TAZORAC, TAZAROTENE  
TAZTIA XT, DILTIAZEM HYDROCHLORIDE  
TECFIDERA, DIMETHYL FUMARATE  
TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR  
TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT  
TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT  
TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT  
TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT  
TECHNETIUM TC 99M GENERATOR, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

**APPENDIX A - PRODUCT NAME INDEX**

\*\* T \*\*

TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT  
TECHNETIUM TC-99M MEBROFENIN, TECHNETIUM TC-99M MEBROFENIN KIT  
TECHNETIUM TC-99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT  
TECHNIVIE, OMBITASVIR  
TEFLARO, CEFTAROLINE FOSAMIL  
TEGRETOL, CARBAMAZEPINE  
TEGRETOL-XR, CARBAMAZEPINE  
TEKTURNA, ALISKIREN HEMIFUMARATE  
TEKTURNA HCT, ALISKIREN HEMIFUMARATE  
TELMISARTAN, TELMISARTAN  
TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE  
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
TEMAZEPAM, TEMAZEPAM  
TEMODAR, TEMOZOLOMIDE  
TEMOZOLOMIDE, TEMOZOLOMIDE  
TENEX, GUANFACINE HYDROCHLORIDE  
TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE  
TENORETIC 100, ATENOLOL  
TENORETIC 50, ATENOLOL  
TENORMIN, ATENOLOL  
TENUATE, DIETHYLPROPION HYDROCHLORIDE  
TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE  
TEPADINA, THIOTEPА  
TERAZOL 3, TERCONAZOLE  
TERAZOL 7, TERCONAZOLE  
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE  
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)  
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE  
TERBUTALINE SULFATE, TERBUTALINE SULFATE  
TERCONAZOLE, TERCONAZOLE  
TERIL, CARBAMAZEPINE  
TERRAMYCIN W/ POLYMYXIN B SULFATE, OXYTETRACYCLINE HYDROCHLORIDE  
TESSALON, BENZONATATE  
TESTIM, TESTOSTERONE  
TESTOPEL, TESTOSTERONE  
TESTOSTERONE, TESTOSTERONE  
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE  
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE  
TESTRED, METHYLTESTOSTERONE  
TETRABENAZINE, TETRABENAZINE  
TETRACAIN HYDROCHLORIDE, TETRACAIN HYDROCHLORIDE  
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE  
TEVETEN, EPROSARTAN MESYLATE  
TEXACORT, HYDROCORTISON  
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201  
THALOMID, THALIDOMIDE  
THAM, TROMETHAMINE  
THEO-24, THEOPHYLLINE  
THEOCHRON, THEOPHYLLINE  
THEOPHYLLINE, THEOPHYLLINE  
THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE  
THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE  
THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE  
THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE  
THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE  
THERMAZENE, SILVER SULFADIAZINE  
THEROXIDIL, MINOXIDIL (OTC)  
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE  
THIOGUANINE, THIOGUANINE  
THIOLA, TIOPRONIN  
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE  
THIOTEPА, THIOTEPА  
THIOTHIXENE, THIOTHIXENE  
THRIVE, NICOTINE POLACRILEX (OTC)  
THYROGEN, THYROTROPIN ALFA

**APPENDIX A - PRODUCT NAME INDEX**

\*\* T \*\*

THYROLAR-0.25, LIOTRIX (T4)  
THYROLAR-0.5, LIOTRIX (T4)  
THYROLAR-1, LIOTRIX (T4)  
THYROLAR-2, LIOTRIX (T4)  
THYROLAR-3, LIOTRIX (T4)  
THYROSAFE, POTASSIUM IODIDE (OTC)  
THYROSHIELD, POTASSIUM IODIDE (OTC)  
TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE  
TIAZAC, DILTIAZEM HYDROCHLORIDE  
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE  
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE  
TIGECYCLINE, TIGECYCLINE  
TIKOSYN, DOFETILIDE  
TIMOLOL, TIMOLOL  
TIMOLOL MALEATE, TIMOLOL MALEATE  
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE  
TIMOPTIC-XE, TIMOLOL MALEATE  
TINDAMAX, TINIDAZOLE  
TINIDAZOLE, TINIDAZOLE  
TIOCONAZOLE, TIOCONAZOLE (OTC)  
TIROSINT, LEVOTHYROXINE SODIUM  
TIS-U-SOL, MAGNESIUM SULFATE  
TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE  
TIVICAY, DOLUTEGRAVIR SODIUM  
TIVORBEX, INDOMETHACIN  
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE  
TOBI, TOBRAMYCIN  
TOBI PODHALER, TOBRAMYCIN  
TOBRADEX, DEXAMETHASONE  
TOBRADEX ST, DEXAMETHASONE  
TOBRAMYCIN, TOBRAMYCIN  
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE  
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE  
TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE  
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE  
TOBREX, TOBRAMYCIN  
TODAY, NONOXYNOL-9 (OTC)  
TOFRANIL, IMIPRAMINE HYDROCHLORIDE  
TOLAK, FLUOROURACIL  
TOLAZAMIDE, TOLAZAMIDE  
TOLBUTAMIDE, TOLBUTAMIDE  
TOLCAPONE, TOLCAPONE  
TOLMETIN SODIUM, TOLMETIN SODIUM  
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE  
TOPAMAX, TOPIRAMATE  
TOPICORT, DESOXIMETASONE  
TOPIRAMATE, TOPIRAMATE  
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE  
TOPROL-XL, METOPROLOL SUCCINATE  
TORISEL, TEMSIROLIMUS  
TORSEMIDE, TORSEMIDE  
TOTECT, DEXRAZOXANE HYDROCHLORIDE  
TOUJEO SOLOSTAR, INSULIN GLARGINE RECOMBINANT  
TOVIAZ, FESOTERODINE FUMARATE  
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
TRACLEER, BOSENTAN  
TRADJENTA, LINAGLIPTIN  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN  
TRANDATE, LABETALOL HYDROCHLORIDE  
TRANDOLAPRIL, TRANDOLAPRIL  
TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL  
TRANEXAMIC ACID, TRANEXAMIC ACID  
TRANSDERM SCOP, SCOPOLAMINE  
TRANXENE, CLORAZEPATE DIPOTASSIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* T \*\*

TRANLYCYPROMINE SULFATE, TRANLYCYPROMINE SULFATE  
TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS  
TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS  
TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS  
TRAVATAN Z, TRAVOPROST  
TRAVOPROST, TRAVOPROST  
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE  
TREANDA, BENDAMUSTINE HYDROCHLORIDE  
TRECATOR, ETHIONAMIDE  
TRELEGY ELLIPTA, FLUTICASONE FUROATE  
TRELSTAR, TRIPTORELIN PAMOATE  
TREPROSTINIL, TREPROSTINIL  
TRESIBA, INSULIN DEGLUDEC  
TRETINOIN, TRETINOIN  
TREXALL, METHOTREXATE SODIUM  
TREXIMET, NAPROXEN SODIUM  
TREZIX, ACETAMINOPHEN  
TRI LO SPRINTEC, ETHINYL ESTRADIOL  
TRI-ESTARYLLA, ETHINYL ESTRADIOL  
TRI-LEGEST 21, ETHINYL ESTRADIOL  
TRI-LEGEST FE, ETHINYL ESTRADIOL  
TRI-LINYAH, ETHINYL ESTRADIOL  
TRI-LO-ESTARYLLA, ETHINYL ESTRADIOL  
TRI-LO-MILI, ETHINYL ESTRADIOL  
TRI-LUMA, FLUOCINOLONE ACETONIDE  
TRI-MILI, ETHINYL ESTRADIOL  
TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL  
TRI-PREVIFEM, ETHINYL ESTRADIOL  
TRI-SPRINTEC, ETHINYL ESTRADIOL  
TRIACIN-C, CODEINE PHOSPHATE  
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)  
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE  
TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE  
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
TRIAZOLAM, TRIAZOLAM  
TRIBENZOR, AMLODIPINE BESYLATE  
TRICOR, FENOFIBRATE  
TRIDERM, TRIAMCINOLONE ACETONIDE  
TRIDIOME, TRIMETHADIONE  
TRIESENCE, TRIAMCINOLONE ACETONIDE  
TRIFERIC, FERRIC PYROPHOSPHATE CITRATE  
TRIFERIC, FERRIC PYROPHOSPHATE CITRATE  
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE  
TRIFLURIDINE, TRIFLURIDINE  
TRIGLIDE, FENOFIBRATE  
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE  
TRILEPTAL, OXCARBAZEPINE  
TRILIPIX, CHOLINE FENOFIBRATE  
TRILYTE, POLYETHYLENE GLYCOL 3350  
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE  
TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE, TRIMETHOBENZAMIDE HYDROCHLORIDE  
TRIMETHOPRIM, TRIMETHOPRIM  
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE  
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE  
TRINTELLIX, VORTioxetine HYDROBROMIDE  
TRIOSTAT, LIOTHYRONINE SODIUM  
TRIPTODUR KIT, TRIPTORELIN PAMOATE  
TRISENOX, ARSENIC TRIOXIDE  
TRIUMEQ, ABACAVIR SULFATE  
TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)  
TRIVORA-28, ETHINYL ESTRADIOL  
TRIZIVIR, ABACAVIR SULFATE  
TROKENDI XR, TOPIRAMATE  
TROPHAMINE, AMINO ACIDS  
TROPHAMINE 10%, AMINO ACIDS

**APPENDIX A - PRODUCT NAME INDEX**

\*\* T \*\*

TROPICACYL, TROPICAMIDE  
TROPICAMIDE, TROPICAMIDE  
TROSPiUM CHLORiDE, TROSPiUM CHLORiDE  
TRULANCE, PLECANATiDE  
TRUSOPT, DORZOLAMiDE HYDROCHLORiDE  
TRUVADA, EMTRiCITABiNE  
TUDORZA PRESSAIR, ACLiDiNiUM BROMiDE  
TUSSiCAPS, CHLORPHENiRAMiNE POLiSTiREX  
TUSSiGON, HOMATROPiNE METHYLBRoMiDE  
TUSSIONEX PENNKiNETiC, CHLORPHENiRAMiNE POLiSTiREX  
TUZiSTRA XR, CHLORPHENiRAMiNE POLiSTiREX  
TWYNSTA, AMLODIPiNE BESiLYATE  
TYBOST, COBiCISTAT  
TYDEMAY, DROSpIRENONE  
TYGACiL, TiGECCYCLiNE  
TYKERB, LAPATiNiB DiTOSiLYATE  
TYLENOL, ACETAMiNOPhEN (OTC)  
TYLENOL W/ CODEiNE NO. 3, ACETAMiNOPhEN  
TYLENOL W/ CODEiNE NO. 4, ACETAMiNOPhEN  
TYMLOS, ABALOPARATiDE  
TYVASO, TREPROSTiNiL  
TYZiNE, TETRAHYDROZOLiNE HYDROCHLORiDE

\*\* U \*\*

U-CORT, HYDROCORTiSONE ACETATE  
UCERiS, BUDESONiDE  
ULESFIA, BENZYL ALCOHOL  
ULORiC, FEBUXOSTAT  
ULTACAN, ARTiCAiNE HYDROCHLORiDE  
ULTACAN FORTE, ARTiCAiNE HYDROCHLORiDE  
ULTANE, SEVOFLURANE  
ULTiVA, REMiFENTANiL HYDROCHLORiDE  
ULTRA-TECHNEKOW FM, TECHNETiUM TC-99M SODiUM PERTECHNETATE GENERATOR  
ULTRACET, ACETAMiNOPhEN  
ULTRAM, TRAMADOL HYDROCHLORiDE  
ULTRATAG, TECHNETiUM TC-99M RED BLOOD CELL KiT  
ULTRAVATE, HALOBETASOL PROPIONATE  
ULTRAViST (PHARMACY BULK), iOPROMiDE  
ULTRAViST 240, iOPROMiDE  
ULTRAViST 300, iOPROMiDE  
ULTRAViST 370, iOPROMiDE  
ULTRESA, PANCRELiPASE (AMYLASE)  
UNASyN, AMPiCiLLiN SODiUM  
UNiSiOM, DOXYLAMiNE SUCCiNATE (OTC)  
UNiTHROiD, LEVOTHYRoXiNE SODiUM \*\*  
UPTRAVi, SELEXiPAG  
URECHOLiNE, BETHANECHOL CHLORiDE  
UREX, METHENAMiNE HiPPURATE  
UROCiT-K, POTASSiUM CiTRATE  
UROXATRAL, ALFUZOSiN HYDROCHLORiDE  
URSO 250, URSoDiOL  
URSO FORTE, URSoDiOL  
URSoDiOL, URSoDiOL  
UTiBRON, GLYCOPYRROLATE  
UVADEX, METHOXALEN

\*\* V \*\*

VABOMERE, MEROPENEM  
VAGiFEM, ESTRADiOL  
VAGiSTAT-1, TIOCONAZOLE (OTC)  
VALACYCLOCViR HYDROCHLORiDE, VALACYCLOCViR HYDROCHLORiDE  
VALCHLOR, MECHLORETHAMiNE HYDROCHLORiDE  
VALCYTE, VALGANCiCLOCViR HYDROCHLORiDE  
VALGANCiCLOCViR HYDROCHLORiDE, VALGANCiCLOCViR HYDROCHLORiDE  
VALiUM, DIAZEPAM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* V \*\*

VALNAC, BETAMETHASONE VALERATE  
VALPROATE SODIUM, VALPROATE SODIUM  
VALPROIC ACID, VALPROIC ACID  
VALSARTAN, VALSARTAN  
VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
VALSTAR PRESERVATIVE FREE, VALRUBICIN  
VALTREX, VALACYCLOVIR HYDROCHLORIDE  
VALTROPIN, SOMATROPIN RECOMBINANT  
VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE  
VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE  
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE  
VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE  
VANDAZOLE, METRONIDAZOLE  
VANIQA, EFLORNITHINE HYDROCHLORIDE  
VANOS, FLUOCINONIDE  
VANTAS, HISTRELIN ACETATE  
VANTRELA ER, HYDROCODONE BITARTRATE  
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE  
VARIBAR, BARIUM SULFATE  
VARIBAR NECTAR, BARIUM SULFATE  
VARITHENA, POLIDOCANOL  
VARUBI, ROLAPITANT HYDROCHLORIDE  
VASCEPA, ICOSAPENT ETHYL  
VASERETIC, ENALAPRIL MALEATE  
VASO STRICT, VASOPRESSIN  
VASOTEC, ENALAPRIL MALEATE  
VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE  
VECTICAL, CALCITRIOL  
VECURONIUM BROMIDE, VECURONIUM BROMIDE  
VELCADE, BORTEZOMIB  
VELETRI, EPOPROSTENOL SODIUM  
VELIVET, DESOGESTREL  
VELPHORO, SUCROFERRIC OXYHYDROXIDE  
VELTASSA, PATIROMER SORBITEX CALCIUM  
VELTIN, CLINDAMYCIN PHOSPHATE  
VEMOLIDY, TENOFOVIR ALAFENAMIDE FUMARATE  
VENCLEXTA, VENETOCLAX  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE  
VENOFER, IRON SUCROSE  
VENTAVIS, ILOPROST  
VENTOLIN HFA, ALBUTEROL SULFATE  
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE  
VERDESO, DESONIDE  
VEREGEN, SINECATECHINS  
VERELAN, VERAPAMIL HYDROCHLORIDE  
VERELAN PM, VERAPAMIL HYDROCHLORIDE  
VERMOX, MEBENDAZOLE  
VERSACLOZ, CLOZAPINE  
VERZENIO, ABEMACICLIB  
VESICARE, SOLIFENACIN SUCCINATE  
VFEND, VORICONAZOLE  
VIAGRA, SILDENAFIL CITRATE  
VIBATIV, TELAVANCIN HYDROCHLORIDE  
VIBERZI, ELUXADOLINE  
VIBISONE, CYANOCOBALAMIN  
VIBRAMYCIN, DOXYCYCLINE  
VIBRAMYCIN, DOXYCYCLINE CALCIUM  
VIBRAMYCIN, DOXYCYCLINE HYCLATE  
VICTOZA, LIRAGLUTIDE RECOMBINANT  
VIDAZA, AZACITIDINE  
VIDEX, DIDANOSINE  
VIDEX EC, DIDANOSINE  
VIEKIRA PAK (COPACKAGED), DASABUVIR SODIUM  
VIEKIRA XR, DASABUVIR SODIUM  
VIENVA, ETHINYLMESTRADIOL

**APPENDIX A - PRODUCT NAME INDEX**

\*\* V \*\*

VIGABATRIN, VIGABATRIN  
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE  
 VIIBRYD, VILAZODONE HYDROCHLORIDE  
 VIMOVO, ESOMEPRAZOLE MAGNESIUM  
 VIMPAT, LACOSAMIDE  
 VINBLASTINE SULFATE, VINBLASTINE SULFATE  
 VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE  
 VINORELBINE TARTRATE, VINORELBINE TARTRATE  
 VIOKACE, PANCRELIPASE (AMYLASE)  
 VIORELE, DESOGESTREL  
 VIRACEPT, NELFINAVIR MESYLATE  
 VIRAMUNE, NEVIRAPINE  
 VIRAMUNE XR, NEVIRAPINE  
 VIRazole, RIBAVIRIN  
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE  
 VIROPTIC, TRIFLURIDINE  
 VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)  
 VISINE-A, NAPHAZOLINE HYDROCHLORIDE (OTC)  
 VISIONBLUE, TRYPLAN BLUE  
 VISIPAQUE 270, IODIXANOL  
 VISIPAQUE 320, IODIXANOL  
 VISTARIL, HYDROXYZINE PAMOATE  
 VISTOGARD, URIDINE TRIACETATE  
 VISUDYNE, VERTEPORFIN  
 VITAMIN D, ERGOCALCIFEROL  
 VITAMIN K1, PHYTONADIONE  
 VITRASE, HYALURONIDASE  
 VITUZ, CHLORPHENIRAMINE MALEATE  
 VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE  
 VIVELLE-DOT, ESTRADIOL  
 VIVITROL, NALTREXONE  
 VIVLODEX, MELOXICAM  
 VIZAMYL, FLUTEMETAMOL F-18  
 VOGELXO, TESTOSTERONE  
 VOLNEA, DESOGESTREL  
 VOLTAREN, DICLOFENAC SODIUM  
 VORICONAZOLE, VORICONAZOLE  
 VOSEVI, SOFOSBUVIR  
 VOSOL, ACETIC ACID, GLACIAL  
 VOSOL HC, ACETIC ACID, GLACIAL  
 VOSPIRE ER, ALBUTEROL SULFATE  
 VOTRIENT, PAZOPANIB HYDROCHLORIDE  
 VPRIV, VELAGLUCERASE ALFA  
 VRAYLAR, CARIPRAZINE HYDROCHLORIDE  
 VUSION, MICONAZOLE NITRATE  
 VYFEMLA, ETHINYLMESTRADIOL  
 VYTORIN, EZETIMIBE  
 VYVANSE, LISDEXAMFETAMINE DIMESYLATE  
 VYXEOS, CYTARABINE  
 VYZULTA, LATANOPROSTENE BUNOD

\*\* W \*\*

WARFARIN SODIUM, WARFARIN SODIUM  
 WELCHOL, COLESEVELAM HYDROCHLORIDE  
 WELLBUTRIN SR, BUPROPION HYDROCHLORIDE  
 WELLBUTRIN XL, BUPROPION HYDROCHLORIDE  
 WERA, ETHINYLMESTRADIOL  
 WOMEN'S ROGAINE, MINOXIDIL (OTC)

\*\* X \*\*

XADAGO, SAFINAMIDE MESYLATE  
 XALATAN, LATANOPROST  
 XALKORI, CRIZOTINIB  
 XANAX, ALPRAZOLAM  
 XANAX XR, ALPRAZOLAM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* X \*\*

XARELTO, RIVAROXABAN  
XATMEP, METHOTREXATE SODIUM  
XELJANZ, TOFACITINIB CITRATE  
XELJANZ XR, TOFACITINIB CITRATE  
XELODA, CAPECITABINE  
XENAZINE, TETRABENAZINE  
XENICAL, ORLISTAT  
XENON XE 133, XENON XE-133  
XEPI, OZENOXACIN  
XERESE, ACYCLOVIR  
XERMELO, TELOTRISTAT ETIPRATE  
XHANCE, FLUTICASONE PROPIONATE  
XIFAXAN, RIFAXIMIN  
XIGDUO XR, DAPAGLIFLOZIN PROPANEDIOL  
XIIDRA, LIFITEGRAST  
XIMINO, MINOCYCLINE HYDROCHLORIDE  
XOFIGO, RADIUM RA-223 DICHLORIDE  
XOLEGEL, KETOCONAZOLE  
XOPENEX, LEVALBUTEROL HYDROCHLORIDE  
XOPENEX HFA, LEVALBUTEROL TARTRATE  
XTAMPZA ER, OXYCODONE  
XTANDI, ENZALUTAMIDE  
XTORO, FINAFLOXACIN  
XULANE, ETHINYL ESTRADIOL  
XULTOPHY 100/3.6, INSULIN DEGLUDEC  
XURIDEN, URIDINE TRIACETATE  
XYLOCAINE, LIDOCAINE HYDROCHLORIDE  
XYLOCAINE 4% PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE  
XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE  
XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE  
XYREM, SODIUM OXYBATE  
XYZAL, LEVO CETIRIZINE DIHYDROCHLORIDE  
XYZAL ALLERGY 24HR, LEVO CETIRIZINE DIHYDROCHLORIDE (OTC)

\*\* Y \*\*

YABELA, DROSPIRENONE  
YASMIN, DROSPIRENONE  
YAZ, DROSPIRENONE  
YONDELIS, TRABECTEDIN  
YOSPRALA, ASPIRIN

\*\* Z \*\*

ZAFIRLUKAST, ZAFIRLUKAST  
ZALEPLON, ZALEPLON  
ZANAFLEX, TIZANIDINE HYDROCHLORIDE  
ZANOSAR, STREPTOZOZOCIN  
ZANTAC, RANITIDINE HYDROCHLORIDE  
ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)  
ZANTAC 150, RANITIDINE HYDROCHLORIDE  
ZANTAC 300, RANITIDINE HYDROCHLORIDE  
ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)  
ZARONTIN, ETHOSUXIMIDE  
ZAROXOLYN, METOLAZONE  
ZAVESCA, MIGLUSTAT  
ZEGERID, OMEPRAZOLE  
ZEGERID OTC, OMEPRAZOLE (OTC)  
ZEJULA, NIRAPARIB TOSYLATE  
ZELAPAR, SELEGILINE HYDROCHLORIDE  
ZELBORAF, VEMURAFENIB  
ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE  
ZEMPLAR, PARICALCITOL  
ZENATANE, ISOTRETINOIN  
ZENPEP, PANCRELIPASE (AMYLASE)  
ZEPATIER, ELBASVIR  
ZERBAXA, CEFTOLOZANE SULFATE

**APPENDIX A - PRODUCT NAME INDEX**

\*\* Z \*\*

ZERIT, STAVUDINE  
ZERVIA, CETIRIZINE HYDROCHLORIDE  
ZESTORETIC, HYDROCHLOROTHIAZIDE  
ZESTRIL, LISINOPRIL  
ZETIA, EZETIMIBE  
ZETONNA, CICLESONIDE  
ZIAC, BISOPROLOL FUMARATE  
ZIAGEN, ABACAVIR SULFATE  
ZIANA, CLINDAMYCIN PHOSPHATE  
ZIDOVUDINE, ZIDOVUDINE  
ZILEUTON, ZILEUTON  
ZILRETTA, TRIAMCINOLONE ACETONIDE  
ZINACEF, CEFUROXIME SODIUM  
ZINACEF IN PLASTIC CONTAINER, CEFUROXIME SODIUM  
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE  
ZINECARD, DEXRAZOXANE HYDROCHLORIDE  
ZINGO, LIDOCAINE HYDROCHLORIDE  
ZIOPTAN, TAFLUPROST  
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE  
ZIPSOR, DICLOFENAC POTASSIUM  
ZIRGAN, GANCICLOVIR  
ZITHROMAX, AZITHROMYCIN  
ZMAX, AZITHROMYCIN  
ZOCOR, SIMVASTATIN  
ZOFRAN, ONDANSETRON HYDROCHLORIDE  
ZOFRAN ODT, ONDANSETRON  
ZOHYDRO ER, HYDROCODONE BITARTRATE  
ZOLADEX, GOSERELIN ACETATE  
ZOLEDRONIC ACID, ZOLEDRONIC ACID  
ZOLINZA, VORINOSTAT  
ZOLMITRIPTAN, ZOLMITRIPTAN  
ZOLOFT, SERTRALINE HYDROCHLORIDE  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE  
ZOLPIMIST, ZOLPIDEM TARTRATE  
ZOMACTON, SOMATROPIN RECOMBINANT  
ZOMETA, ZOLEDRONIC ACID  
ZOMIG, ZOLMITRIPTAN  
ZOMIG-ZMT, ZOLMITRIPTAN  
ZONALON, DOXEPPIN HYDROCHLORIDE  
ZONEGRAN, ZONISAMIDE  
ZONISAMIDE, ZONISAMIDE  
ZONTIVITY, VORAPAXAR SULFATE  
ZORBTIVE, SOMATROPIN RECOMBINANT  
ZORTRESS, EVEROLIMUS  
ZORVOLEX, DICLOFENAC  
ZOSYN, PIPERACILLIN SODIUM  
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM  
ZOVIA 1/35E-28, ETHINYL ESTRADIOL  
ZOVIA 1/50E-28, ETHINYL ESTRADIOL  
ZOVIRAX, ACYCLOVIR  
ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE  
ZUPLENZ, ONDANSETRON  
ZURAMPIC, LESINURAD  
ZUTRIPRO, CHLORPHENIRAMINE MALEATE  
ZYBAN, BUPROPION HYDROCHLORIDE  
ZYCLARA, IMIQUIMOD  
ZYDELIG, IDELALISIB  
ZYFLO, ZILEUTON  
ZYFLO CR, ZILEUTON  
ZYKADIA, CERITINIB  
ZYLET, LOTEPREDNOL ETABONATE  
ZYLOPRIM, ALLOPURINOL  
ZYMAR, GATIFLOXACIN  
ZYMAXID, GATIFLOXACIN  
ZYPITAMAG, PITAVASTATIN MAGNESIUM

**APPENDIX A - PRODUCT NAME INDEX**

\*\* Z \*\*

ZYPREXA, OLANZAPINE  
ZYPREXA RELPREVV, OLANZAPINE PAMOATE  
ZYPREXA ZYDIS, OLANZAPINE  
ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)  
ZYTIGA, ABIRATERONE ACETATE  
ZYVOX, LINEZOLID

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* 3 \*\*

**3D IMAGING DRUG**

- \* 3D IMAGING DRUG DESIGN AND DEVELOPMENT LLC  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**3M**

- \* 3M CO  
PERIDEX, CHLORHEXIDINE GLUCONATE
- \* 3M HEALTH CARE INC  
AVAGARD, ALCOHOL (OTC)  
DURAPREP, IODINE POVACRYLEX (OTC)

**3M DRUG DELIVERY**

- \* 3M DRUG DELIVERY SYSTEMS  
FENTANYL-100, FENTANYL  
FENTANYL-12, FENTANYL  
FENTANYL-25, FENTANYL  
FENTANYL-50, FENTANYL  
FENTANYL-75, FENTANYL  
PROVENTIL-HFA, ALBUTEROL SULFATE

\*\* A \*\*

**AAA USA INC**

- \* ADVANCED ACCELERATOR APPLICATIONS USA INC  
NETSPOT, GALLIUM DOTATATE GA-68

**AAIPHARMA LLC**

- \* AAIPHARMA LLC  
AZASAN, AZATHIOPRINE

**ABBVIE**

- \* ABBVIE INC  
ANDROGEL, TESTOSTERONE  
BIAXIN, CLARITHROMYCIN  
CREON, PANCRELIPASE (AMYLASE)  
CYCLOSPORINE, CYCLOSPORINE  
DEPACON, VALPROATE SODIUM  
DEPAKENE, VALPROIC ACID  
DEPAKOTE ER, DIVALPROEX SODIUM  
DEPAKOTE, DIVALPROEX SODIUM  
GENGRAF, CYCLOSPORINE  
K-TAB, POTASSIUM CHLORIDE  
KALETRA, LOPINAVIR  
MARINOL, DRONABINOL  
MAVIK, TRANDOLAPRIL  
MIVACRON, MIVACURIUM CHLORIDE  
NIASPAN, NIACIN  
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE  
NIMBEX, CISATRACURIUM BESYLATE  
NORVIR, RITONAVIR  
SURVANTA, BERACTANT  
SYNTHROID, LEVOTHYROXINE SODIUM \*\*  
TARKA, TRANDOLAPRIL  
TEVETEN, EPROSARTAN MESYLATE  
TRICOR, FENOFIBRATE  
TRIDIONE, TRIMETHADIONE  
TRILIPIX, CHOLINE FENOFIBRATE  
ULTANE, SEVOFLURANE  
ZEMPLAR, PARICALCITOL

**ABBVIE ENDOCRINE**

- \* ABBVIE ENDOCRINE INC  
LUPANETA PACK, LEUPROLIDE ACETATE

**ABBVIE ENDOCRINE INC**

- \* ABBVIE ENDOCRINE INC  
LUPRON DEPOT, LEUPROLIDE ACETATE  
LUPRON DEPOT-PED, LEUPROLIDE ACETATE

**ABBVIE INC**

- \* ABBVIE INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ABBVIE INC
  - DUOPA, CARBIDOPA
  - Mavyret, GLECAPREVIR
  - NORVIR, RITONAVIR
  - TECHNIVIE, OMBITASVIR
  - VENCLEXTA, VENETOCLAX
  - VIEKIRA PAK (COPACKAGED), DASABUVIR SODIUM
  - VIEKIRA XR, DASABUVIR SODIUM

**ABHAI INC**

- \* ABHAI INC
  - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

**ABHAI LLC**

- \* ABHAI LLC
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

**ABON PHARMS LLC**

- \* ABON PHARMACEUTICALS LLC
  - CLOFARABINE, CLOFARABINE

**ABRAXIS BIOSCIENCE**

- \* ABRAXIS BIOSCIENCE LLC
  - ABRAXANE, PACLITAXEL

**ABRAXIS PHARM**

- \* ABRAXIS PHARMACEUTICAL PRODUCTS
  - CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE

**ACADIA PHARMS INC**

- \* ACADIA PHARMACEUTICALS INC
  - NUPLAZID, PIMAVANSERIN TARTRATE

**ACCELRX LABS**

- \* ACCELRX LABS LLC
  - CARISOPRODOL, CARISOPRODOL

**ACCORD HLTHCARE**

- \* ACCORD HEALTHCARE INC
  - ALLOPURINOL, ALLOPURINOL
  - AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - ANASTROZOLE, ANASTROZOLE
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
  - BICALUTAMIDE, BICALUTAMIDE
  - BIVALIRUDIN, BIVALIRUDIN
  - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
  - CAPECITABINE, CAPECITABINE
  - CARBIDOPA AND LEVODOPA, CARBIDOPA
  - CARBOPLATIN, CARBOPLATIN
  - CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
  - CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
  - CISPLATIN, CISPLATIN
  - CLONAZEPAM, CLONAZEPAM
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - CLOZAPINE, CLOZAPINE
  - DECITABINE, DECITABINE
  - DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DOCETAXEL, DOCETAXEL
  - DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
  - DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
  - ENTECAVIR, ENTECAVIR
  - EPLERENONE, EPLERENONE
  - EPTIFIBATIDE, EPTIFIBATIDE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ACCORD HEALTHCARE INC
  - ETOPOSIDE, ETOPOSIDE
  - FINASTERIDE, FINASTERIDE
  - FLUOROURACIL, FLUOROURACIL
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - GLIMEPIRIDE, GLIMEPIRIDE
  - GLIPIZIDE, GLIPIZIDE
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
  - ITRACONAZOLE, ITRACONAZOLE
  - LETROZOLE, LETROZOLE
  - LEVETIRACETAM, LEVETIRACETAM
  - LISINOPRIL, LISINOPRIL
  - METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
  - METHYLDOPA, METHYLDOPA
  - MITOMYCIN, MITOMYCIN
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
  - MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
  - NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
  - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
  - NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - NORETHINDRONE, NORETHINDRONE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXALIPLATIN, OXALIPLATIN
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - PARICALCITOL, PARICALCITOL
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RAMIPRIL, RAMIPRIL
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - ROSVASTATIN CALCIUM, ROUVASTATIN CALCIUM
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SIMVASTATIN, SIMVASTATIN
  - SPIRONOLACTONE, SPIRONOLACTONE
  - TACROLIMUS, TACROLIMUS
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - TOPIRAMATE, TOPIRAMATE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**ACELLA PHARMS LLC**

- \* ACELLA PHARMACEUTICALS LLC
  - BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
  - GABAPENTIN, GABAPENTIN
  - HYDROCODONE BITARTRATE AND CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE
  - PHENYTOIN SODIUM, PHENYTOIN SODIUM

**ACI HEALTHCARE LTD**

- \* ACI HEALTHCARE LTD
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - GABAPENTIN, GABAPENTIN
  - LEVETIRACETAM, LEVETIRACETAM
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

**ACIC FINE CHEMS**

- \* ACIC FINE CHEMICALS INC
  - TRANEXAMIC ACID, TRANEXAMIC ACID

**ACIC PHARMS**

- \* ACIC PHARMACEUTICALS INC
  - LEVETIRACETAM, LEVETIRACETAM
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

**ACLARIS THERAPS INC**

- \* ACLARIS THERAPEUTICS INC  
ESKATA, HYDROGEN PEROXIDE

**ACORDA**

- \* ACORDA THERAPEUTICS INC  
AMPYRA, DALFAMPRIDINE  
QUTENZA, CAPSAICIN

**ACS DOBFAR**

- \* ACS DOBFAR SPA  
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM  
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM  
CEFEPEMIE HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE  
CEFOXITIN, CEFOXITIN SODIUM  
CEFTAZIDIME, CEFTAZIDIME  
CEFTRIAXONE, CEFTRIAXONE SODIUM  
IMIPENEM AND CILASTATIN, CILASTATIN SODIUM  
KEFZOL, CEFAZOLIN SODIUM  
MEROPENEM, MEROPENEM

**ACS DOBFAR INFO SA**

- \* ACS DOBFAR INFO SA  
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE  
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN  
ZOLEDRONIC ACID, ZOLEDRONIC ACID

**ACS DOBFAR SPA**

- \* ACS DOBFAR SPA  
AMPICILLIN SODIUM, AMPICILLIN SODIUM  
CEFUROXIME SODIUM, CEFUROXIME SODIUM

**ACTAVIS ELIZABETH**

- \* ACTAVIS ELIZABETH LLC  
ALPRAZOLAM, ALPRAZOLAM  
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
CARBIDOPA AND LEVODOPA, CARBIDOPA  
CLONAZEPAM, CLONAZEPAM  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
DEFERASIROX, DEFERASIROX  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
DEXTROMETHORPHAN HYDROBROMIDE AND QUINIDINE SULFATE, DEXTROMETHORPHAN HYDROBROMIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXE PIN HYDROCHLORIDE, DOXE PIN HYDROCHLORIDE  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
FENOFIBRIC ACID, CHOLINE FENOFIBRATE  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
GABAPENTIN, GABAPENTIN  
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE  
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
INDAPAMIDE, INDAPAMIDE  
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
LOVASTATIN, LOVASTATIN  
METOPROLOL SUCCINATE, METOPROLOL SUCCINATE  
NIFEDIPINE, NIFEDIPINE  
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)  
OXAZEPAM, OXAZEPAM  
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ACTAVIS ELIZABETH LLC
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
  - PROPYLTHIOURACIL, PROPYLTHIOURACIL
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SPIRONOLACTONE, SPIRONOLACTONE
  - TEMAZEPAM, TEMAZEPAM
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- \* ACTAVIS ELIZABETH LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - ALPRAZOLAM, ALPRAZOLAM
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
  - LAMOTRIGINE, LAMOTRIGINE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

**ACTAVIS INC**

- \* ACTAVIS INC
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - METHOXSALEN, METHOXSALEN
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**ACTAVIS LABS**

- \* ACTAVIS LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - PERMETHRIN, PERMETHRIN

**ACTAVIS LABS FL**

- \* ACTAVIS LABORATORIES FL INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - DESVENLAFAKINE SUCCINATE, DESVENLAFAKINE SUCCINATE
  - GUAIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
  - GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
  - GUAIFENESIN, GUAIFENESIN (OTC)

**ACTAVIS LABS FL INC**

- \* ACTAVIS LABORATORIES FL INC
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CABERGOLINE, CABERGOLINE
  - CARTIA XT, DILTIAZEM HYDROCHLORIDE
  - CLARITHROMYCIN, CLARITHROMYCIN
  - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
  - DALFAMPRIDINE, DALFAMPRIDINE
  - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
  - DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
  - DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE
  - DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE
  - DUTASTERIDE, DUTASTERIDE
  - FENTANYL CITRATE, FENTANYL CITRATE
  - FLUTAMIDE, FLUTAMIDE
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - KETOPROFEN, KETOPROFEN
  - LEVETIRACETAM, LEVETIRACETAM
  - LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
  - LORATADINE, LORATADINE (OTC)
  - METAXALONE, METAXALONE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
  - MIRTAZAPINE, MIRTAZAPINE
  - NAPROXEN SODIUM, NAPROXEN SODIUM
  - NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
  - NITROGLYCERIN, NITROGLYCERIN
  - OMEPRAZOLE, OMEPRAZOLE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ACTAVIS LABORATORIES FL INC  
OXYCODONE AND ASPIRIN, ASPIRIN  
PALIPERIDONE, PALIPERIDONE  
PAROXETINE MESYLATE, PAROXETINE MESYLATE  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
PREDNISONE, PREDNISONE  
RAMELTEON, RAMELTEON  
RISPERIDONE, RISPERIDONE  
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE  
TAZTIA XT, DILTIAZEM HYDROCHLORIDE  
TETRABENAZINE, TETRABENAZINE  
TRANEXAMIC ACID, TRANEXAMIC ACID  
TROSPiUM CHLORIDE, TROSPiUM CHLORIDE  
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**ACTAVIS LABS NY INC**

- \* ACTAVIS LABORATORIES NY INC  
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)

**ACTAVIS LABS UT INC**

- \* ACTAVIS LABORATORIES UT INC  
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
CLONIDINE, CLONIDINE  
EMLA, LIDOCAINE  
FIORICET W/ CODEINE, ACETAMINOPHEN  
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE  
LIDOCAINE, LIDOCAINE  
NORINYL 1+50 28-DAY, MESTRANOL  
PROGESTERONE, PROGESTERONE  
TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE  
TENUATE, DIETHYLPROPION HYDROCHLORIDE  
TESTOSTERONE, TESTOSTERONE
- \* ACTAVIS LABORATORIES UT INC INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC  
FENTANYL-100, FENTANYL  
FENTANYL-25, FENTANYL  
FENTANYL-50, FENTANYL  
FENTANYL-75, FENTANYL  
TESTOSTERONE, TESTOSTERONE

**ACTAVIS LLC**

- \* ACTAVIS LLC  
AZACITIDINE, AZACITIDINE  
DAPSONE, DAPSONE  
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE  
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM  
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
- \* ACTAVIS LLC AN INDIRECT WHOLLY-OWNED SUB OF TEVA PHARMACEUTICALS USA INC  
BUSULFAN, BUSULFAN  
DOCETAXEL, DOCETAXEL  
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN

**ACTAVIS MID ATLANTIC**

- \* ACTAVIS MID ATLANTIC LLC  
ACETASOL HC, ACETIC ACID, GLACIAL  
ACYCLOVIR, ACYCLOVIR  
ADAPALENE AND BENZOYL PEROXIDE, ADAPALENE  
ADAPALENE, ADAPALENE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
CICLOPIROX, CICLOPIROX  
CLINDAMYCIN PHOSPHATE AND TRETINOIN, CLINDAMYCIN PHOSPHATE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLOTrimazole AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CLOTrimazole, CLOTrimazole (OTC)  
DESOXIMETASONE, DESOXIMETASONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ACTAVIS MID ATLANTIC LLC
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM  
ENULOSE, LACTULOSE
  - FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
  - GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
  - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
  - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
  - HYDROCORTISONE, HYDROCORTISONE
  - IBUPROFEN, IBUPROFEN (OTC)
  - LEVETIRACETAM, LEVETIRACETAM
  - MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
  - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
  - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
  - NITROFURANTOIN, NITROFURANTOIN
  - NYSTATIN, NYSTATIN
  - PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
  - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - VALNAC, BETAMETHASONE VALERATE
- \* ACTAVIS MID ATLANTIC LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - IBUPROFEN, IBUPROFEN
  - M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
  - PERMETHRIN, PERMETHRIN (OTC)

**ACTAVIS PHARMA**

- \* ACTAVIS PHARMA INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - MICONAZOLE NITRATE, MICONAZOLE NITRATE
  - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)

**ACTAVIS TOTOWA**

- \* ACTAVIS TOTOWA LLC
  - DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
  - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
  - FINASTERIDE, FINASTERIDE
  - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - OXALIPLATIN, OXALIPLATIN
  - PACLITAXEL, PACLITAXEL
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - REPAGLINIDE, REPAGLINIDE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - VINORELBINE TARTRATE, VINORELBINE TARTRATE

**ACTAVIS TOTOWA TEVA**

- \* ACTAVIS TOTOWA LLC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - FINASTERIDE, FINASTERIDE

**ACTELION PHARMS**

- \* ACTELION PHARMACEUTICALS LTD
  - TRACLEER, BOSENTAN

**ACTELION PHARMS LTD**

- \* ACTELION PHARMACEUTICALS LTD
  - OPSUMIT, MACITENTAN
  - TRACLEER, BOSENTAN
  - UPTRAVI, SELEXIPAG
  - VALCHLOR, MECHLORETHAMINE HYDROCHLORIDE
  - VELETRI, EPOPROSTENOL SODIUM
  - VENTAVIS, ILOPROST
  - ZAVESCA, MIGLUSTAT

**ACTIENT PHARMS**

- \* ACTIENT PHARMACEUTICALS LLC
  - THEO-24, THEOPHYLLINE

**ADAMAS PHARMA**

- \* ADAMAS PHARMA LLC
  - GOCOVRI, AMANTADINE HYDROCHLORIDE

**ADAMIS PHARMS CORP**

- \* ADAMIS PHARMACEUTICALS CORP
  - SYMJEPI, EPINEPHRINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

**ADAPT**

- \* ADAPT PHARMA OPERATIONS LTD  
NARCAN, NALOXONE HYDROCHLORIDE

**ADARE PHARMS INC**

- \* ADARE PHARMACEUTICALS INC  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

**ADDMEDICA SAS**

- \* ADDMEDICA SAS  
SIKLOS, HYDROXYUREA

**ADIENNE SA**

- \* ADIENNE SA  
TEPADINA, THIOTEPA

**AEGERION**

- \* AEGERION PHARMACEUTICALS INC  
JUXTAPID, LOMITAPIDE MESYLATE

**AERIE PHARMS INC**

- \* AERIE PHARMACEUTICALS INC  
RHOPRESSA, NETARSUDIL DIMESYLADE

**AETERNA ZENTARIS**

- \* AETERNA ZENTARIS GMBH  
MACRILEN, MACIMORELIN ACETATE

**AGOURON PHARMS**

- \* AGOURON PHARMACEUTICALS LLC  
VIRACEPT, NELFINAVIR MESYLATE

**AILEX PHARMS LLC**

- \* AILEX PHARMACEUTICALS LLC  
CROMOLYN SODIUM, CROMOLYN SODIUM  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)  
SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

**AIPIING PHARM INC**

- \* AIPIING PHARMACEUTICAL INC  
FOLIC ACID, FOLIC ACID  
IBUPROFEN, IBUPROFEN  
IBUPROFEN, IBUPROFEN (OTC)

**AJANTA PHARMA**

- \* AJANTA PHARMA LTD  
LEVETIRACETAM, LEVETIRACETAM

**AJANTA PHARMA LTD**

- \* AJANTA PHARMA LTD  
ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE  
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLADE  
ARIPIPRAZOLE, ARIPIPRAZOLE  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE  
ENTACAPONE, ENTACAPONE  
LANSOPRAZOLE, LANSOPRAZOLE  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
OLANZAPINE, OLANZAPINE  
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE  
RISPERIDONE, RISPERIDONE  
RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE  
VORICONAZOLE, VORICONAZOLE  
ZOLMITRIPTAN, ZOLMITRIPTAN

**AKORN**

- \* AKORN INC  
ADENOSINE, ADENOSINE  
AK-FLUOR 10%, FLUORESCEIN SODIUM  
AK-FLUOR 25%, FLUORESCEIN SODIUM  
AKBETA, LEVOBUNOLOL HYDROCHLORIDE  
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AKORN INC
  - AKTEN, LIDOCAINE HYDROCHLORIDE
  - AKTOB, TOBRAMYCIN
  - ALFENTA, ALFENTANIL HYDROCHLORIDE
  - AMINOCAPROIC ACID, AMINOCAPROIC ACID
  - AMINOCAPROIC, AMINOCAPROIC ACID
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - ATROPINE SULFATE, ATROPINE SULFATE
  - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
  - BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
  - BACITRACIN, BACITRACIN
  - BAL, DIMERCAPROL
  - BALANCED SALT, CALCIUM CHLORIDE
  - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
  - BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
  - CALCITRIOL, CALCITRIOL
  - CAPASTAT SULFATE, CAPREOMYCIN SULFATE
  - CARBOPLATIN, CARBOPLATIN
  - CICLOPIROX, CICLOPIROX
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - CROMOLYN SODIUM, CROMOLYN SODIUM
  - DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
  - DESOXIMETASONE, DESOXIMETASONE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
  - ENDOSOL EXTRA, CALCIUM CHLORIDE
  - EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE
  - EPTIFIBATIDE, EPTIFIBATIDE
  - ERYTHRHYMYCIN, ERYTHRHYMYCIN
  - ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
  - ETHOSUXIMIDE, ETHOSUXIMIDE
  - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
  - GENTAK, GENTAMICIN SULFATE
  - GENTAMICIN SULFATE, GENTAMICIN SULFATE
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - IC-GREEN, INDOCYANINE GREEN
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)
  - LATANOPROST, LATANOPROST
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
  - LORAZEPAM, LORAZEPAM
  - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
  - NEDOCROMIL SODIUM, NEDOCROMIL SODIUM
  - NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
  - NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - OFLOXACIN, OFLOXACIN
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
  - PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
  - PARICALCITOL, PARICALCITOL
  - PYRAZINAMIDE, PYRAZINAMIDE
  - RIFAMPIN, RIFAMPIN
  - SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
  - SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
  - SUFENTA PRESERVATIVE FREE, SUFENTANIL CITRATE
  - SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
  - TERBUTALINE SULFATE, TERBUTALINE SULFATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AKORN INC
  - TIMOLOL MALEATE, TIMOLOL MALEATE
  - TIMOLOL, TIMOLOL
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - TRANEXAMIC ACID, TRANEXAMIC ACID
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
  - TROPICACYL, TROPICAMIDE
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VORICONAZOLE, VORICONAZOLE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**AKORN INC**

- \* AKORN INC
  - ACETYLCYSTEINE, ACETYLCYSTEINE
  - APRACLONIDINE HYDROCHLORIDE, APRACLONIDINE HYDROCHLORIDE
  - CEFTRIAXONE, CEFTRIAXONE SODIUM
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
  - CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
  - DOXERCALCIFEROL, DOXERCALCIFEROL
  - DRONABINOL, DRONABINOL
  - EPHEDRINE SULFATE, EPHEDRINE SULFATE
  - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - INAPSINE, DROPERIDOL
  - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
  - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
  - MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
  - NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)
  - NAPHAZOLINE HYDROCHLORIDE, NAPHAZOLINE HYDROCHLORIDE
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
  - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
  - PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
  - ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
  - TOBRAMYCIN, TOBRAMYCIN
  - TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**ALCON**

- \* ALCON LABORATORIES INC
  - BETOPTIC, BETAXOLOL HYDROCHLORIDE
  - BSS PLUS, CALCIUM CHLORIDE
  - BSS, CALCIUM CHLORIDE
  - CROMOLYN SODIUM, CROMOLYN SODIUM
  - MIOSTAT, CARBACHOL
  - NAPHCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)

**ALCON PHARMS LTD**

- \* ALCON PHARMACEUTICALS LTD
  - BETADINE, POVIDONE-IODINE
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - KETOTIFEN FUMARATE, KETOTIFEN FUMARATE (OTC)

**ALEMBIC LTD**

- \* ALEMBIC LTD
  - LITHIUM CARBONATE, LITHIUM CARBONATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

**ALEMBIC PHARMS LTD**

- \* ALEMBIC PHARMACEUTICALS LTD
  - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
  - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ALEMBIC PHARMACEUTICALS LTD
  - CELECOXIB, CELECOXIB
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
  - DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
  - DESVENLAFAKINE SUCCINATE, DESVENLAFAKINE SUCCINATE
  - DESVENLAFAKINE, DESVENLAFAKINE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DOXYCYCLINE, DOXYCYCLINE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - FAMOTIDINE, FAMOTIDINE
  - FENOFIBRIC ACID, CHOLINE FENOFIBRATE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IRBESARTAN, IRBESARTAN
  - ITRACONAZOLE, ITRACONAZOLE
  - LAMOTRIGINE, LAMOTRIGINE
  - LEFLUNOMIDE, LEFLUNOMIDE
  - LINEZOLID, LINEZOLID
  - LITHIUM CARBONATE, LITHIUM CARBONATE
  - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - MEPROBAMATE, MEPROBAMATE
  - METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - METRONIDAZOLE, METRONIDAZOLE
  - MODAFINIL, MODAFINIL
  - OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
  - TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TELMISARTAN, TELMISARTAN
  - THEOPHYLLINE, THEOPHYLLINE
  - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - VALSARTAN, VALSARTAN
  - VENLAFAKINE HYDROCHLORIDE, VENLAFAKINE HYDROCHLORIDE
  - ZOLMITRIPTAN, ZOLMITRIPTAN

**ALIMERA SCIENCES INC**

- \* ALIMERA SCIENCES INC
  - ILUVIEN, FLUOCINOLONE ACETONIDE

**ALKEM**

- \* ALKEM LABORATORIES LTD
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - GABAPENTIN, GABAPENTIN
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**ALKEM LABS LTD**

- \* ALKEM LABORATORIES LTD
  - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
  - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
  - CAPECITABINE, CAPECITABINE
  - CEFUROXIME AXETIL, CEFUROXIME AXETIL
  - CEPHALEXIN, CEPHALEXIN
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - EZETIMIBE AND SIMVASTATIN, EZETIMIBE
  - EZETIMIBE, EZETIMIBE
  - FINASTERIDE, FINASTERIDE
  - GABAPENTIN, GABAPENTIN
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - ITRACONAZOLE, ITRACONAZOLE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ALKEM LABORATORIES LTD
  - LAMOTRIGINE, LAMOTRIGINE
  - LIDOCaine, LIDOCaine
  - LINEZOLID, LINEZOLID
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
  - OLANZAPINE, OLANZAPINE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RASAGILINE MESYLATE, RASAGILINE MESYLATE
  - RILUZOLE, RILUZOLE
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - ROSVASTATIN CALCIUM, ROUVASTATIN CALCIUM
  - TAMSULOSIN HYDROCHLORIDE , TAMSULOSIN HYDROCHLORIDE
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**ALKERMES**

- \* ALKERMES INC
  - VIVITROL, NALTREXONE

**ALKERMES INC**

- \* ALKERMES INC
  - ARISTADA, ARIPIPRAZOLE LAUROXIL

**ALLEGIANCE HLTHCARE**

- \* ALLEGIANCE HEALTHCARE CORP
  - POVIDONE IODINE, POVIDONE-IODINE (OTC)

**ALLEGIS**

- \* ALLEGIS HOLDINGS LLC
  - PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE

**ALLERGAN**

- \* ALLERGAN
  - ACULAR LS, KETOROLAC TROMETHAMINE
  - ALPHAGAN P, BRIMONIDINE TARTRATE
  - BLEPH-10, SULFACETAMIDE SODIUM
  - GENOPTIC, GENTAMICIN SULFATE
  - ZYMAXID, GATIFLOXACIN
- \* ALLERGAN INC
  - ACULAR, KETOROLAC TROMETHAMINE
  - ACUVAIL, KETOROLAC TROMETHAMINE
  - ACZONE, DAPSONE
  - ALOCRIL, NEDOCROMIL SODIUM
  - ALPHAGAN P, BRIMONIDINE TARTRATE
  - AVAGE, TAZAROTENE
  - AZELEX, AZELAIC ACID
  - COMBIGAN, BRIMONIDINE TARTRATE
  - ELESTAT, EPINASTINE HYDROCHLORIDE
  - LASTACAFT, ALCAFTADINE
  - LATISSE, BIMATOPROST
  - LUMIGAN, BIMATOPROST
  - OCUFLOX, OFLOXACIN
  - OZURDEX, DEXAMETHASONE
  - POLYTRIM, POLYMYXIN B SULFATE
  - RESTASIS MULTIDOSE, CYCLOSPORINE
  - RESTASIS, CYCLOSPORINE
  - TAZORAC, TAZAROTENE
  - ZYMAR, GATIFLOXACIN
- \* ALLERGAN PHARMACEUTICAL
  - BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
  - BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
  - BLEPHAMIDE, PREDNISOLONE ACETATE
  - FML FORTE, FLUOROMETHOLONE
  - FML, FLUOROMETHOLONE
  - OCUFEN, FLURBIPROFEN SODIUM
  - PRED FORTE, PREDNISOLONE ACETATE
  - PRED MILD, PREDNISOLONE ACETATE
  - PRED-G, GENTAMICIN SULFATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

**ALLERGAN HOLDINGS**

- \* ALLERGAN HOLDINGS UNLTD CO  
VIBERZI, ELUXADOLINE

**ALLERGAN INC**

- \* ALLERGAN INC  
ACZONE, DAPSONE  
RHOFADE, OXYMETAZOLINE HYDROCHLORIDE

**ALLERGAN SALES LLC**

- \* ALLERGAN SALES LLC  
ACTIGALL, URSDIOL  
ALORA, ESTRADIOL  
ANDRODERM, TESTOSTERONE  
BREVICON 28-DAY, ETHINYL ESTRADIOL  
CONDYLOX, PODOFILOX  
CORDRAN, FLURANDRENOLIDE  
CRINONE, PROGESTERONE  
DALVANCE, DALBAVANCIN HYDROCHLORIDE  
ESTRACE, ESTRADIOL  
FIORINAL W/CODEINE, ASPIRIN  
FIORINAL, ASPIRIN  
GELNIQUE, OXYBUTYNIN CHLORIDE  
INFED, IRON DEXTRAN  
KADIAN, MORPHINE SULFATE  
MICROZIDE, HYDROCHLOROTHIAZIDE  
NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL  
NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL  
OXYTROL FOR WOMEN, OXYBUTYNIN (OTC)  
OXYTROL, OXYBUTYNIN  
RAPAFLO, SILODOSIN  
SAVELLA, MILNACIPRAN HYDROCHLORIDE  
TRELSTAR, TRIPTORELIN PAMOATE

**ALLERQUEST**

- \* ALLERQUEST LLC  
PRE-PEN, BENZYL PENICILLOYL POLYLYSINE

**ALLIED PHARMA INC**

- \* ALLIED PHARMA INC  
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CLARITHROMYCIN, CLARITHROMYCIN  
LEVETIRACETAM, LEVETIRACETAM  
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

**ALLOS**

- \* ALLOS THERAPEUTICS INC  
FOLOTYN, PRALATREXATE

**ALPHARMA PHARMS**

- \* ALPHARMA PHARMACEUTICALS LLC  
EMBEDA, MORPHINE SULFATE

**ALRA**

- \* ALRA LABORATORIES INC  
CHOLAC, LACTULOSE  
CONSTILAC, LACTULOSE  
GEN-XENE, CLORAZEPATE DIPOTASSIUM  
IBU-TAB 200, IBUPROFEN (OTC)  
IBU-TAB, IBUPROFEN

**ALTAIRE PHARMS INC**

- \* ALTAIRE PHARMACEUTICALS INC  
ALTAFLUOR BENOX, BENOXINATE HYDROCHLORIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
NAPHAZOLINE HYDROCHLORIDE AND PHENIRAMINE MALEATE, NAPHAZOLINE HYDROCHLORIDE (OTC)  
OFLOXACIN, OFLOXACIN

**ALTATHERA PHARMS LLC**

- \* ALTATHERA PHARMACEUTICALS LLC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ALTATHERA PHARMACEUTICALS LLC  
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

**ALVOGEN**

- \* ALVOGEN GROUP HOLDINGS 2 LLC  
DAPSONE, DAPSONE  
OFLOXACIN, OFLOXACIN
- \* ALVOGEN GROUP HOLDINGS 3 LLC  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
FORFIVO XL, BUPROPION HYDROCHLORIDE
- \* ALVOGEN GROUP HOLDINGS LLC  
ADALAT CC, NIFEDIPINE
- \* ALVOGEN INC  
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

**ALVOGEN INC**

- \* ALVOGEN INC  
ACETYLCYSTEINE, ACETYLCYSTEINE  
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE  
VORICONAZOLE, VORICONAZOLE

**ALVOGEN MALTA**

- \* ALVOGEN MALTA OPERATIONS LTD  
ATENOLOL AND CHLORTHALIDONE, ATENOLOL  
ATENOLOL, ATENOLOL  
BUDESONIDE, BUDESONIDE  
CARBIDOPA, CARBIDOPA  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DISULFIRAM, DISULFIRAM  
EXEMESTANE, EXEMESTANE  
FELBAMATE, FELBAMATE  
MACROBID, NITROFURANTOIN  
MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE  
MELPHALAN, MELPHALAN  
NAPRELAN, NAPROXEN SODIUM  
NATEGLINIDE, NATEGLINIDE  
NEVIRAPINE, NEVIRAPINE  
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE  
RIVASTIGMINE, RIVASTIGMINE  
SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE  
SPECTAZOLE, ECONAZOLE NITRATE  
TENORETIC 100, ATENOLOL  
TENORETIC 50, ATENOLOL  
TENORMIN, ATENOLOL  
ZESTORETIC, HYDROCHLOROTHIAZIDE  
ZESTRIL, LISINOPRIL

**ALVOGEN PINE BROOK**

- \* ALVOGEN PINE BROOK LLC  
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE

**AM ANTIBIOTICS**

- \* AMERICAN ANTIBIOTICS INC  
AMOXICILLIN, AMOXICILLIN

**AMAG PHARMA USA**

- \* AMAG PHARMA USA INC  
MAKENA PRESERVATIVE FREE, HYDROXYPROGESTERONE CAPROATE  
MAKENA, HYDROXYPROGESTERONE CAPROATE

**AMAG PHARMS INC**

- \* AMAG PHARMACEUTICALS INC  
FERAHME, FERUMOXYTOL  
INTRAROSA, PRASTERONE

**AMARIN PHARMS**

- \* AMARIN PHARMACEUTICALS IRELAND LTD  
VASCEPA, ICOSAPENT ETHYL

**AMERIGEN PHARMS LTD**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AMERIGEN PHARMACEUTICALS LTD  
CARBIDOPA, CARBIDOPA  
INDAPAMIDE, INDAPAMIDE  
TEMOZOLOMIDE, TEMOZOLOMIDE

**AMGEN**

- \* AMGEN INC  
SENSIPAR, CINACALCET HYDROCHLORIDE

**AMGEN INC**

- \* AMGEN INC  
CORLANOR, IVABRADINE HYDROCHLORIDE

**AMNEAL PHARM**

- \* AMNEAL PHARMACEUTICAL  
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE  
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE  
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE  
FLECAINIDE ACETATE, FLECAINIDE ACETATE  
FOLIC ACID, FOLIC ACID  
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
PRIMIDONE, PRIMIDONE

**AMNEAL PHARMS**

- \* AMNEAL PHARMACEUTICALS  
ACYCLOVIR, ACYCLOVIR  
ALBUTEROL SULFATE, ALBUTEROL SULFATE  
ARIPIPRAZOLE, ARIPIPRAZOLE  
ATOVAQUONE, ATOVAQUONE  
CALCITRIOL, CALCITRIOL  
CALCIUM ACETATE, CALCIUM ACETATE  
CAPECITABINE, CAPECITABINE  
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE  
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE  
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
ENTECAVIR, ENTECAVIR  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
ESTRADIOL, ESTRADIOL  
FELBAMATE, FELBAMATE  
GABAPENTIN, GABAPENTIN  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
INDOMETHACIN, INDOMETHACIN  
ITRACONAZOLE, ITRACONAZOLE  
LEVETIRACETAM, LEVETIRACETAM  
LIDOCAINE, LIDOCAINE  
LINEZOLID, LINEZOLID  
LORAZEPAM, LORAZEPAM  
MEMANTINE HYDROCHLORIDE AND DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
MEROPENEM, MEROPENEM  
METAZALONE, METAZALONE  
MILNACIPRAN HYDROCHLORIDE, MILNACIPRAN HYDROCHLORIDE  
MOMETASONE FUROATE, MOMETASONE FUROATE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
NIACIN, NIACIN  
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN  
NITROFURANTOIN, NITROFURANTOIN  
NIZATIDINE, NIZATIDINE  
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
OXCARBAZEPINE, OXCARBAZEPINE  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

## \* AMNEAL PHARMACEUTICALS

POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
 PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE  
 PROMETHAZINE HYDROCHLORIDE AND PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE  
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
 PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE  
 QUININE SULFATE, QUININE SULFATE  
 RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM  
 RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE  
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
 RISPERIDONE, RISPERIDONE  
 SILDENAFIL CITRATE, SILDENAFIL CITRATE  
 TELMISARTAN, TELMISARTAN  
 TEMAZEPAM, TEMAZEPAM  
 TEMOZOLOMIDE, TEMOZOLOMIDE  
 TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN  
 VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE  
 VORICONAZOLE, VORICONAZOLE  
 WARFARIN SODIUM, WARFARIN SODIUM

## \* AMNEAL PHARMACEUTICALS HOLDINGS GMBH

DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

## \* AMNEAL PHARMACEUTICALS OF NEW YORK LLC

ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE  
 ASPIRIN AND DIPYRIDAMOLE, ASPIRIN  
 BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
 BUDESONIDE, BUDESONIDE  
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
 BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
 DUTASTERIDE, DUTASTERIDE  
 EPTIFIBATIDE, EPTIFIBATIDE  
 IBUPROFEN, IBUPROFEN (OTC)  
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE  
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
 METHOTREXATE SODIUM, METHOTREXATE SODIUM  
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL  
 NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL  
 NORETHINDRONE, NORETHINDRONE  
 NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
 OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS  
 OSeltamivir Phosphate, OSeltamivir Phosphate  
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
 PARICALCITOL, PARICALCITOL  
 SPIRONOLACTONE, SPIRONOLACTONE  
 TOBRAMYCIN, TOBRAMYCIN  
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
 VALSARTAN, VALSARTAN

**AMNEAL PHARMS CO**

## \* AMNEAL PHARMACEUTICALS CO GMBH

BUMETANIDE, BUMETANIDE  
 BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE  
 BUSULFAN, BUSULFAN  
 CLOFARABINE, CLOFARABINE  
 DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE  
 DOXEPEPIN HYDROCHLORIDE, DOXEPEPIN HYDROCHLORIDE  
 DOXERCALCIFEROL, DOXERCALCIFEROL  
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
 EZETIMIBE AND SIMVASTATIN, EZETIMIBE  
 EZETIMIBE, EZETIMIBE  
 FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM  
 FUROSEMIDE, FUROSEMIDE  
 GLYCOPYRROLATE, GLYCOPYRROLATE  
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM  
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AMNEAL PHARMACEUTICALS CO GMBH  
NADOLOL, NADOLOL  
OXAPROZIN, OXAPROZIN  
PARICALCITOL, PARICALCITOL  
SEVELAMER CARBONATE, SEVELAMER CARBONATE  
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE  
TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE  
TRANEXAMIC ACID, TRANEXAMIC ACID  
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**AMNEAL PHARMS LLC**

- \* AMNEAL PHARMACEUTICALS LLC  
ACTIVELLA, ESTRADIOL  
AZATHIOPRINE, AZATHIOPRINE  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)

**AMNEAL PHARMS NY**

- \* AMNEAL PHARMACEUTICALS NY LLC  
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN  
ALPRAZOLAM, ALPRAZOLAM  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM  
GABAPENTIN, GABAPENTIN  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE  
IBUPROFEN, IBUPROFEN  
IBUPROFEN, IBUPROFEN (OTC)  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
NAPROXEN SODIUM, NAPROXEN SODIUM  
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)  
NAPROXEN, NAPROXEN  
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
REPREXAIN, HYDROCODONE BITARTRATE  
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

- \* AMNEAL PHARMACEUTICALS OF NY LLC  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
ISOTRETINOIN, ISOTRETINOIN  
PROGESTERONE, PROGESTERONE

**AMPHASTAR PHARM**

- \* AMPHASTAR PHARMACEUTICAL INC  
AMPHADASE, HYALURONIDASE  
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

**AMPHASTAR PHARMS INC**

- \* AMPHASTAR PHARMACEUTICALS INC  
CORTROSYN, COSYNTROPIN  
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE  
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE  
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

**AMRING PHARMS**

- \* AMRING PHARMACEUTICALS INC  
BROMFENAC SODIUM, BROMFENAC SODIUM  
LATANOPROST, LATANOPROST  
NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN  
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE

**ANACOR PHARMS INC**

- \* ANACOR PHARMACEUTICALS INC  
EUCRISA, CRISABOROLE  
KERYDIN, TAVABOROLE

**ANBEX**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ANBEX INC  
IOSAT, POTASSIUM IODIDE (OTC)

**ANBISON LAB CO LTD**

- \* ANBISON LABORATORY CO LTD  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM

**ANCHEM PHARMS**

- \* ANCHEM PHARMACEUTICALS INC  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
DUTASTERIDE AND TAMSULOSIN HYDROCHLORIDE, DUTASTERIDE  
FENOFIBRIC ACID, CHOLINE FENOFIBRATE  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE  
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE  
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
TRETINOIN, TRETINOIN  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- \* ANCHEM PHARMACEUTICALS TAIWAN INC  
DIVALPROEX SODIUM, DIVALPROEX SODIUM
- \* ANCHEM PHARMACEUTICALS, INC  
ALPRAZOLAM, ALPRAZOLAM  
CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

**ANDA REPOSITORY**

- \* ANDA REPOSITORY LLC  
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
PRIMIDONE, PRIMIDONE

**ANDRX LABS LLC**

- \* ANDRX LABS LLC  
FORTAMET, METFORMIN HYDROCHLORIDE

**ANI PHARMS**

- \* ANI PHARMACEUTICALS INC  
CORTELEMMA, HYDROCORTISONE  
LACTULOSE, LACTULOSE  
LUVOX, FLUVOXAMINE MALEATE  
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE

**ANI PHARMS INC**

- \* ANI PHARMACEUTICALS INC  
ALPRAZOLAM, ALPRAZOLAM  
CEFUROXIME AXETIL, CEFUROXIME AXETIL  
CHLORPROPAMIDE, CHLORPROPAMIDE  
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE  
DESIPIRAMINE HYDROCHLORIDE, DESIPIRAMINE HYDROCHLORIDE  
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE  
ETODOLAC, ETODOLAC  
FLECAINIDE ACETATE, FLECAINIDE ACETATE  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
GLIPIZIDE, GLIPIZIDE  
GLUCAMIDE, CHLORPROPAMIDE  
GUANABENZ ACETATE, GUANABENZ ACETATE  
INDAPAMIDE, INDAPAMIDE  
INDERAL LA, PROPRANOLOL HYDROCHLORIDE  
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE  
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE  
LITHOBID, LITHIUM CARBONATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ANI PHARMACEUTICALS INC
  - LORAZEPAM, LORAZEPAM
  - METHAZOLAMIDE, METHAZOLAMIDE
  - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
  - NILUTAMIDE, NILUTAMIDE
  - NIZATIDINE, NIZATIDINE
  - OXCARBAZEPINE, OXCARBAZEPINE
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
  - POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
  - PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
  - PROPRANOLOL HYDROCHLORIDE AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE
  - RISPERIDONE, RISPERIDONE
  - TESTOSTERONE, TESTOSTERONE
  - TRIAMTERENE AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE
  - VALPROIC ACID, VALPROIC ACID
  - VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

**ANTARES PHARMA INC**

- \* ANTARES PHARMA INC
  - OTREXUP, METHOTREXATE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

**ANTIBIOTICE**

- \* ANTIBIOTICE SA
  - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
  - AMPICILLIN SODIUM, AMPICILLIN SODIUM
  - NAFCILLIN SODIUM, NAFCILLIN SODIUM

**ANTRIM PHARMS LLC**

- \* ANTRIM PHARMACEUTICALS LLC
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE

**APEX PHARMS INC**

- \* APEX PHARMACEUTICALS INC
  - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE

**APGDI**

- \* ASTELLAS PHARMA GLOBAL DEVELOPMENT INC
  - MYRBETRIQ, MIRABEGRON

**APICORE US**

- \* APICORE US LLC
  - TETRABENAZINE, TETRABENAZINE

**APII**

- \* ALLERGAN PHARMACEUTICALS INTERNATIONAL LTD
  - ACTONEL, RISEDRONATE SODIUM
  - ASACOL HD, MESALAMINE
  - ATELVIA, RISEDRONATE SODIUM
  - DELZICOL, MESALAMINE
  - ENABLEX, DARIFENACIN HYDROBROMIDE
  - ESTROSTEP FE, ETHINYL ESTRADIOL
  - FEMCON FE, ETHINYL ESTRADIOL
  - FEMHRT, ETHINYL ESTRADIOL
  - FEMRING, ESTRADIOL ACETATE
  - LO LOESTRIN FE, ETHINYL ESTRADIOL
  - LO MINASTRIN FE, ETHINYL ESTRADIOL
  - LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
  - LOESTRIN 21 1/20, ETHINYL ESTRADIOL
  - LOESTRIN 24 FE, ETHINYL ESTRADIOL
  - LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
  - LOESTRIN FE 1/20, ETHINYL ESTRADIOL
  - MINASTRIN 24 FE, ETHINYL ESTRADIOL
  - NOR-QD, NORETHINDRONE
  - NORCO, ACETAMINOPHEN
  - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
  - SARAFEM, FLUOXETINE HYDROCHLORIDE
  - TAYTULLA, ETHINYL ESTRADIOL

**APOLLO PHARMS INC**

- \* APOLLO PHARMACEUTICALS INC
  - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

**APOPHARMA INC**

- \* APOPHARMA INC  
FERRIPROX, DEFERIPRONE

**APOTEX**

- \* APOTEX CORP  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
- \* APOTEX INC  
ALENDRONATE SODIUM, ALENDRONATE SODIUM  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
CARBIDOPA AND LEVODOPA, CARBIDOPA  
CEFUROXIME AXETIL, CEFUROXIME AXETIL  
CIMETIDINE, CIMETIDINE  
CIMETIDINE, CIMETIDINE (OTC)  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
CYCLOSPORINE, CYCLOSPORINE  
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE  
ENALAPRIL MALEATE, ENALAPRIL MALEATE  
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE  
EPLERENONE, EPLERENONE  
ETODOLAC, ETODOLAC  
FAMCICLOVIR, FAMCICLOVIR  
FAMOTIDINE, FAMOTIDINE  
FLUCONAZOLE, FLUCONAZOLE  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
GEMFIBROZIL, GEMFIBROZIL  
GLIPIZIDE, GLIPIZIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LAMIVUDINE, LAMIVUDINE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL  
OMEPRAZOLE, OMEPRAZOLE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE  
PENTOXIFYLLINE, PENTOXIFYLLINE  
PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE  
RAMIPRIL, RAMIPRIL  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE  
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE  
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE  
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE  
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE  
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE  
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE  
TROSPiUM CHLORIDE, TROSPiUM CHLORIDE

**APOTEX CORP**

- \* APOTEX CORP  
AZITHROMYCIN, AZITHROMYCIN  
CABERGOLINE, CABERGOLINE  
CLARITHROMYCIN, CLARITHROMYCIN  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL  
NARatriptan, NARatriptan HYDROCHLORIDE  
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
RILUZOLE, RILUZOLE  
SILDENAFIL CITRATE, SILDENAFIL CITRATE  
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE  
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

**APOTEX INC**

\* APOTEX INC  
ABACAVIR SULFATE, ABACAVIR SULFATE  
ACYCLOVIR, ACYCLOVIR  
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE  
ALKERAN, MELPHALAN  
ALPRAZOLAM, ALPRAZOLAM  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE  
ANASTROZOLE, ANASTROZOLE  
ARIPIPRAZOLE, ARIPIPRAZOLE  
ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE  
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM  
ATOVAQUONE, ATOVAQUONE  
AZELASTINE HYDROCHLORIDE AND FLUTICASONE PROPIONATE, AZELASTINE HYDROCHLORIDE  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE  
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
BENZONATATE, BENZONATATE  
BICALUTAMIDE, BICALUTAMIDE  
BIMATOPROST, BIMATOPROST  
BIVALIRUDIN, BIVALIRUDIN  
BROMFENAC SODIUM, BROMFENAC SODIUM  
BUDESONIDE, BUDESONIDE  
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE  
CALCITONIN-SALMON, CALCITONIN SALMON  
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL  
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL  
CARBAMAZEPINE, CARBAMAZEPINE  
CARBIDOPA AND LEVODOPA, CARBIDOPA  
CEFPROZIL, CEFPROZIL  
CELECOXIB, CELECOXIB  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE  
CICLOPIROX, CICLOPIROX  
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE  
DASATINIB, DASATINIB  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DROSPIRENONE AND ETHINYLMESTRADIOL, DROSPIRENONE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
DUTASTERIDE, DUTASTERIDE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
EZETIMIBE, EZETIMIBE  
FENOFLIBRATE (MICRONIZED), FENOFLIBRATE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)  
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE  
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE  
IBANDRONATE SODIUM, IBANDRONATE SODIUM  
IMATINIB MESYLATE, IMATINIB MESYLATE  
IMIQUIMOD, IMIQUIMOD  
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
IRBESARTAN, IRBESARTAN  
LAMIVUDINE, LAMIVUDINE  
LAMOTRIGINE, LAMOTRIGINE  
LETROZOLE, LETROZOLE  
LEVETIRACETAM, LEVETIRACETAM  
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE  
LEVOFLOXACIN, LEVOFLOXACIN  
LOVASTATIN, LOVASTATIN  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE  
MODAFINIL, MODAFINIL  
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE  
MOMETASONE FUROATE, MOMETASONE FUROATE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT****\*\* A \*\***

- \* APOTEX INC
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
  - NABUMETONE, NABUMETONE
  - NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
  - NEVIRAPINE, NEVIRAPINE
  - OFLOXACIN, OFLOXACIN
  - OLANZAPINE, OLANZAPINE
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
  - OXCARBAZEPINE, OXCARBAZEPINE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RAMIPRIL, RAMIPRIL
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
  - RASAGILINE MESYLATE, RASAGILINE MESYLATE
  - RISEDRONATE SODIUM, RISEDRONATE SODIUM
  - RISPERIDONE, RISPERIDONE
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TIMOLOL MALEATE, TIMOLOL MALEATE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRANEXAMIC ACID, TRANEXAMIC ACID
  - TRAVOPROST, TRAVOPROST
  - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
  - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID
  - ZOLMITRIPTAN, ZOLMITRIPTAN
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

- \* APOTEX INC ETOBICOKE SITE
  - ACYCLOVIR, ACYCLOVIR
  - ALLOPURINOL, ALLOPURINOL
  - BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CARBAMAZEPINE, CARBAMAZEPINE
  - CARVEDILOL, CARVEDILOL
  - CILOSTAZOL, CILOSTAZOL
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
  - DILTZAC, DILTIAZEM HYDROCHLORIDE
  - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
  - ETODOLAC, ETODOLAC
  - FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
  - GABAPENTIN, GABAPENTIN
  - LEFLUNOMIDE, LEFLUNOMIDE
  - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LISINOPRIL, LISINOPRIL
  - LORATADINE, LORATADINE (OTC)
  - MELOXICAM, MELOXICAM
  - MIRTAZAPINE, MIRTAZAPINE
  - OXaprozin, OXaprozin
  - SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
  - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
  - TOPIRAMATE, TOPIRAMATE
  - TORSEMIDE, TORSEMIDE
  - ZONISAMIDE, ZONISAMIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* APOTEX INC RICHMOND HILL
  - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
  - BUDESONIDE, BUDESONIDE (OTC)
  - DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
  - FLUNISOLIDE, FLUNISOLIDE
  - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - RISPERIDONE, RISPERIDONE
- \* APOTEX INC.
  - DILTZAC, DILTIAZEM HYDROCHLORIDE
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

**APOTEX TECHNOLOGIES**

- \* APOTEX TECHNOLOGIES INC
  - PAXIL CR, PAROXETINE HYDROCHLORIDE
  - PAXIL, PAROXETINE HYDROCHLORIDE

**APOTHECON**

- \* APOTHECON INC DIV BRISTOL MYERS SQUIBB
  - KENALOG-10, TRIAMCINOLONE ACETONIDE
  - KENALOG-40, TRIAMCINOLONE ACETONIDE

**APP PHARMS**

- \* APP PHARMACEUTICALS LLC
  - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE

**APPCO PHARMA LLC**

- \* APPCO PHARMA LLC
  - BUDESONIDE, BUDESONIDE
  - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - LAMIVUDINE, LAMIVUDINE
  - METRONIDAZOLE, METRONIDAZOLE
  - MODAFINIL, MODAFINIL
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - VORICONAZOLE, VORICONAZOLE
  - ZOLMITRIPTAN, ZOLMITRIPTAN

**APRECIA PHARMS**

- \* APRECIA PHARMACEUTICALS LLC
  - SPRITAM, LEVETIRACETAM

**AQUA PHARMS**

- \* AQUA PHARMACEUTICALS
  - ACTICLATE CAP, DOXYCYCLINE HYCLATE
  - CORDRAN SP, FLURANDRENOLIDE
  - CORDRAN, FLURANDRENOLIDE
  - MONODOX, DOXYCYCLINE
  - VERDESO, DESONIDE
- \* AQUA PHARMACEUTICALS LLC
  - FLUOROPLEX, FLUOROURACIL
  - XOLEGEL, KETOCONAZOLE

**AQUA PHARMS LLC**

- \* AQUA PHARMACEUTICALS LLC
  - ACTICLATE, DOXYCYCLINE HYCLATE
  - ALTABAX, RETAPAMULIN
  - VELTIN, CLINDAMYCIN PHOSPHATE

**ARALEZ PHARMS**

- \* ARALEZ PHARMACEUTICALS TRADING DAC
  - TOPROL-XL, METOPROLOL SUCCINATE
  - YOSPRALA, ASPIRIN
  - ZONTIVITY, VORAPAXAR SULFATE

**ARALEZ PHARMS INC**

- \* ARALEZ PHARMACEUTICALS INC
  - FIBRICOR, FENOFIBRIC ACID

**ARBOR PHARMS LLC**

- \* ARBOR PHARMACEUTICALS LLC
  - BIDIL, HYDRALAZINE HYDROCHLORIDE
  - CETYLEV, ACETYLCYSTEINE
  - E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ARBOR PHARMACEUTICALS LLC
  - E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
  - EDARBI, AZILSARTAN KAMEDOXOMIL
  - EDARBYCLOR, AZILSARTAN KAMEDOXOMIL
  - ERY-TAB, ERYTHROMYCIN
  - ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
  - ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
  - ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
  - ERYTHROMYCIN, ERYTHROMYCIN
  - EVEKEO, AMPHETAMINE SULFATE
  - GLIADEL, CARMUSTINE
  - HORIZANT, GABAPENTIN ENACARBIL
  - NYMALIZE, NIMODIPINE
  - PCE, ERYTHROMYCIN
  - SKLICE, IVERMECTIN
  - SOTYLIZE, SOTALOL HYDROCHLORIDE
  - TRIPTODUR KIT, TRIPTORELIN PAMOATE

**ARCO PHARMS LLC**

- \* ARCO PHARMACEUTICALS LLC
  - THYROSHIELD, POTASSIUM IODIDE (OTC)

**AREVA PHARMS**

- \* AREVA PHARMACEUTICALS INC
  - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

**ARIAD**

- \* ARIAD PHARMACEUTICALS INC
  - ALUNBRIG, BRIGATINIB
  - ICLUSIG, PONATINIB HYDROCHLORIDE

**ARISE PHARMS**

- \* ARISE PHARMACEUTICALS LLC
  - IBUPROFEN, IBUPROFEN (OTC)

**ASCEND THERAPS US**

- \* ASCEND THERAPEUTICS US LLC
  - ESTROGEL, ESTRADIOL

**ASCENT PHARMS INC**

- \* ASCENT PHARMACEUTICALS INC
  - DUTASTERIDE, DUTASTERIDE
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - IBUPROFEN, IBUPROFEN (OTC)
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

**ASPEN GLOBAL**

- \* ASPEN GLOBAL INC
  - MYLERAN, BUSULFAN

**ASPEN GLOBAL INC**

- \* ASPEN GLOBAL INC
  - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
  - CYCLESSA, DESOGESTREL
  - HYDROXYPROGESTERONE CAPROATE, HYDROXYPROGESTERONE CAPROATE
  - LEUKERAN, CHLORAMBUCIL
  - MYLERAN, BUSULFAN
  - THIOGUANINE, THIOGUANINE

**ASTELLAS**

- \* ASTELLAS PHARMA US INC
  - ADENOCARD, ADENOSINE
  - AMBI SOME, AMPHOTERICIN B
  - ASTAGRAF XL, TACROLIMUS
  - CRESEMBA, ISAVUCONAZONIUM SULFATE
  - LEXISCAN, REGADENOSON
  - MYCAMINE, MICAFUNGIN SODIUM
  - PROGRAF, TACROLIMUS
  - VESICARE, SOLIFENACIN SUCCINATE
  - XTANDI, ENZALUTAMIDE

**ASTRAZENECA**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* ASTRAZENECA LP  
PULMICORT FLEXHALER, BUDESONIDE  
SYMBICORT, BUDESONIDE
- \* ASTRAZENECA PHARMACEUTICALS LP  
ATACAND HCT, CANDESARTAN CILEXETIL  
ATACAND, CANDESARTAN CILEXETIL  
FASLODEX, FULVESTRANT  
ZOMIG, ZOLMITRIPTAN  
ZOMIG-ZMT, ZOLMITRIPTAN
- \* ASTRAZENECA UK LTD  
CALQUENCE, ACALABRUTINIB  
SERQUEL XR, QUETIAPINE FUMARATE

**ASTRAZENECA AB**

- \* ASTRAZENECA AB  
BYDUREON BCISE, EXENATIDE  
BYDUREON PEN, EXENATIDE SYNTHETIC  
BYDUREON, EXENATIDE SYNTHETIC  
BYETTA, EXENATIDE SYNTHETIC  
FARXIGA, DAPAGLIFLOZIN PROPANEDIOL  
KOMBIGLYZE XR, METFORMIN HYDROCHLORIDE  
ONGLYZA, SAXagliptin HYDROCHLORIDE  
QTERN, DAPAGLIFLOZIN PROPANEDIOL  
SYMLIN, PRAMLINTIDE ACETATE  
XIGDUO XR, DAPAGLIFLOZIN PROPANEDIOL

**ASTRAZENECA LP**

- \* ASTRAZENECA LP  
NEXIUM 24HR, ESOMEPRAZOLE MAGNESIUM (OTC)

**ASTRAZENECA PHARMS**

- \* ASTRAZENECA PHARMACEUTICALS LP  
ALVESCO, CICLESONIDE  
ARIMIDEX, ANASTROZOLE  
BEVESPI AEROSPHERE, FORMOTEROL FUMARATE  
BRILINTA, TICAGRELOR  
CASODEX, BICALUTAMIDE  
DALIRESP, ROFLUMILAST  
EPANOVA, OMEGA-3-CARBOXYLIC ACIDS  
IRESSA, GEFITINIB  
LYNPARZA, OLAPARIB  
MOVANTIK, NALOXEGOL OXALATE  
NEXIUM IV, ESOMEPRAZOLE SODIUM  
NEXIUM, ESOMEPRAZOLE MAGNESIUM  
OMNARIS, CICLESONIDE  
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)  
PULMICORT RESPULES, BUDESONIDE  
RHINOCORT ALLERGY, BUDESONIDE (OTC)  
SERQUEL, QUETIAPINE FUMARATE  
TAGRISSO, OSIMERTINIB MESYLATE  
TUDORZA PRESSAIR, ACLIDINIMUM BROMIDE  
ZETONNA, CICLESONIDE

**ATHENEX INC**

- \* ATHENEX INC  
BUMETANIDE, BUMETANIDE  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DIPYRIDAMOLE, DIPYRIDAMOLE  
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE  
ENALAPRILAT, ENALAPRILAT  
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE  
FAMOTIDINE, FAMOTIDINE  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE  
TERBUTALINE SULFATE, TERBUTALINE SULFATE  
VALPROATE SODIUM, VALPROATE SODIUM

**ATLAS PHARMS LLC**

- \* ATLAS PHARMACEUTICALS LLC  
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT****\*\* A \*\***

- \* ATLAS PHARMACEUTICALS LLC  
METHOCARBAMOL, METHOCARBAMOL

**ATNAHS PHARMA US**

- \* ATNAHS PHARMA US LTD  
ANAPROX DS, NAPROXEN SODIUM  
ANAPROX, NAPROXEN SODIUM  
EC-NAPROSYN, NAPROXEN  
NAPROSYN, NAPROXEN

**ATON**

- \* ATON PHARMA INC  
CUPRIMINE, PENICILLAMINE  
EDECRIN, ETHACRYNATE SODIUM  
EDECRIN, ETHACRYNIC ACID  
LACRISERT, HYDROXYPROPYL CELLULOSE  
LODOSYN, CARBIDOPA  
SPRINE, TRIENTINE HYDROCHLORIDE  
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE

**ATON PHARMA VPNA**

- \* ATON PHARMA DIV VALEANT PHARMACEUTICALS NORTH AMERICA LLC  
DEMSEER, METYROSINE

**AUROBINDO**

- \* AUROBINDO PHARMA LTD  
AMOXICILLIN, AMOXICILLIN  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
CLARITHROMYCIN, CLARITHROMYCIN  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LISINOPRIL, LISINOPRIL  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
MIRTAZAPINE, MIRTAZAPINE  
NEVIRAPINE, NEVIRAPINE  
ZIDOVUDINE, ZIDOVUDINE

**AUROBINDO PHARMA**

- \* AUROBINDO PHARMA  
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
- \* AUROBINDO PHARMA LTD  
ALENDRONATE SODIUM, ALENDRONATE SODIUM  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
AMPICILLIN SODIUM, AMPICILLIN SODIUM  
ATENOLOL, ATENOLOL  
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE  
CARISOPRODOL, CARISOPRODOL  
CARVEDILOL, CARVEDILOL  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CEFDINIR, CEFDINIR  
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL  
CEFPROZIL, CEFPROZIL  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
DIDANOSINE, DIDANOSINE  
FINASTERIDE, FINASTERIDE  
FLUCONAZOLE, FLUCONAZOLE  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM  
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE  
GLYBURIDE, GLYBURIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AUROBINDO PHARMA LTD
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - MELOXICAM, MELOXICAM
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON, ONDANSETRON
  - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
  - PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
  - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
  - QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - RIBAVARIN, RIBAVIRIN
  - RIBAVIRIN, RIBAVIRIN
  - RISPERIDONE, RISPERIDONE
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SIMVASTATIN, SIMVASTATIN
  - STAVUDINE, STAVUDINE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - TOPIRAMATE, TOPIRAMATE
  - TORSEMIDE, TORSEMIDE
  - TRANDOLAPRIL, TRANDOLAPRIL
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - ZALEPLON, ZALEPLON
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**AUROBINDO PHARMA LTD**

- \* AUROBINDO PHARMA LIMITED
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - LEVOFLOXACIN, LEVOFLOXACIN
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
- \* AUROBINDO PHARMA LTD
  - ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
  - ABACAVIR SULFATE, ABACAVIR SULFATE
  - ACETAMINOPHEN, ACETAMINOPHEN (OTC)
  - ACETYLCYSTEINE, ACETYLCYSTEINE
  - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
  - ADENOSINE, ADENOSINE
  - AFIRMELLE, ETHINYL ESTRADIOL
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
  - ALPRAZOLAM, ALPRAZOLAM
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
  - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
  - AMOXICILLIN, AMOXICILLIN
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - ATHENTIA NEXT, LEVONORGESTREL (OTC)
  - ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
  - ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
  - ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
  - AUROVELA 1.5/30, ETHINYL ESTRADIOL
  - AUROVELA 1/20, ETHINYL ESTRADIOL
  - AUROVELA 24 FE, ETHINYL ESTRADIOL
  - AUROVELA FE 1.5/30, ETHINYL ESTRADIOL
  - AUROVELA FE 1/20, ETHINYL ESTRADIOL
  - AYUNA, ETHINYL ESTRADIOL
  - AZITHROMYCIN, AZITHROMYCIN
  - BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT****\*\* A \*\***

- \* AUROBINDO PHARMA LTD
  - BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
  - CAFFEINE CITRATE, CAFFEINE CITRATE
  - CEFIXIME, CEFIXIME
  - CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
  - CEFPROZIL, CEFPROZIL
  - CEFUROXIME AXETIL, CEFUROXIME AXETIL
  - CELECOXIB, CELECOXIB
  - CEPHALEXIN, CEPHALEXIN
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - CLOZAPINE, CLOZAPINE
  - CYONANZ, ETHINYL ESTRADIOL
  - DALFAMPRIDINE, DALFAMPRIDINE
  - DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
  - DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
  - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - EFAVIRENZ, EFAVIRENZ
  - ENTACAPONE, ENTACAPONE
  - ENTECAVIR, ENTECAVIR
  - EPTIFIBATIDE, EPTIFIBATIDE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
  - ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
  - ESZOPICLONE, ESZOPICLONE
  - ETOMIDATE, ETOMIDATE
  - EZETIMIBE, EZETIMIBE
  - FAMCICLOVIR, FAMCICLOVIR
  - FAMOTIDINE, FAMOTIDINE
  - FAMOTIDINE, FAMOTIDINE (OTC)
  - FELODIPINE, FELODIPINE
  - FENOFIBRATE, FENOFIBRATE
  - FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
  - FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
  - FINASTERIDE, FINASTERIDE
  - FLECAINIDE ACETATE, FLECAINIDE ACETATE
  - FLUCONAZOLE, FLUCONAZOLE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
  - FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
  - GABAPENTIN, GABAPENTIN
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - GEMFIBROZIL, GEMFIBROZIL
  - GLIMEPIRIDE, GLIMEPIRIDE
  - GLIPIZIDE, GLIPIZIDE
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - GUAIIFENESIN AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
  - IBUPROFEN, IBUPROFEN (OTC)
  - INCASSIA, NORETHINDRONE
  - INDOMETHACIN, INDOMETHACIN
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
  - IRBESARTAN AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE
  - IRBESARTAN, IRBESARTAN
  - ISOSULFAN BLUE, ISOSULFAN BLUE
  - KALLIGA, DESOGESTREL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AUROBINDO PHARMA LTD
  - LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
  - LAMIVUDINE, LAMIVUDINE
  - LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
  - LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
  - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
  - LINEZOLID, LINEZOLID
  - LO SIMPESSE, ETHINYL ESTRADIOL
  - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - MEROPENEM, MEROPENEM
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
  - METHOCARBAMOL, METHOCARBAMOL
  - METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
  - METRONIDAZOLE, METRONIDAZOLE
  - MILI, ETHINYL ESTRADIOL
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MIRTAZAPINE, MIRTAZAPINE
  - MODAFINIL, MODAFINIL
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - NADOLOL, NADOLOL
  - NAFCILLIN SODIUM, NAFCILLIN SODIUM
  - NAPROXEN SODIUM, NAPROXEN SODIUM
  - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
  - NAPROXEN, NAPROXEN
  - NEVIRAPINE, NEVIRAPINE
  - NEXESTA FE, ETHINYL ESTRADIOL
  - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
  - NYLIA 1/35, ETHINYL ESTRADIOL
  - NYLIA 7/7/7, ETHINYL ESTRADIOL
  - OLANZAPINE, OLANZAPINE
  - OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - OMEPRAZOLE, OMEPRAZOLE
  - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXACILLIN SODIUM, OXACILLIN SODIUM
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PARICALCITOL, PARICALCITOL
  - PHENYTOIN SODIUM, PHENYTOIN SODIUM
  - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
  - PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM
  - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - PRASUGREL, PRASUGREL HYDROCHLORIDE
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
  - PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
  - RAMIPRIL, RAMIPRIL
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
  - REPAGLINIDE, REPAGLINIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AUROBINDO PHARMA LTD
  - RISEDRONATE SODIUM, RISEDRONATE SODIUM
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
  - ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - SEVELAMER CARBONATE, SEVELAMER CARBONATE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - SIMPESSE, ETHINYLMESTRADIOL
  - SPIRONOLACTONE, SPIRONOLACTONE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TELMISARTAN, TELMISARTAN
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - TRANEXAMIC ACID, TRANEXAMIC ACID
  - TRI-LO-MILI, ETHINYLMESTRADIOL
  - TRI-MILI, ETHINYLMESTRADIOL
  - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
  - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - VALSARTAN, VALSARTAN
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VORICONAZOLE, VORICONAZOLE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID
  - ZOLMITRIPTAN, ZOLMITRIPTAN
- \* AUROBINDO PHARMA LTD INC
  - ZIDOVUDINE, ZIDOVUDINE

**AUROLIFE PHARMA LLC**

- \* AUROLIFE PHARMA LLC
  - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
  - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
  - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
  - DUTASTERIDE, DUTASTERIDE
  - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - LORAZEPAM, LORAZEPAM
  - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
  - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

**AUSTARPHARMA LLC**

- \* AUSTARPHARMA LLC
  - ALENDRONATE SODIUM, ALENDRONATE SODIUM
  - METHOCARBAMOL, METHOCARBAMOL
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE

**AUXILIUM PHARMS INC**

- \* AUXILIUM PHARMACEUTICALS INC
  - TESTOPEL, TESTOSTERONE
  - THEO-24, THEOPHYLLINE

**AUXILIUM PHARMS LLC**

- \* AUXILIUM PHARMACEUTICALS LLC
  - DILATRATE-SR, ISOSORBIDE DINITRATE
  - EDEX, ALPROSTADIL
  - ROBAXIN, METHOCARBAMOL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

- \* AUXILIUM PHARMACEUTICALS LLC  
ROBAXIN-750, METHOCARBAMOL  
SEMPREX-D, ACRIVASTINE  
STRIANT, TESTOSTERONE  
TESTIM, TESTOSTERONE  
THEO-24, THEOPHYLLINE

**AVACOR PRODS**

- \* AVACOR PRODUCTS LLC  
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)

**AVADEL LEGACY**

- \* AVADEL LEGACY PHARMACEUTICALS LLC  
VAZCULEP, PHENYLEPHRINE HYDROCHLORIDE

**AVADEL PHARMS**

- \* AVADEL PHARMACEUTICALS USA INC  
ACIPHEX SPRINKLE, RABEPRAZOLE SODIUM

**AVADEL SPECLT**

- \* AVADEL SPECIALTY PHARMACEUTICALS LLC  
NOCTIVA, DESMOPRESSIN ACETATE

**AVANIR PHARMS**

- \* AVANIR PHARMACEUTICALS  
ONZETRA XSAIL, SUMATRIPTAN SUCCINATE
- \* AVANIR PHARMACEUTICALS INC  
NUDEXTA, DEXTROMETHORPHAN HYDROBROMIDE

**AVANTHI INC**

- \* AVANTHI INC  
CHLORPHENIRAMINE MALEATE, CHLORPHENIRAMINE MALEATE (OTC)  
DEXBROMPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE SULFATE, DEXBROMPHENIRAMINE MALEATE  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
DIETHYLPROPION HYDROCHLORIDE, DIETHYLPROPION HYDROCHLORIDE  
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE  
INDOMETHACIN, INDOMETHACIN  
LOMAIRA, PHENTERMINE HYDROCHLORIDE  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE

**AVEDRO INC**

- \* AVEDRO INC  
PHOTREXA VISCOSUS IN DEXTRAN 20%, RIBOFLAVIN 5'-PHOSPHATE SODIUM  
PHOTREXA, RIBOFLAVIN 5'-PHOSPHATE SODIUM

**AVEMA PHARMA**

- \* AVEMA PHARMA SOLUTIONS  
IBUPROFEN, IBUPROFEN (OTC)

**AVENT**

- \* AVENT INC  
PYTEST KIT, UREA, C-14  
PYTEST, UREA, C-14

**AVEVA**

- \* AVEVA DRUG DELIVERY SYSTEMS INC  
CLONIDINE, CLONIDINE  
FENTANYL-100, FENTANYL  
FENTANYL-12, FENTANYL  
FENTANYL-25, FENTANYL  
FENTANYL-37, FENTANYL  
FENTANYL-50, FENTANYL  
FENTANYL-62, FENTANYL  
FENTANYL-75, FENTANYL  
FENTANYL-87, FENTANYL  
NICOTINE, NICOTINE (OTC)

**AVID RADIOPHARMS INC**

- \* AVID RADIOPHARMACEUTICALS INC  
AMYVID, FLORBETAPIR F-18

**AVONDALE PHARMS**

- \* AVONDALE PHARMACEUTICALS LLC  
NIACOR, NIACIN

**AYTU BIOSCIENCE INC**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* A \*\*

\* AYTU BIOSCIENCE INC  
NATESTO, TESTOSTERONE

\*\* B \*\*

**B BRAUN**

\* B BRAUN MEDICAL INC  
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL  
AMINO ACIDS, AMINO ACIDS  
BALANCED SALT, CALCIUM CHLORIDE  
CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM  
CEFEPIME AND DEXTROSE IN DUPLEX CONTAINER, CEFEPIME HYDROCHLORIDE  
CEFOTETAN AND DEXTROSE IN DUPLEX CONTAINER, CEFOTETAN DISODIUM  
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER, CEFOXITIN SODIUM  
CEFTAZIDIME IN DEXTROSE CONTAINER, CEFTAZIDIME  
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM  
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM  
DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM  
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE  
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE  
FREAMEINE HBC 6.9%, AMINO ACIDS  
FREAMEINE III 10%, AMINO ACIDS  
FREAMEINE III 3% W/ ELECTROLYTES, AMINO ACIDS  
FREAMEINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS  
FREAMEINE III 8.5%, AMINO ACIDS  
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE  
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN  
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPATAMINE 8%, AMINO ACIDS  
ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE  
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL  
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE  
NEPHRAMINE 5.4%, AMINO ACIDS  
NUTRILIPID 10%, SOYBEAN OIL  
NUTRILIPID 20%, SOYBEAN OIL  
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

\* B BRAUN MEDICAL INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* B BRAUN MEDICAL INC
  - SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
  - STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
  - STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION
  - THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
  - THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
  - THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
  - THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
  - TROPHAMINE 10%, AMINO ACIDS
  - TROPHAMINE, AMINO ACIDS

**B BRAUN MEDICAL INC**

- \* B BRAUN MEDICAL INC
  - CLOROTEKAL, CHLOROPROCAINE HYDROCHLORIDE
  - MEROPENEM AND SODIUM CHLORIDE IN DUPLEX CONTAINER, MEROPENEM

**BAJAJ MEDICAL LLC**

- \* BAJAJ MECICAL LLC
  - DYNA-HEX, CHLORHEXIDINE GLUCONATE (OTC)

**BARR**

- \* BARR LABORATORIES INC
  - AMILORIDE HYDROCHLORIDE AND HYDROCHLORTIAZIDE, AMILORIDE HYDROCHLORIDE
  - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
  - ARANELLE, ETHINYL ESTRADIOL
  - ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
  - BALZIVA-28, ETHINYL ESTRADIOL
  - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
  - CHLORZOXAZONE, CHLORZOXAZONE
  - CLONAZEPAM, CLONAZEPAM
  - DANAZOL, DANAZOL
  - DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
  - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
  - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
  - DIAZEPAM, DIAZEPAM
  - DIDANOSINE, DIDANOSINE
  - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
  - DIPYRIDAMOLE, DIPYRIDAMOLE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
  - DUTASTERIDE, DUTASTERIDE
  - ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
  - ESTRADIOL AND NORGESTIMATE, ESTRADIOL
  - ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
  - FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
  - FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - HYDROXYUREA, HYDROXYUREA
  - HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
  - ISONIAZID, ISONIAZID
  - JUNEL 1.5/30, ETHINYL ESTRADIOL
  - JUNEL 1/20, ETHINYL ESTRADIOL
  - JUNEL FE 1.5/30, ETHINYL ESTRADIOL
  - JUNEL FE 1/20, ETHINYL ESTRADIOL
  - KARIVA, DESOGESTREL
  - KELNOR, ETHINYL ESTRADIOL
  - LEFLUNOMIDE, LEFLUNOMIDE
  - LESSINA-28, ETHINYL ESTRADIOL
  - MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
  - MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
  - MEGESTROL ACETATE, MEGESTROL ACETATE
  - MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHOTREXATE SODIUM, METHOTREXATE SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BARR LABORATORIES INC
  - NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE NIACIN, NIACIN
  - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
  - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
  - NORTREL 0.5/35-28, ETHINYL ESTRADIOL
  - NORTREL 1/35-21, ETHINYL ESTRADIOL
  - NORTREL 1/35-28, ETHINYL ESTRADIOL
  - NORTREL 7/7/7, ETHINYL ESTRADIOL
  - ONDANSETRON, ONDANSETRON
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PORTIA-28, ETHINYL ESTRADIOL
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - SPRINTEC, ETHINYL ESTRADIOL
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - TREXALL, METHOTREXATE SODIUM
  - TRI-LEGEST 21, ETHINYL ESTRADIOL
  - TRI-LEGEST FE, ETHINYL ESTRADIOL
  - TRI-SPRINTEC, ETHINYL ESTRADIOL
  - WARFARIN SODIUM, WARFARIN SODIUM

- \* BARR PHARMACEUTICALS LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

**BARR LABS DIV TEVA**

- \* BARR LABORATORIES INC SUB TEVA PHARMACEUTICALS USA
  - ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
  - BUDESONIDE, BUDESONIDE
  - OXYBUTYNIN, OXYBUTYNIN

**BARR LABS INC**

- \* BARR LABORATORIES INC
  - ACITRETIN, ACITRETIN
  - CLOZAPINE, CLOZAPINE
  - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
  - ESTRADIOL, ESTRADIOL
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
  - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - OLANZAPINE, OLANZAPINE
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - OXYCODONE HYDROCHLORIDE AND IBUPROFEN, IBUPROFEN
  - TRETINOIN, TRETINOIN
  - TRI LO SPRINTEC, ETHINYL ESTRADIOL

**BAUSCH AND LOMB**

- \* BAUSCH AND LOMB INC
  - ALAWAY, KETOTIFEN FUMARATE (OTC)
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - ALREX, LOTEPEREDNOL ETABONATE
  - BESIVANCE, BESIFLOXACIN HYDROCHLORIDE
  - CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
  - DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
  - FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
  - ISTALOL, TIMOLOL MALEATE
  - LATANOPROST, LATANOPROST
  - LOTEMAX, LOTEPEREDNOL ETABONATE
  - MIOCHOL-E, ACETYLCHOLINE CHLORIDE
  - OFLOXACIN, OFLOXACIN
  - OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
  - PROLENSA, BROMFENAC SODIUM
  - RETISERT, FLUOCINOLONE ACETONIDE
  - SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
  - TIMOLOL MALEATE, TIMOLOL MALEATE
  - VITRASE, HYALURONIDASE
  - VYZULTA, LATANOPROSTENE BUNOD

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BAUSCH AND LOMB INC  
ZIRGAN, GANCICLOVIR  
ZYLET, LOTEPREDNOL ETABONATE
- \* BAUSCH AND LOMB PHARMACEUTICALS INC  
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC  
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE  
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)  
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE  
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE  
DEXASPORIN, DEXAMETHASONE  
ERYTHROMYCIN, ERYTHROMYCIN  
FLUNISOLIDE, FLUNISOLIDE  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE  
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE  
NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC  
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE  
NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN  
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE  
NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC  
OFLOXACIN, OFLOXACIN  
OPTIPRANOL, METIPRANOL HYDROCHLORIDE  
OTICAIR, HYDROCORTISONE  
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE  
PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE  
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM  
TIMOLOL MALEATE, TIMOLOL MALEATE  
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE  
TOBRAMYCIN, TOBRAMYCIN  
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE  
TROPICAMIDE, TROPICAMIDE

**BAUSCH AND LOMB INC**

- \* BAUSCH AND LOMB INC  
BEPREVE, BEPOTASTINE BESILATE  
LOTEMAX, LOTEPREDNOL ETABONATE  
LUMIFY, BRIMONIDINE TARTRATE (OTC)

**BAXTER HLTHCARE**

- \* BAXTER HEALTHCARE CORP  
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL  
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE  
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM  
BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM  
REVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE  
REVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE  
REVIBLOC, ESMOLOL HYDROCHLORIDE  
CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
CEFEPIME IN PLASTIC CONTAINER, CEFEPIME HYDROCHLORIDE  
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM  
CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS  
CLINIMIX E 2.75/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC  
CLINIMIX E 2.75/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC  
CLINIMIX E 2.75/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,  
CLINIMIX E 4.25/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BAXTER HEALTHCARE CORP
  - CLINIMIX E 4.25/20 SULFITE FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC
  - CLINIMIX E 4.25/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC
  - CLINIMIX E 4.25/5 SULFITE FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER,
  - CLINIMIX E 5/10 SULFITE FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER,
  - CLINIMIX E 5/15 SULFITE FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER,
  - CLINIMIX E 5/20 SULFITE FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER,
  - CLINIMIX E 5/25 SULFITE FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER,
  - CLINIMIX E 5/35 SULFITE FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER,
  - CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
  - CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
  - DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND ELECTROLYTE NO. 48 IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
  - DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER,
  - DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER,
  - DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER,
  - DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER,
  - DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER,
  - DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER,
  - DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC
  - DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 60% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
  - DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
  - DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
  - EXTRANEAL, ICODEXTRIN
  - FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
  - FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
  - FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BAXTER HEALTHCARE CORP  
FORANE, ISOFLURANE  
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE  
HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN  
HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN  
IFEX, IFOSFAMIDE  
LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE  
MESNEX, MESNA  
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE  
NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM  
NEXTERONE, AMIODARONE HYDROCHLORIDE  
NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN  
OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 10% IN WATER, MANNITOL  
OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 15% IN WATER, MANNITOL  
OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 20% IN WATER, MANNITOL  
OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL  
OSMITROL 5% IN WATER, MANNITOL  
PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM  
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM  
POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM  
POTASSIUM CHLORIDE 0.224% IN SODIUM CHLORIDE 0.9%, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 0.3% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS  
PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS  
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
SEVOFLURANE, SEVOFLURANE  
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL  
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION  
STERILE WATER, STERILE WATER FOR IRRIGATION

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BAXTER HEALTHCARE CORP  
SUPRANE, DESFLURANE  
TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE  
TIS-U-SOL, MAGNESIUM SULFATE  
TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS  
TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS  
TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS  
VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
- \* BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV  
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

**BAXTER HLTHCARE CORP**

- \* BAXTER HEALTHCARE CORP  
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE  
CEFAZOLIN IN PLASTIC CONTAINER, CEFAZOLIN SODIUM  
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN  
CIPROFLOXACIN, CIPROFLOXACIN  
CLINDAMYCIN PHOSPHATE IN 0.9% SODIUM CHLORIDE, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE  
FUROSEMIDE, FUROSEMIDE  
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN  
LEVOFLOXACIN, LEVOFLOXACIN  
METOPROLOL TARTRATE, METOPROLOL TARTRATE  
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE  
NOREpinephrine Bitartrate, NOREpinephrine Bitartrate  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL B22GK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL B22GK 4/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL B22GK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 2/0 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 4/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BK 0/0/1.2 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
- \* BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE  
PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
- \* BAXTER HEALTHCARE CORP ANESTHESIA CRITICAL CARE  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

**BAYER**

- \* BAYER HEALTHCARE LLC  
ALEVE, NAPROXEN SODIUM (OTC)  
ALEVE-D SINUS & COLD, NAPROXEN SODIUM (OTC)

**BAYER HEALTHCARE**

- \* BAYER HEALTHCARE PHARMACEUTICALS INC  
ALIQOPA, COPANLISIB DIHYDROCHLORIDE

**BAYER HEALTHCARE LLC**

- \* BAYER HEALTHCARE LLC  
CHILDREN'S CLARITIN, LORATADINE (OTC)  
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)  
CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)  
CLARITIN HIVES RELIEF, LORATADINE (OTC)  
CLARITIN REDITABS, LORATADINE (OTC)  
CLARITIN, LORATADINE (OTC)  
CLARITIN-D 24 HOUR, LORATADINE (OTC)  
CLARITIN-D, LORATADINE (OTC)  
LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)  
MIRALAX, POLYETHYLENE GLYCOL 3350 (OTC)  
MYCELEX-7 COMBINATION PACK, CLOTRIMAZOLE (OTC)  
MYCELEX-7, CLOTRIMAZOLE (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BAYER HEALTHCARE LLC  
ZEGERID OTC, OMEPRAZOLE (OTC)

**BAYER HLTHCARE**

- \* BAYER HEALTHCARE CONSUMER CARE  
ALEVE PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
- \* BAYER HEALTHCARE PHARMACEUTICALS INC  
ADEMPAS, RIOCIGUAT  
ANGELIQ, DROSPIRENONONE  
AVELOX, MOXIFLOXACIN HYDROCHLORIDE  
BEYAZ, DROSPIRENONE  
BILTRICIDE, PRAZIQUANTEL  
CIPRO, CIPROFLOXACIN  
CIPRO, CIPROFLOXACIN HYDROCHLORIDE  
CLIMARA PRO, ESTRADIOL  
CLIMARA, ESTRADIOL  
DESONATE, DESONIDE  
EOVIST, GADOXETATE DISODIUM  
FINACEA, AZELAIC ACID  
FINACEA, AZELAIC ACID  
GADAVIST, GADOBUTROL  
KYLEENA, LEVONORGESTREL  
LEVITRA, VARDENAFIL HYDROCHLORIDE  
MAGNEVIST, GADOPENTETATE DIMEGLUMINE  
MENOSTAR, ESTRADIOL  
MIRENA, LEVONORGESTREL  
NATAZIA, DIENOGEST  
NEXAVAR, SORAFENIB TOSYLATE  
PRECPOSE, ACARBOSE  
SAFYRAL, DROSPIRENONE  
SKYLA, LEVONORGESTREL  
STAXYN, VARDENAFIL HYDROCHLORIDE  
STIVARGA, REGORAFENIB  
ULTRAVIST (PHARMACY BULK), IOPROMIDE  
ULTRAVIST 240, IOPROMIDE  
ULTRAVIST 300, IOPROMIDE  
ULTRAVIST 370, IOPROMIDE  
XOFIGO, RADIUM RA-223 DICHLORIDE  
YASMIN, DROSPIRENONE  
YAZ, DROSPIRENONE

**BAYSHORE PHARMS LLC**

- \* BAYSHORE PHARMACEUTICALS LLC  
METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE  
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE

**BDSI**

- \* BIODELIVERY SCIENCES INTERNATIONAL INC  
BELBUCA, BUPRENORPHINE HYDROCHLORIDE  
BUNAVAIL, BUPRENORPHINE HYDROCHLORIDE

**TECTON DICKINSON**

- \* BECTON DICKINSON AND CO  
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)  
E-Z SCRUB 201, POVIDONE-IODINE (OTC)  
E-Z SCRUB 241, POVIDONE-IODINE (OTC)

**TECTON DICKINSON CO**

- \* BECTON DICKINSON AND CO  
CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP TRIPLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)

**BEDFORD**

- \* BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC  
CEFTRIAXONE, CEFTRIAXONE SODIUM

**BEDFORD LABS**

- \* BEDFORD LABORATORIES

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BEDFORD LABORATORIES  
LORAZEPAM PRESERVATIVE FREE, LORAZEPAM

**BELCHER PHARMS**

- \* BELCHER PHARMACEUTICALS LLC  
CEPHALEXIN, CEPHALEXIN  
DESLORATADINE, DESLORATADINE

**BELCHER PHARMS LLC**

- \* BELCHER PHARMACEUTICALS LLC  
CEFIXIME, CEFIXIME  
EPINEPHRINE, EPINEPHRINE  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE  
TACROLIMUS, TACROLIMUS

**BEXIMCO PHARMS USA**

- \* BEXIMCO PHARMACEUTICALS USA INC  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
METHOCARBAMOL, METHOCARBAMOL  
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

**BEXIMCO USA**

- \* BEXIMCO PHARMACEUTICALS USA INC  
CARVEDILOL, CARVEDILOL

**BI-COASTAL PHARMA**

- \* BI-COASTAL PHARMA INTERNATIONAL LLC  
DUVOID, BETHANECHOL CHLORIDE

**BIO NUCLEONICS**

- \* BIO NUCLEONICS INC  
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE SR-89

**BIO PHARM INC**

- \* BIO PHARM INC  
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
RISPERIDONE, RISPERIDONE

**BIO-PHARM INC**

- \* BIO-PHARM INC  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE  
HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,  
LACTULOSE, LACTULOSE  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

**BIOCON LIMITED**

- \* BIOCON LIMITED  
SIMVASTATIN, SIMVASTATIN

**BIOCON LTD**

- \* BIOCON LTD  
ROSVASTATIN CALCIUM, ROUVASTATIN CALCIUM

**BIOCRYST**

- \* BIOCRYST PHARMACEUTICALS INC  
RAPIVAB, PERAMIVIR

**BIOFRONTERA**

- \* BIOFRONTERA BIOSCIENCE GMBH  
AMELUZ, AMINOLEVULINIC ACID HYDROCHLORIDE

**BIOGEN IDEC**

- \* BIOGEN IDEC INC  
SPINRAZA, NUSINERSEN SODIUM

**BIOGEN IDEC INC**

- \* BIOGEN IDEC INC  
TECFIDERA, DIMETHYL FUMARATE

**BIOMARIN PHARM**

- \* BIOMARIN PHARMACEUTICAL INC  
KUVAN, SAPROPTERIN DIHYDROCHLORIDE

**BIOMEDICAL RES FDN**

- \* BIOMEDICAL RESEARCH FOUNDATION NORTHWEST LOUISIANA  
AMMONIA N 13, AMMONIA N-13

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BIOMEDICAL RESEARCH FOUNDATION NORTHWEST LOUISIANA  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**BIONPHARMA INC**

- \* BIONPHARMA INC  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
BENZONATATE, BENZONATATE  
BEXAROTENE, BEXAROTENE  
CALCITRIOL, CALCITRIOL  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM  
DUTASTERIDE, DUTASTERIDE  
ETHOSUXIMIDE, ETHOSUXIMIDE  
GRANisetron HYDROCHLORIDE PRESERVATIVE FREE, GRANisetron HYDROCHLORIDE  
GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE  
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)  
IBUPROFEN, IBUPROFEN (OTC)  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)  
MIDOL LIQUID GELS, IBUPROFEN (OTC)  
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)  
NIMODIPINE, NIMODIPINE  
PARICALCITOL, PARICALCITOL  
PROGESTERONE, PROGESTERONE  
TETRABENAZINE, TETRABENAZINE  
VALPROIC ACID, VALPROIC ACID  
VITAMIN D, ERGOCALCIFEROL  
ZONISAMIDE, ZONISAMIDE

**BLAIREX**

- \* BLAIREX LABORATORIES INC  
BRONCHO SALINE, SODIUM CHLORIDE (OTC)

**BLUE EARTH**

- \* BLUE EARTH DIAGNOSTICS LTD  
AXUMIN, FLUCICLOVINE F-18

**BOEHRINGER INGELHEIM**

- \* BOEHRINGER INGELHEIM  
CATAPRES, CLONIDINE HYDROCHLORIDE  
CATAPRES-TTS-1, CLONIDINE  
CATAPRES-TTS-2, CLONIDINE  
CATAPRES-TTS-3, CLONIDINE  
GILOTrif, AFATINIB DIMALEATE  
GLYXAMBI, EMPAGLIFLOZIN  
MICARDIS HCT, HYDROCHLOROTHIAZIDE  
MICARDIS, TELMISARTAN  
MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
- \* BOEHRINGER INGELHEIM PHARMACEUTICALS INC  
AGGRENOX, ASPIRIN  
APTIVUS, TIPRANAVIR  
ATROVENT HFA, IPRATROPIUM BROMIDE  
ATROVENT, IPRATROPIUM BROMIDE  
COMBIVENT RESPIMAT, ALBUTEROL SULFATE  
FLOMAX, TAMSULOSIN HYDROCHLORIDE  
JARDIANCE, EMPAGLIFLOZIN  
JENTADUETO XR, LINAGLIPTIN  
JENTADUETO, LINAGLIPTIN  
MIRAPEX ER, PRAMIPEXOLE DIHYDROCHLORIDE  
MOBIC, MELOXICAM  
OFEV, NINTEDANIB ESYLATE  
PERSANTINE, DIPYRIDAMOLE  
PRADAXA, DABIGATRAN ETEXILATE MESYLATE  
SPIRIVA RESPIMAT, TIOTROPIUM BROMIDE  
SPIRIVA, TIOTROPIUM BROMIDE  
STIOLTO RESPIMAT, OLODATEROL HYDROCHLORIDE  
STRIVERDI RESPIMAT, OLODATEROL HYDROCHLORIDE  
SYNJARDY XR, EMPAGLIFLOZIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BOEHRINGER INGELHEIM PHARMACEUTICALS INC  
SYNJARDY, EMPAGLIFLOZIN  
TRADJENTA, LINAGLIPTIN  
TWYNSTA, AMLODIPIINE BESYLATE  
VIRAMUNE XR, NEVIRAPINE  
VIRAMUNE, NEVIRAPINE

**BOSCOGEN**

- \* BOSCOGEN INC  
ACYCLOVIR, ACYCLOVIR  
REPAGLINIDE, REPAGLINIDE  
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

**BPI LABS LLC**

- \* BPI LABS LLC  
ZOLEDRONIC ACID, ZOLEDRONIC ACID

**BRACCO**

- \* BRACCO DIAGNOSTICS INC  
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82  
CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT  
CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE  
CYSTOGRAFIN, DIATRIZOATE MEGLUMINE  
E-Z-HD, BARIUM SULFATE  
E-Z-PAQUE, BARIUM SULFATE  
GASTROGRAFIN, DIATRIZOATE MEGLUMINE  
ISOVUE-200, IOPAMIDOL  
ISOVUE-250, IOPAMIDOL  
ISOVUE-300, IOPAMIDOL  
ISOVUE-370, IOPAMIDOL  
ISOVUE-M 200, IOPAMIDOL  
ISOVUE-M 300, IOPAMIDOL  
KINEVAC, SINCALIDE  
LIQUID E-Z-PAQUE, BARIUM SULFATE  
LUMASON, SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES  
MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE  
MULTIHANCE, GADOBENATE DIMEGLUMINE  
PROHANCE MULTIPACK, GADOTERIDOL  
PROHANCE, GADOTERIDOL  
READI-CAT 2 SMOOTHIES, BARIUM SULFATE  
READI-CAT 2, BARIUM SULFATE  
RENOGRAFIN-76, DIATRIZOATE MEGLUMINE  
SINOGRAFIN, DIATRIZOATE MEGLUMINE  
TAGITOL V, BARIUM SULFATE  
VARIBAR NECTAR, BARIUM SULFATE  
VARIBAR, BARIUM SULFATE

**BRAEBURN PHARMS INC**

- \* BRAEBURN PHARMACEUTICALS INC  
PROBUPHINE, BUPRENORPHINE HYDROCHLORIDE

**BRAINTREE**

- \* BRAINTREE LABORATORIES INC  
GOLYTELY, POLYETHYLENE GLYCOL 3350  
NULYTLY, POLYETHYLENE GLYCOL 3350  
NULYTLY-FLAVORED, POLYETHYLENE GLYCOL 3350

**BRAINTREE LABS**

- \* BRAINTREE LABORATORIES INC  
SUPREP BOWEL PREP KIT, MAGNESIUM SULFATE

**BRECKENRIDGE PHARM**

- \* BRECKENRIDGE PHARMACEUTICAL INC  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE  
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CILOSTAZOL, CILOSTAZOL  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
DUTASTERIDE, DUTASTERIDE  
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE  
ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL  
LEVETIRACETAM, LEVETIRACETAM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* B \*\*

- \* BRECKENRIDGE PHARMACEUTICAL INC
  - MEFENAMIC ACID, MEFENAMIC ACID
  - MEGESTROL ACETATE, MEGESTROL ACETATE
  - MELOXICAM, MELOXICAM
  - METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
  - METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - NEOMYCIN SULFATE, NEOMYCIN SULFATE
  - OMEPRAZOLE, OMEPRAZOLE
  - OXCARBAZEPINE, OXCARBAZEPINE
  - PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE , POLYETHYLENE
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**BRIGHAM WOMENS**

- \* BRIGHAM AND WOMENS HOSP
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**BRIGHAM WOMENS HOSP**

- \* BRIGHAM AND WOMENS HOSP INC
  - AMMONIA N 13, AMMONIA N-13

**BRISTOL MYERS SQUIBB**

- \* BRISTOL MYERS SQUIBB
  - AZACTAM, AZTREONAM
  - BARACLUDE, ENTECAVIR
  - LYSODREN, MITOTANE
  - PRAVACHOL, PRAVASTATIN SODIUM
- \* BRISTOL MYERS SQUIBB CO
  - AZACTAM IN PLASTIC CONTAINER, AZTREONAM
  - DROXIA, HYDROXYUREA
  - GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
  - HYDREA, HYDROXYUREA
  - REYATAZ, ATAZANAVIR SULFATE
  - SPRYCEL, DASATINIB
  - SUSTIVA, EFAVIRENZ
  - VIDEX EC, DIDANOSINE
- \* BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
  - ELIQUIS, APIXABAN
  - ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
  - GLUCOPHAGE, METFORMIN HYDROCHLORIDE
  - ZERIT, STAVUDINE
- \* BRISTOL MYERS SQUIBB PHARMA CO
  - COUMADIN, WARFARIN SODIUM

**BRISTOL-MYERS SQUIBB**

- \* BRISTOL-MYERS SQUIBB CO
  - DAKLINZA, DACLATASVIR DIHYDROCHLORIDE
  - EVOTAZ, ATAZANAVIR SULFATE
  - VIDEX, DIDANOSINE

\*\* C \*\*

**CADILA PHARMS LTD**

- \* CADILA PHARMACEUTICALS LTD
  - ACYCLOVIR, ACYCLOVIR
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - FOLIC ACID, FOLIC ACID
  - GEMFIBROZIL, GEMFIBROZIL
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - METRONIDAZOLE, METRONIDAZOLE
  - OFLOXACIN, OFLOXACIN
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - TELMISARTAN, TELMISARTAN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

- \* CADILA PHARMACEUTICALS LTD  
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**CADISTA PHARMS**

- \* CADISTA PHARMACEUTICALS INC  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM

**CALL INC**

- \* CALL INC DBA ROCHESTER PHARMACEUTICALS  
ADAPALENE, ADAPALENE

**CARDINAL HEALTH 414**

- \* CARDINAL HEALTH 414 LLC CARDINAL HEALTH NUCLEAR PHARMACY SERVICES  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
LYMPHOSEEK KIT, TECHNETIUM TC-99M TILMANOCEPT  
MDP-BRACCO, TECHNETIUM TC-99M MEDRONATE KIT  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18  
TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

**CARDINAL HEALTH 418**

- \* CARDINAL HEALTH 418 INC  
SODIUM IODIDE I 123, SODIUM IODIDE I-123

**CARIBE HOLDINGS**

- \* CARIBE HOLDINGS CAYMAN CO LTD DBA PURACAP CARIBE  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
GEMFIBROZIL, GEMFIBROZIL

**CARLSBAD**

- \* CARLSBAD TECHNOLOGY INC  
ACYCLOVIR, ACYCLOVIR  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
FAMOTIDINE, FAMOTIDINE  
GLIMEPIRIDE, GLIMEPIRIDE  
LOVASTATIN, LOVASTATIN

**CARLSBAD TECHNOLOGY**

- \* CARLSBAD TECHNOLOGY INC  
ACYCLOVIR, ACYCLOVIR

**CARMEL BIOSCIENCES**

- \* CARMEL BIOSCIENCES INC  
PREXXARTAN, VALSARTAN

**CASPER PHARMA LLC**

- \* CASPER PHARMA LLC  
CASPORYN HC, HYDROCORTISONE  
FURADANTIN, NITROFURANTOIN  
NEOSPORIN, BACITRACIN ZINC  
PERMAPEN, PENICILLIN G BENZATHINE  
ROBINUL FORTE, GLYCOPYRROLATE  
ROBINUL, GLYCOPYRROLATE  
TERRAMYCIN W/ POLYMYXIN B SULFATE, OXYTETRACYCLINE HYDROCHLORIDE  
ZYLOPRIM, ALLOPURINOL

**CATALENT**

- \* CATALENT PHARMA SOLUTIONS LLC  
VALPROIC ACID, VALPROIC ACID

**CEDIPROF INC**

- \* CEDIPROF INC  
LEVO-T, LEVOTHYROXINE SODIUM \*\*

**CELATOR PHARMS**

- \* CELATOR PHARMACEUTICALS INC  
VYXEOS, CYTARABINE

**CELERITY PHARMS**

- \* CELERITY PHARMACEUTICALS LLC  
BIVALIRUDIN IN 0.9% SODIUM CHLORIDE, BIVALIRUDIN

**CELGENE**

- \* CELGENE CORP  
ISTODAX, ROMIDEPSIN  
POMALYST, POMALIDOMIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

- \* CELGENE CORP  
REVLIMID, LENALIDOMIDE  
THALOMID, THALIDOMIDE  
VIDAZA, AZACITIDINE

**CELGENE CORP**

- \* CELGENE CORP  
IDHIFA, ENASIDENIB MESYLATE  
OTEZLA, APREMILAST

**CEPHALON**

- \* CEPHALON INC  
ACTIQ, FENTANYL CITRATE  
FENTORA, FENTANYL CITRATE  
GABITRIL, TIAGABINE HYDROCHLORIDE  
NUVIGIL, ARMODAFINIL  
PROVIGIL, MODAFINIL  
TREANDA, BENDAMUSTINE HYDROCHLORIDE  
TRISENOX, ARSENIC TRIOXIDE

**CEREXA**

- \* CEREXA INC  
TEFLARO, CEFTAROLINE FOSAMIL
- \* CEREXA INC A SUB OF FOREST LABORATORIES LLC  
AVYCAZ, AVIBACTAM SODIUM

**CFT PHARMS LLC**

- \* CFT PHARMACEUTICALS LLC  
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

**CHANGZHOU PHARM**

- \* CHANGZHOU PHARMACEUTICAL FACTORY  
ROSVUSTATIN CALCIUM, ROUVUSTATIN CALCIUM

**CHARTWELL LIFE SCI**

- \* CHARTWELL LIFE SCIENCE LLC  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
DOXYCYCLINE, DOXYCYCLINE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**CHARTWELL MOLECULES**

- \* CHARTWELL MOLECULES LLC  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
DISULFIRAM, DISULFIRAM  
GEMFIBROZIL, GEMFIBROZIL  
NABUMETONE, NABUMETONE  
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

**CHARTWELL RX**

- \* CHARTWELL RX SCIENCES LLC  
CALCIUM ACETATE, CALCIUM ACETATE  
CILOSTAZOL, CILOSTAZOL  
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE  
INDOMETHACIN, INDOMETHACIN  
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE

**CHARTWELL TETRA**

- \* CHARTWELL TETRA LLC  
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

**CHATTEM**

- \* CHATTEM INC  
SELSUN, SELENIUM SULFIDE  
UNISOM, DOXYLAMINE SUCCINATE (OTC)

**CHEMI SPA**

- \* CHEMI SPA  
DECITABINE, DECITABINE  
TEMOZOLOMIDE, TEMOZOLOMIDE

**CHEMISCH FBRK KRSSLR**

- \* CHEMISCHE FABRIK KREUSSLER & CO. GMBH  
ASCLERA, POLIDOCANOL

**CHEMO RESEARCH SL**

- \* CHEMO RESEARCH SL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

\* CHEMO RESEARCH SL  
 BENZNIDAZOLE, BENZNIDAZOLE  
 ECOZA, ECONAZOLE NITRATE  
 NUVESSA, METRONIDAZOLE  
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN

**CHEPLAPHARM**

\* CHEPLAPHARM ARZNEIMITTEL GMBH  
 XENICAL, ORLISTAT

**CHIESI USA INC**

\* CHIESI USA INC  
 BETHKIS, TOBRAMYCIN  
 CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
 CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
 CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
 CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER, NICARDIPINE HYDROCHLORIDE  
 CARDENE, NICARDIPINE HYDROCHLORIDE  
 CLEVIPREX, CLEVIDIPINE  
 CUROSURF, PORACTANT ALFA  
 KENGREAL, CANGRELOR  
 ZYFLO CR, ZILEUTON  
 ZYFLO, ZILEUTON

**CHILDRENS HOSP MI**

\* CHILDRENS HOSP MICHIGAN  
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**CHINA RESOURCES**

\* CHINA RESOURCES SAIKE PHARMACEUTICAL CO LTD  
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE

**CHIRHOCLIN**

\* CHIRHOCLIN INC  
 CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN

**CINTEX SVCS**

\* CINTEX SERVICES LLC  
 FLURANDRENOLIDE, FLURANDRENOLIDE

**CIPHER PHARMS INC**

\* CIPHER PHARMACEUTICALS INC  
 CONZIP, TRAMADOL HYDROCHLORIDE  
 LIPOFEN, FENOFLIBRATE

**CIPLA**

\* CIPLA LTD  
 NEVIRAPINE, NEVIRAPINE  
 RISPERIDONE, RISPERIDONE  
 ZIDOVUDINE, ZIDOVUDINE

**CIPLA LTD**

\* CIPLA LTD  
 ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE  
 ABACAVIR SULFATE, ABACAVIR SULFATE  
 ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE  
 ALENDRONATE SODIUM, ALENDRONATE SODIUM  
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
 ANASTROZOLE, ANASTROZOLE  
 BUDESONIDE, BUDESONIDE  
 CARBOPLATIN, CARBOPLATIN  
 CARVEDILOL, CARVEDILOL  
 CELECOXIB, CELECOXIB  
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
 CICLOPIROX, CICLOPIROX  
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
 DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE  
 DECITABINE, DECITABINE  
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
 ENTECAVIR, ENTECAVIR  
 EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE  
 FAMCICLOVIR, FAMCICLOVIR  
 FENOFLIBRATE, FENOFLIBRATE  
 FINASTERIDE, FINASTERIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

\* CIPLA LTD  
 FLUTAMIDE, FLUTAMIDE  
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE  
 GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE  
 GRISEOFULVIN, GRISEOFULVIN, MICROSIZE  
 IRBESARTAN, IRBESARTAN  
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
 LAMIVUDINE, LAMIVUDINE  
 LAMOTRIGINE, LAMOTRIGINE  
 LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE  
 LEVOFLOXACIN, LEVOFLOXACIN  
 MELOXICAM, MELOXICAM  
 MONTELUKAST SODIUM, MONTELUKAST SODIUM  
 NEVIRAPINE, NEVIRAPINE  
 OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE  
 OXALIPLATIN, OXALIPLATIN  
 PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE  
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM  
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
 STAVUDINE, STAVUDINE  
 TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE  
 TOPIRAMATE, TOPIRAMATE  
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE  
 VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE  
 ZALEPLON, ZALEPLON  
 ZIDOVUDINE, ZIDOVUDINE  
 ZOLEDRONIC ACID, ZOLEDRONIC ACID  
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**CLINIGEN HLTHCARE**

\* CLINIGEN HEALTHCARE LTD  
 ETHYOL, AMIFOSTINE  
 FOSCAVIR, FOSCARNET SODIUM  
 TOTECT, DEXRAZOXANE HYDROCHLORIDE

**CLOVER PHARMS**

\* CLOVER PHARMACEUTICALS CORP  
 AMICAR, AMINOCAPROIC ACID

**CLOVIS ONCOLOGY INC**

\* CLOVIS ONCOLOGY INC  
 RUBRACA, RUCAPARIB CAMSYLATE

**CMP DEV LLC**

\* CMP DEVELOPMENT LLC  
 CAROSPIR, SPIRONOLACTONE

**CMP PHARMA INC**

\* CMP PHARMA INC  
 AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
 HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE  
 ISONIAZID, ISONIAZID  
 SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE  
 SPS, SODIUM POLYSTYRENE SULFONATE  
 TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE

**CNTY LINE PHARMS**

\* COUNTY LINE PHARMACEUTICALS LLC  
 BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN  
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
 CICLOPIROX, CICLOPIROX  
 DYNACIN, MINOCYCLINE HYDROCHLORIDE  
 FENOFLIBRATE (MICRONIZED), FENOFLIBRATE  
 FENOFLIBRATE, FENOFLIBRATE  
 FLUOCINONIDE, FLUOCINONIDE  
 LIDEX, FLUOCINONIDE  
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
 TAMBOCOR, FLECAINIDE ACETATE  
 TRANDATE, LABETALOL HYDROCHLORIDE  
 UREX, METHENAMINE HIPPURATE

**COLGATE PALMOLIVE**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

- \* COLGATE PALMOLIVE  
COLGATE TOTAL, SODIUM FLUORIDE (OTC)

**COLGATE PALMOLIVE CO**

- \* COLGATE PALMOLIVE CO  
PERIOGARD, CHLORHEXIDINE GLUCONATE

**COLGATE-PALMOLIVE CO**

- \* COLGATE-PALMOLIVE CO  
PERIOGARD, CHLORHEXIDINE GLUCONATE

**COLLEGIUM PHARM INC**

- \* COLLEGIUM PHARMACEUTICAL INC  
XTAMPZA ER, OXYCODONE

**COMBE**

- \* COMBE INC  
VAGISTAT-1, TIOCONAZOLE (OTC)

**CONCORDIA LABS INC**

- \* CONCORDIA LABORATORIES INC  
PHOTOFRIN, PORFIMER SODIUM

**CONCORDIA PHARMS INC**

- \* CONCORDIA PHARMACEUTICALS INC  
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE  
DUTOPROL, HYDROCHLOROTHIAZIDE  
DYRENium, TRIAMTERENE  
KAPVAY, CLONIDINE HYDROCHLORIDE  
LANOXIN, DIGOXIN  
NILANDRON, NILUTAMIDE  
ORAPRED ODT, PREDNISOLONE SODIUM PHOSPHATE  
PARNATE, TRANLYCYPROMINE SULFATE  
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE  
UROXATRAL, ALFUZOSIN HYDROCHLORIDE

**CONTRACT PHARMACAL**

- \* CONTRACT PHARMACAL CORP  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
IBUPROFEN, IBUPROFEN  
IBUPROFEN, IBUPROFEN (OTC)  
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)  
PROFEN, IBUPROFEN (OTC)

**COOPERSURGICAL**

- \* COOPERSURGICAL INC  
PARAGARD T 380A, COPPER

**CORCEPT THERAP**

- \* CORCEPT THERAPEUTICS INC  
KORLYM, MIFEPRISTONE

**CORDEN PHARMA**

- \* CORDEN PHARMA LATINA SPA  
GLEOSTINE, LOMUSTINE

**COREPHARMA**

- \* COREPHARMA LLC  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE  
GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE  
METAXALONE, METAXALONE  
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
POTASSIUM CITRATE, POTASSIUM CITRATE

**COVIS PHARMA BV**

- \* COVIS PHARMA BV  
ALTOPREV, LOVASTATIN  
BETAPACE AF, SOTALOL HYDROCHLORIDE  
BETAPACE, SOTALOL HYDROCHLORIDE  
LANOXIN PEDIATRIC, DIGOXIN  
LANOXIN, DIGOXIN  
PRILOSEC, OMEPRAZOLE MAGNESIUM  
RILUTEK, RILUZOLE  
SULAR, NISOLDIPINE  
ZANAFLEX, TIZANIDINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

**CPDC**

- \* CENTRE FOR PROBE DEVELOPMENT AND COMMERCIALIZATION  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**CPPI CV**

- \* CP PHARMACEUTICALS INTERNATIONAL CV  
SUTENT, SUNITINIB MALATE

**CRANE PHARMS LLC**

- \* CRANE PHARMACEUTICALS LLC  
DAPTOMYCIN, DAPTOMYCIN

**CROSSMEDIKA SA**

- \* CROSSMEDIKA SA  
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE  
TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

**CROWN LABS**

- \* CROWN LABORATORIES INC  
ALA-CORT, HYDROCORTISONE  
ALA-SCALP, HYDROCORTISONE  
TRIDERM, TRIAMCINOLONE ACETONIDE

**CROWN LABS INC**

- \* CROWN LABORATORIES INC  
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN  
NYSTATIN, NYSTATIN

**CSPC NBP PHARM CO**

- \* CSPC NBP PHARMACEUTICAL CO LTD  
BENZONATATE, BENZONATATE

**CSPC OUYI PHARM CO**

- \* CSPC OUYI PHARMACEUTICAL CO LTD  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
GABAPENTIN, GABAPENTIN  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

**CUBIST PHARMS**

- \* CUBIST PHARMACEUTICALS INC  
ENTEREG, ALVIMOPAN

**CUBIST PHARMS LLC**

- \* CUBIST PHARMACEUTICALS LLC  
CUBICIN RF, DAPTOMYCIN  
CUBICIN, DAPTOMYCIN  
DIFICID, FIDAXOMICIN  
SIVEXTRO, TEDIZOLID PHOSPHATE  
ZERBAXA, CEFTOLOZANE SULFATE

**CUMBERLAND PHARMS**

- \* CUMBERLAND PHARMACEUTICALS INC  
ACETADOTE, ACETYL CYSTEINE  
CALDOLOR, IBUPROFEN  
LACTULOSE, LACTULOSE  
VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER, CONIVAPTAN HYDROCHLORIDE

**CUSTOPHARM INC**

- \* CUSTOPHARM INC  
ACETAMINOPHEN, ACETAMINOPHEN  
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE  
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM

**CUTANEA**

- \* CUTANEA LIFE SCIENCES INC  
AKTIPAK, BENZOYL PEROXIDE

**CUTIS HEALTH LLC**

- \* CUTIS HEALTH LLC  
DORAL, QUAZEPAM

**CYCLE PHARMS LTD**

- \* CYCLE PHARMACEUTICALS LTD  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* C \*\*

\* CYCLE PHARMACEUTICALS LTD  
NITYR, NITISINONE

**CYNDEA PHARMA**

\* CYNDEA PHARMA SL  
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

**CYPRESS PHARM**

\* CYPRESS PHARMACEUTICAL INC  
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE  
ELIPHOS, CALCIUM ACETATE  
REZIRA, HYDROCODONE BITARTRATE  
VITUZ, CHLORPHENIRAMINE MALEATE  
ZUTRIPRO, CHLORPHENIRAMINE MALEATE

\*\* D \*\*

**DAEWOONG PHARM CO**

\* DAEWOONG PHARMACEUTICAL CO LTD  
MEROPENEM, MEROPENEM

**DAIICHI SANKYO**

\* DAIICHI SANKYO INC  
AZOR, AMLODIPINE BESYLATE  
BENICAR HCT, HYDROCHLOROTHIAZIDE  
BENICAR, OLMESARTAN MEDOXOMIL  
TRIBENZOR, AMLODIPINE BESYLATE  
WELCHOL, COLESEVELAM HYDROCHLORIDE

**DAIICHI SANKYO INC**

\* DAIICHI SANKYO INC  
EVOXAC, CEVIMELINE HYDROCHLORIDE  
MORPHABOND ER, MORPHINE SULFATE  
ROXYBOND, OXYCODONE HYDROCHLORIDE  
SAVAYSA, EDOXABAN TOSYLATE

**DANCO LABS LLC**

\* DANCO LABORATORIES LLC  
MIFEPREX, MIFEPRISTONE

**DAVA INTL INC**

\* DAVA INTERNATIONAL INC  
ALPRAZOLAM, ALPRAZOLAM

**DAVA PHARMS INC**

\* DAVA PHARMACEUTICALS INC  
ACYCLOVIR, ACYCLOVIR  
AMOXICILLIN, AMOXICILLIN  
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE  
ATENOLOL, ATENOLOL  
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE  
DIAZEPAM, DIAZEPAM  
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE  
GLYBURIDE (MICRONIZED), GLYBURIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
METHOTREXATE SODIUM, METHOTREXATE SODIUM  
MORPHINE SULFATE, MORPHINE SULFATE  
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM  
PROPYLTHIOURACIL, PROPYLTHIOURACIL  
PYRAZINAMIDE, PYRAZINAMIDE  
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE  
VOSPIRE ER, ALBUTEROL SULFATE

**DAVIS AND GECK**

\* DAVIS AND GECK DIV AMERICAN CYANAMID CO  
PRE-OP II, HEXACHLOROPHENENE  
PRE-OP, HEXACHLOROPHENENE

**DENTSPLY PHARM**

\* DENTSPLY PHARMACEUTICAL INC  
CITANESE FORTE DENTAL, EPINEPHRINE BITARTRATE  
ORAQIX, LIDOCAINE

**DEPOMED INC**

\* DEPOMED INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* D \*\*

- \* DEPOMED INC
  - CAMBIA, DICLOFENAC POTASSIUM
  - GRALISE, GABAPENTIN
  - NUCYNTA ER, TAPENTADOL HYDROCHLORIDE
  - NUCYNTA, TAPENTADOL HYDROCHLORIDE
  - ZIPSOR, DICLOFENAC POTASSIUM

**DEPROCO**

- \* DEPROCO INC
  - LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
  - LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
  - SCANDONEST L, LEVONORDEFRIN
  - SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
  - SEPTOCAIN, ARTICAINE HYDROCHLORIDE

**DEVA HOLDING AS**

- \* DEVA HOLDING AS
  - ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
  - TEMOZOLOMIDE, TEMOZOLOMIDE

**DEXCEL LTD**

- \* DEXCEL LTD
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

**DEXCEL PHARMA**

- \* DEXCEL PHARMA TECHNOLOGIES LTD
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
  - LEVETIRACETAM, LEVETIRACETAM
  - OMEPRAZOLE, OMEPRAZOLE (OTC)
  - PERIOCHIP, CHLORHEXIDINE GLUCONATE

**DFB ONCOLOGY LTD**

- \* DFB ONCOLOGY LTD
  - DOCETAXEL, DOCETAXEL

**DIAGNOSTIC GREEN**

- \* DIAGNOSTIC GREEN GMBH
  - INDOCYANINE GREEN, INDOCYANINE GREEN

**DIALYSIS SUPS**

- \* DIALYSIS SUPPLIES INC
  - NORMOCARB HF 25, MAGNESIUM CHLORIDE
  - NORMOCARB HF 35, MAGNESIUM CHLORIDE

**DIGESTIVE CARE INC**

- \* DIGESTIVE CARE INC
  - PERTZYE, PANCRELIPIASE (AMYLASE)

**DORC**

- \* DORC INTERNATIONAL BV
  - MEMBRANEBLUE, TRYPLAN BLUE
  - VISIONBLUE, TRYPLAN BLUE

**DOUGLAS PHARMS**

- \* DOUGLAS PHARMACEUTICALS AMERICA LTD
  - MYORISAN, ISOTRETINOIN

**DOW PHARM**

- \* DOW PHARMACEUTICAL SCIENCES
  - ACANYA, BENZOYL PEROXIDE
  - ATRALIN, TRETINOIN
  - JUBLIA, EFINACONAZOLE
  - ONEXTON, BENZOYL PEROXIDE
  - OXSORALEN-ULTRA, METHOXSALEN

**DR REDDYS LA**

- \* DR REDDYS LABORATORIES LOUISIANA LLC
  - IBUPROFEN, IBUPROFEN
  - IBUPROFEN, IBUPROFEN (OTC)
  - LOPURIN, ALLOPURINOL
  - SSD AF, SILVER SULFADIAZINE
  - SSD, SILVER SULFADIAZINE

**DR REDDYS LABS INC**

- \* DR REDDYS LABORATORIES INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* D \*\*

- \* DR REDDYS LABORATORIES INC
  - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
  - AMOXIL, AMOXICILLIN
  - AUGMENTIN '125', AMOXICILLIN
  - AUGMENTIN '250', AMOXICILLIN
  - AUGMENTIN '875', AMOXICILLIN
  - AUGMENTIN XR, AMOXICILLIN
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - FINASTERIDE, FINASTERIDE
  - FLUCONAZOLE, FLUCONAZOLE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - IBUPROFEN, IBUPROFEN
  - IBUPROFEN, IBUPROFEN (OTC)
  - LAROTID, AMOXICILLIN
  - LEVOFLOXACIN, LEVOFLOXACIN
  - MELOXICAM, MELOXICAM
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
  - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
  - NITROGLYCERIN, NITROGLYCERIN
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PROGESTERONE, PROGESTERONE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - SIMVASTATIN, SIMVASTATIN
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**DR REDDYS LABS INTL**

- \* DR REDDYS LABORATORIES INTERNATIONAL SA
  - RAMELTEON, RAMELTEON

**DR REDDYS LABS LTD**

- \* DR REDDYS LABORATORIES LIMITED
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- \* DR REDDYS LABORATORIES LTD
  - ALENDRONATE SODIUM, ALENDRONATE SODIUM
  - AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - ANASTROZOLE, ANASTROZOLE
  - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - AZACITIDINE, AZACITIDINE
  - BIVALIRUDIN, BIVALIRUDIN
  - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
  - CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
  - CARVEDILOL, CARVEDILOL
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - CLOFARABINE, CLOFARABINE
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - DECITABINE, DECITABINE
  - DESLORATADINE AND PSEUDOEPHEDRINE SULFATE 24 HOUR, DESLORATADINE
  - DESLORATADINE, DESLORATADINE
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - DOCETAXEL, DOCETAXEL
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* D \*\*

- \* DR REDDYS LABORATORIES LTD
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - ESZOPICLONE, ESZOPICLONE
  - FAMOTIDINE, FAMOTIDINE
  - FAMOTIDINE, FAMOTIDINE (OTC)
  - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
  - FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
  - FINASTERIDE, FINASTERIDE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FONDAPARINUX SODIUM, FONDAPARINUX SODIUM
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - GLIMEPIRIDE, GLIMEPIRIDE
  - GLCOPYRROLATE, GLCOPYRROLATE
  - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
  - GUAIFENESIN AND PSEUDOEPHEDRINE HYDROCHLORIDE, GUAIFENESIN (OTC)
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
  - IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
  - IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IRBESARTAN, IRBESARTAN
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - LAMOTRIGINE, LAMOTRIGINE
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
  - LATANOPROST, LATANOPROST
  - LETROZOLE, LETROZOLE
  - LEVETIRACETAM, LEVETIRACETAM
  - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MINOLIRA, MINOCYCLINE HYDROCHLORIDE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - NAPROXEN AND ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - NAPROXEN SODIUM, NAPROXEN SODIUM
  - NATEGLINIDE, NATEGLINIDE
  - NIZATIDINE, NIZATIDINE
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - OFLOXACIN, OFLOXACIN
  - OLANZAPINE, OLANZAPINE
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
  - OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
  - OMEPRAZOLE, OMEPRAZOLE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXaprozin, OXaprozin
  - PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PARICALCITOL, PARICALCITOL
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - RAMIPRIL, RAMIPRIL
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
  - RISPERIDONE, RISPERIDONE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SEVELAMER CARBONATE, SEVELAMER CARBONATE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - SIROLIMUS, SIROLIMUS
  - TACROLIMUS, TACROLIMUS

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* D \*\*

- \* DR REDDYS LABORATORIES LTD
  - TETRABENAZINE, TETRABENAZINE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - VINORELBINE TARTRATE, VINORELBINE TARTRATE
  - ZAFIRLUKAST, ZAFIRLUKAST
  - ZEMBRACE SYMTOUCH, SUMATRIPTAN SUCCINATE
  - ZENATANE, ISOTRETINOIN
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**DR REDDYS LABS SA**

- \* DR REDDYS LABORATORIES SA
  - EZETIMIBE AND SIMVASTATIN, EZETIMIBE
  - FENOFIBRATE (MICRONIZED), FENOFIBRATE
  - HABITROL, NICOTINE (OTC)
  - MERCAPTOPURINE, MERCAPTOPURINE

**DRAIMAGE**

- \* DRAIMAGE INC
  - DTPA, TECHNETIUM TC-99M PENTETATE KIT
  - TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

**DUCHESNAY**

- \* DUCHESNAY INC
  - BONJESTA, DOXYLAMINE SUCCINATE
  - DICLEGIS, DOXYLAMINE SUCCINATE
  - OSPHENA, OSPEMIFENE

**DURAMED PHARMS BARR**

- \* DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
  - AVIANE-28, ETHINYL ESTRADIOL
  - CRYSELLE, ETHINYL ESTRADIOL
  - DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
  - ENPRESSE-28, ETHINYL ESTRADIOL
  - METHYLPREDNISOLONE, METHYLPREDNISOLONE
  - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - VELIVET, DESOGESTREL

**DURAMED RES**

- \* DURAMED RESEARCH INC
  - AYGESTIN, NORETHINDRONE ACETATE

**DUSA**

- \* DUSA PHARMACEUTICALS INC
  - LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

**REDDYS**

- \* DOCTOR REDDYS LABORATORIES LTD
  - DESLORATADINE, DESLORATADINE
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE

\*\* E \*\*

**EAGLE PHARMS**

- \* EAGLE PHARMACEUTICALS INC
  - ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
  - BENDEKA, BENDAMUSTINE HYDROCHLORIDE
  - DOCETAXEL, DOCETAXEL
  - RYANODEX, DANTROLENE SODIUM

**EASTMAN KODAK**

- \* EASTMAN KODAK CO
  - LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE

**ECI PHARMS LLC**

- \* ECI PHARMACEUTICALS LLC
  - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
  - CALCIUM ACETATE, CALCIUM ACETATE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - LAMIVUDINE, LAMIVUDINE
  - LEVETIRACETAM, LEVETIRACETAM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* E \*\*

\* ECI PHARMACEUTICALS LLC  
METHIMAZOLE, METHIMAZOLE  
PARICALCITOL, PARICALCITOL  
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE  
VALPROIC ACID, VALPROIC ACID

**ECLAT PHARMS LLC**

\* ECLAT PHARMACEUTICALS LLC  
BLOXIVERZ, NEOSTIGMINE METHYLSULFATE

**ECOLAB**

\* ECOLAB INC  
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)  
CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

**ECR**

\* ECR PHARMACEUTICALS  
DEXAMETHASONE, DEXAMETHASONE

**ECR PHARMA**

\* ECR PHARMA  
TUSSICAPS, CHLORPHENIRAMINE POLISTIREX

**EDENBRIDGE PHARMS**

\* EDENBRIDGE PHARMACEUTICALS LLC  
CARBIDOPA, CARBIDOPA  
ETHACRYNIC ACID, ETHACRYNIC ACID  
IVERMECTIN, IVERMECTIN  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE  
TINIDAZOLE, TINIDAZOLE

**EDISON THERAPS LLC**

\* EDISON THERAPEUTICS LLC  
METHERGINE, METHYLERGONOVINE MALEATE

**EGALET**

\* EGALET CORP  
ARYMO ER, MORPHINE SULFATE

**EGALET US INC**

\* EGALET US INC  
OXAYDO, OXYCODONE HYDROCHLORIDE  
SPRIX, KETOROLAC TROMETHAMINE

**EI INC**

\* EI INC  
THEROXIDIL, MINOXIDIL (OTC)

**EISAI INC**

\* EISAI INC  
ACIPHEX, RABEPRAZOLE SODIUM  
ARICEPT ODT, DONEPEZIL HYDROCHLORIDE  
ARICEPT, DONEPEZIL HYDROCHLORIDE  
BANZEL, RUFINAMIDE  
BELVIQ XR, LORCASERIN HYDROCHLORIDE  
BELVIQ, LORCASERIN HYDROCHLORIDE  
FYCOMPA, PERAMPANEL  
HALAVEN, ERIBULIN MESYLATE  
HEXALEN, ALTRETAMINE  
LENVIMA, LENVATINIB MESYLATE  
PANRETIN, ALITRETNINOIN  
SALAGEN, PILOCARPINE HYDROCHLORIDE

**ELEFSEE PHARMS INTL**

\* ELEFSEE PHARMACEUTICALS INTERNATIONAL LTD  
LAZANDA, FENTANYL CITRATE

**ELI LILLY AND CO**

\* ELI LILLY AND CO  
BASAGLAR, INSULIN GLARGINE  
EFFIENT, PRASUGREL HYDROCHLORIDE  
HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT  
PROZAC, FLUOXETINE HYDROCHLORIDE  
VERZENIO, ABEMACICLIB

**ELI LILLY CO**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* E \*\*

\* ELI LILLY CO  
 ADCIRCA, TADALAFIL  
 ZYPREXA RELPREVV, OLANZAPINE PAMOATE

**ELITE LABS**

\* ELITE LABORATORIES INC  
 HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE  
 NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE  
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

**ELITE LABS INC**

\* ELITE LABORATORIES INC  
 DANTROLENE SODIUM, DANTROLENE SODIUM  
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
 ISRADIPINE, ISRADIPINE  
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE  
 PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE  
 PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE  
 TRIMIPRAMINE MALEATE, TRIMIPRAMINE MALEATE

**EMCURE PHARMS**

\* EMCURE PHARMACEUTICALS LTD  
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE

**EMCURE PHARMS INDIA**

\* EMCURE PHARMACEUTICALS LTD INDIA  
 FOSINOPRIL SODIUM AND HYDROCHLORTIAZIDE, FOSINOPRIL SODIUM

**EMCURE PHARMS LTD**

\* EMCURE PHARMACEUTICALS LTD  
 ACARBOSE, ACARBOSE  
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM  
 ADENOSINE, ADENOSINE  
 AMIKACIN SULFATE, AMIKACIN SULFATE  
 BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE  
 BICNU, CARMUSTINE  
 CIDOFOVIR, CIDOFOVIR  
 COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM  
 ETOMIDATE, ETOMIDATE  
 FUROSEMIDE, FUROSEMIDE  
 GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE  
 IBANDRONATE SODIUM, IBANDRONATE SODIUM  
 IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
 LEVOFLOXACIN, LEVOFLOXACIN  
 METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE  
 ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE  
 RIFAMPIN, RIFAMPIN  
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
 TRANEXAMIC ACID, TRANEXAMIC ACID  
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE  
 ZOLEDRONIC ACID, ZOLEDRONIC ACID

**EMD SERONO**

\* EMD SERONO INC  
 GONAL-F RFF REDI-JECT, FOLLITROPIN ALFA/BETA  
 GONAL-F RFF, FOLLITROPIN ALFA/BETA  
 GONAL-F, FOLLITROPIN ALFA/BETA  
 OVIDREL, CHORIOGONADOTROPIN ALFA  
 SAIZEN, SOMATROPIN RECOMBINANT  
 SEROSTIM, SOMATROPIN RECOMBINANT  
 ZORBTIVE, SOMATROPIN RECOMBINANT

**EMD SERONO INC**

\* EMD SERONO INC  
 CETROTIDE, CETRORELIX

**EMERALD INTL LTD**

\* EMERALD INTERNATIONAL LTD  
 BACLOFEN, BACLOFEN

**EMMAUS MEDCL**

\* EMMAUS MEDICAL INC  
 ENDARI, L-GLUTAMINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* E \*\*

- \* EMMAUS MEDICAL INC  
NUTRESTORE, L-GLUTAMINE

**ENDO PHARM**

- \* ENDO PHARMACEUTICAL SOLUTIONS INC  
SUPPRELIN LA, HISTRELIN ACETATE  
VALSTAR PRESERVATIVE FREE, VALRUBICIN  
VANTAS, HISTRELIN ACETATE

**ENDO PHARMS**

- \* ENDO PHARMACEUTICALS INC  
DELATESTRYL, TESTOSTERONE ENANTHATE  
FORTESTA, TESTOSTERONE  
FROVA, FROVATRIPTAN SUCCINATE  
OPANA, OXYMORPHONE HYDROCHLORIDE  
PERCODAN, ASPIRIN

**ENDO PHARMS INC**

- \* ENDO PHARMACEUTICALS INC  
AVEED, TESTOSTERONE UNDECANOATE  
COLY-MYCIN S, COLISTIN SULFATE  
MEGACE ES, MEGESTROL ACETATE  
NASCOBAL, CYANOCOBALAMIN  
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE

**ENDO VENTURES LTD**

- \* ENDO VENTURES LTD IRELAND  
SUMAVEL DOSEPRO, SUMATRIPTAN SUCCINATE

**EPI HLTH**

- \* EPI HEALTH LLC  
SITAVIG, ACYCLOVIR

**EPIC PHARMA**

- \* EPIC PHARMA INC  
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE  
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
- \* EPIC PHARMA LLC  
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE  
SULINDAC, SULINDAC  
TRANDOLAPRIL, TRANDOLAPRIL  
URSODIOL, URSODIOL

**EPIC PHARMA INC**

- \* EPIC PHARMA INC  
ESTRADIOL, ESTRADIOL  
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

**EPIC PHARMA LLC**

- \* EPIC PHARMA LLC  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE  
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
GABAPENTIN, GABAPENTIN  
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE  
GLYBURIDE, GLYBURIDE  
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE  
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE  
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE  
MOLINDONE HYDROCHLORIDE, MOLINDONE HYDROCHLORIDE  
MORPHINE SULFATE, MORPHINE SULFATE  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE  
PHENYTOIN, PHENYTOIN  
PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE  
SULINDAC, SULINDAC  
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* E \*\*

**ESSENTIAL ISOTOPES**

- \* ESSENTIAL ISOTOPES LLC  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**ETHYPHARM**

- \* ETHYPHARM  
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

**ETHYPHARM USA CORP**

- \* ETHYPHARM USA CORP  
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

**EUROHLTH INTL SARL**

- \* EUROHEALTH INTERNATIONAL SARL  
DROPERIDOL, DROPERIDOL  
MORPHINE SULFATE, MORPHINE SULFATE  
NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE

**EXALENZ BIOSCIENCE**

- \* EXALENZ BIOSCIENCE LTD  
IDKIT:HP, CITRIC ACID

**EXELA HOLDINGS**

- \* EXELA HOLDINGS INC  
DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM

**EXELA PHARMA SCIENCE**

- \* EXELA PHARMA SCIENCES  
CAFFEINE CITRATE, CAFFEINE CITRATE  
IBUPROFEN LYSINE, IBUPROFEN LYSINE  
NICARDIPINE HYDROCHLORIDE IN 0.9% SODIUM CHLORIDE, NICARDIPINE HYDROCHLORIDE  
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE  
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE

**EXELA PHARMA SCS LLC**

- \* EXELA PHARMA SCIENCES LLC  
CAFFEINE CITRATE, CAFFEINE CITRATE  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
GANCICLOVIR, GANCICLOVIR  
MAGNESIUM SULFATE, MAGNESIUM SULFATE  
NIPRIDE RTU IN SODIUM CHLORIDE 0.9%, SODIUM NITROPRUSSIDE  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM  
POTASSIUM ACETATE, POTASSIUM ACETATE  
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**EXELIXIS**

- \* EXELIXIS INC  
COMETRIQ, CABOZANTINIB S-MALATE

**EXELIXIS INC**

- \* EXELIXIS INC  
CABOMETYX, CABOZANTINIB S-MALATE

**EXELTIS USA INC**

- \* EXELTIS USA INC  
ESTRASORB, ESTRADIOL HEMIHYDRATE

**EYEVANCE PHARMS**

- \* EYEVANCE PHARMACEUTICALS LLC  
ZERVIALE, CETIRIZINE HYDROCHLORIDE

**LILLY**

- \* ELI LILLY AND CO  
ALIMTA, PEMETREXED DISODIUM  
CIALIS, TADALAFIL  
CYMBALTA, DULOXETINE HYDROCHLORIDE  
EVISTA, RALOXIFENE HYDROCHLORIDE  
FORTEO, TERIPARATIDE RECOMBINANT HUMAN  
GEMZAR, GEMCITABINE HYDROCHLORIDE  
GLUCAGON, GLUCAGON RECOMBINANT  
HUMALOG KWIKPEN, INSULIN LISPRO RECOMBINANT  
HUMALOG MIX 50/50 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT  
HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT  
HUMALOG MIX 75/25 KWIKPEN, INSULIN LISPRO PROTAMINE RECOMBINANT

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* E \*\*

- \* ELI LILLY AND CO
  - HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
  - HUMALOG, INSULIN LISPRO RECOMBINANT
  - HUMATROPE, SOMATROPIN RECOMBINANT
  - HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)
  - HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
  - HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
  - HUMULIN R KWIKPEN, INSULIN HUMAN
  - HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)
  - HUMULIN R, INSULIN HUMAN
  - HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
  - PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
  - STRATTERA, ATOMOXETINE HYDROCHLORIDE
  - SYMBYAX, FLUOXETINE HYDROCHLORIDE
  - ZYPREXA ZYDIS, OLANZAPINE
  - ZYPREXA, OLANZAPINE

\*\* F \*\*

**FDC LTD**

- \* FDC LTD
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - LATANOPROST, LATANOPROST
  - OFLOXACIN, OFLOXACIN
  - TIMOLOL MALEATE, TIMOLOL MALEATE

**FDN CONSUMER**

- \* FOUNDATION CONSUMER HEALTHCARE LLC
  - LEVONORGESTREL, LEVONORGESTREL (OTC)
  - PLAN B ONE-STEP, LEVONORGESTREL (OTC)

**FEINSTEIN**

- \* FEINSTEIN INSTITUTE MEDICAL RESEARCH
  - AMMONIA N 13, AMMONIA N-13
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**FERA PHARMS**

- \* FERA PHARMACEUTICALS LLC
  - TOBRAMYCIN, TOBRAMYCIN

**FERA PHARMS LLC**

- \* FERA PHARMACEUTICALS LLC
  - LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM

**FERRER INTERNACIONAL**

- \* FERRER INTERNACIONAL SA
  - XEPI, OZENOXACIN

**FERRING**

- \* FERRING PHARMACEUTICALS INC
  - ACTHREL, CORTICORELIN OVINE TRIFLUTATE
  - BRAVELLE, UROFOLLITROPIN
  - CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
  - ENDOMETRIN, PROGESTERONE
  - FIRMAGON, DEGARELIX ACETATE
  - MENOPUR, MENOTROPINS (FSH)
  - MINIRIN, DESMOPRESSIN ACETATE
  - ZOMACTON, SOMATROPIN RECOMBINANT

**FERRING PHARMS INC**

- \* FERRING PHARMACEUTICALS INC
  - CERVIDIL, DINOPROSTONE
  - CLENPIQ, CITRIC ACID
  - DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
  - DDAVP, DESMOPRESSIN ACETATE
  - LYSTEDA, TRANEXAMIC ACID
  - PREPOPIK, CITRIC ACID
  - STIMATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE

**FLAMEL IRELAND LTD**

- \* FLAMEL IRELAND LIMITED
  - AKOVAZ, EPHEDRINE SULFATE

**FLAMINGO PHARMS**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* F \*\*

- \* FLAMINGO PHARMACEUTICALS LTD  
METRONIDAZOLE, METRONIDAZOLE  
PIROXICAM, PIROXICAM

**FLEXION THERAPS INC**

- \* FLEXION THERAPEUTICS INC  
ZILRETTA, TRIAMCINOLONE ACETONIDE

**FOREST LABS**

- \* FOREST LABORATORIES INC  
BYSTOLIC, NEBIVOLOL HYDROCHLORIDE  
CELEXA, CITALOPRAM HYDROBROMIDE  
LEXAPRO, ESCITALOPRAM OXALATE  
THYROLAR-0.25, LIOTRIX (T4)  
THYROLAR-0.5, LIOTRIX (T4)  
THYROLAR-1, LIOTRIX (T4)  
THYROLAR-2, LIOTRIX (T4)  
THYROLAR-3, LIOTRIX (T4)

**FOREST LABS INC**

- \* FOREST LABORATORIES INC  
BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE  
BENTYL, DICYCLOMINE HYDROCHLORIDE  
CARAFATE, SUCRALFATE  
FETZIMA, LEVOMILNACIPRAN HYDROCHLORIDE  
RECTIV, NITROGLYCERIN  
ULTRESA, PANCRELIPIASE (AMYLASE)  
URSO 250, URSODIOL  
URSO FORTE, URSODIOL  
VIOKACE, PANCRELIPIASE (AMYLASE)  
ZENPEP, PANCRELIPIASE (AMYLASE)

**FOREST LABS LLC**

- \* FOREST LABORATORIES LLC  
BYVALSON, NEBIVOLOL HYDROCHLORIDE  
CANASA, MESALAMINE  
LINZESS, LINACLOTIDE  
NAMENDA XR, MEMANTINE HYDROCHLORIDE  
NAMENDA, MEMANTINE HYDROCHLORIDE  
NAMZARIC, DONEPEZIL HYDROCHLORIDE  
PYLERA, BISMUTH SUBCITRATE POTASSIUM  
SAPHRIS, ASENAPINE MALEATE  
VIIBRYD, VILAZODONE HYDROCHLORIDE

**FOREST RES INST INC**

- \* FOREST RESEARCH INSTITUTE INC  
VRAYLAR, CARIPRAZINE HYDROCHLORIDE

**FOUGERA PHARMS**

- \* FOUGERA PHARMACEUTICALS INC  
ADAPALENE, ADAPALENE  
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE  
AMCINONIDE, AMCINONIDE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CALCIPOTRIENE, CALCIPOTRIENE  
CICLOPIROX, CICLOPIROX  
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
CLOBETASOL PROPIONATE (EMOLlient), CLOBETASOL PROPIONATE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CLOTRIMAZOLE, CLOTRIMAZOLE  
CUTIVATE, FLUTICASONE PROPIONATE  
DESONIDE, DESONIDE  
DESOXIMETASONE, DESOXIMETASONE  
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE  
ECONAZOLE NITRATE, ECONAZOLE NITRATE  
ERYTHROMYCIN, ERYTHROMYCIN  
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE  
FLUOCINONIDE, FLUOCINONIDE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* F \*\*

- \* FOUGERA PHARMACEUTICALS INC
  - HYDROCORTISONE, HYDROCORTISONE
  - IMIQUIMOD, IMIQUIMOD
  - KETOCONAZOLE, KETOCONAZOLE
  - LIDOCAINE AND PRILOCAINE, LIDOCAINE
  - METRONIDAZOLE, METRONIDAZOLE
  - MOMETASONE FUROATE, MOMETASONE FUROATE
  - MUPIROCIN, MUPIROCIN
  - NYSTATIN, NYSTATIN
  - OXISTAT, OXICONAZOLE NITRATE
  - PANDEL, HYDROCORTISONE PROBUTATE
  - PREDNICARBATE, PREDNICARBATE
  - SOLARAZE, DICLOFENAC SODIUM
  - SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
  - TERCONAZOLE, TERCONAZOLE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
  - TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

**FOUGERA PHARMS INC**

- \* FOUGERA PHARMACEUTICALS INC
  - ACYCLOVIR, ACYCLOVIR
  - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
  - BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
  - FLUOCINONIDE, FLUOCINONIDE
  - GENTAMICIN SULFATE, GENTAMICIN SULFATE
  - HYDROCORTISONE, HYDROCORTISONE
  - LIDOCAINE, LIDOCAINE
  - NITROGLYCERIN, NITROGLYCERIN
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - NYSTATIN, NYSTATIN
  - TACROLIMUS, TACROLIMUS

**FRESENIUS**

- \* FRESENIUS KABI DEUTSCHLAND GMBH
  - INTRALIPID 10%, SOYBEAN OIL
  - INTRALIPID 20%, SOYBEAN OIL
  - INTRALIPID 30%, SOYBEAN OIL

**FRESENIUS KABI**

- \* FRESENIUS KABI AUSTRIA GMBH
  - LACTULOSE, LACTULOSE

**FRESENIUS KABI ONCOL**

- \* FRESENIUS KABI ONCOLOGY PLC
  - ANASTROZOLE, ANASTROZOLE
  - BICALUTAMIDE, BICALUTAMIDE
  - CARBOPLATIN, CARBOPLATIN
  - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - LETROZOLE, LETROZOLE
  - OXALIPLATIN, OXALIPLATIN
  - PACLITAXEL, PACLITAXEL

**FRESENIUS KABI USA**

- \* FRESENIUS KABI USA LLC
  - ACETAMINOPHEN, ACETAMINOPHEN
  - ACETYLCYSTEINE, ACETYLCYSTEINE
  - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
  - ADENOSINE, ADENOSINE
  - AMIKACIN SULFATE, AMIKACIN SULFATE
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - ARGATROBAN, ARGATROBAN
  - ASTRAMORPH PF, MORPHINE SULFATE
  - AZITHROMYCIN, AZITHROMYCIN
  - AZTREONAM, AZTREONAM
  - BACITRACIN, BACITRACIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* F \*\*

- \* FRESENIUS KABI USA LLC
  - BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
  - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
  - BIVALIRUDIN, BIVALIRUDIN
  - BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
  - BORTEZOMIB, BORTEZOMIB
  - CAFFEINE CITRATE, CAFFEINE CITRATE
  - CALCIUM GLUCONATE, CALCIUM GLUCONATE
  - CARBOPLATIN, CARBOPLATIN
  - CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
  - CEFOTETAN, CEFOTETAN DISODIUM
  - CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
  - CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
  - CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
  - CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
  - CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
  - CISPLATIN, CISPLATIN
  - CLADRBINE, CLADRBINE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
  - COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
  - CYTARABINE, CYTARABINE
  - DACARBAZINE, DACARBAZINE
  - DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
  - DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
  - DEXAMETHASONE SODIUM PHOSPHATE PRESERVATIVE FREE, DEXAMETHASONE SODIUM PHOSPHATE
  - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
  - DILAUDID, HYDROMORPHONE HYDROCHLORIDE
  - DILAUDID-HP, HYDROMORPHONE HYDROCHLORIDE
  - DIMENHYDRINATE, DIMENHYDRINATE
  - DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
  - DIPRIVAN, PROPOFOL
  - DIPYRIDAMOLE, DIPYRIDAMOLE
  - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
  - DOXY 100, DOXYCYCLINE HYCLATE
  - DOXY 200, DOXYCYCLINE HYCLATE
  - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
  - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
  - ETOPOSIDE, ETOPOSIDE
  - FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
  - FAMOTIDINE, FAMOTIDINE
  - FLOXURIDINE, FLOXURIDINE
  - FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
  - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
  - FLUMAZENIL, FLUMAZENIL
  - FLUOROURACIL, FLUOROURACIL
  - FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
  - FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
  - FOLIC ACID, FOLIC ACID
  - FOSAPREPITANT DIMEGLUMINE, FOSAPREPITANT DIMEGLUMINE
  - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
  - FUROSEMIDE, FUROSEMIDE
  - GANCICLOVIR, GANCICLOVIR SODIUM
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - GENTAMICIN SULFATE, GENTAMICIN SULFATE
  - GLUCAGON, GLUCAGON HYDROCHLORIDE
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - GRANISETRON HYDROCHLORIDE PRESERVATIVE FREE, GRANISETRON HYDROCHLORIDE
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
  - HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* F \*\*

- \* FRESENIUS KABI USA LLC
  - HEPARIN SODIUM, HEPARIN SODIUM
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
  - IFOSFAMIDE, IFOSFAMIDE
  - INDOMETHACIN, INDOMETHACIN
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - KABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
  - KANAMYCIN SULFATE, KANAMYCIN SULFATE
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
  - LEVOOTHYROXINE SODIUM, LEVOOTHYROXINE SODIUM
  - LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
  - LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
  - LINEZOLID, LINEZOLID
  - MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
  - MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
  - MAGNESIUM SULFATE, MAGNESIUM SULFATE
  - MAGNESIUM SULFATE, MAGNESIUM SULFATE
  - MANNITOL 25%, MANNITOL
  - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
  - MESNA, MESNA
  - METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
  - METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
  - METHOTREXATE SODIUM, METHOTREXATE SODIUM
  - METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
  - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE
  - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
  - MILRINONE LACTATE, MILRINONE LACTATE
  - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - NAROPIN, ROPIVACAINE HYDROCHLORIDE
  - NEBUPENT, PENTAMIDINE ISETHIONATE
  - NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
  - NESACAIN, CHLOROPROCAINE HYDROCHLORIDE
  - NESACAIN-MPF, CHLOROPROCAINE HYDROCHLORIDE
  - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
  - OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
  - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXALIPLATIN, OXALIPLATIN
  - OXYTOCIN, OXYTOCIN
  - PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE
  - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
  - PENTAM, PENTAMIDINE ISETHIONATE
  - PERIKABIVEN IN PLASTIC CONTAINER, AMINO ACIDS
  - POLOCAINE, MEPIVACAINE HYDROCHLORIDE
  - POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
  - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
  - PROGESTERONE, PROGESTERONE
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
  - PROTAMINE SULFATE, PROTAMINE SULFATE
  - PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
  - RIFAMPIN, RIFAMPIN
  - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
  - SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
  - SENSORCAINE, BUPIVACAINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* F \*\*

- \* FRESENIUS KABI USA LLC
  - SMOFLIPID 20%, FISH OIL
  - SODIUM ACETATE, SODIUM ACETATE
  - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
  - STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION
  - STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TERBUTALINE SULFATE, TERBUTALINE SULFATE
  - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
  - TIGECYCLINE, TIGECYCLINE
  - TOBRAMYCIN SULFATE (PHARMACY BULK), TOBRAMYCIN SULFATE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - TRANEXAMIC ACID, TRANEXAMIC ACID
  - VALPROATE SODIUM, VALPROATE SODIUM
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VIBISONE, CYANOCOBALAMIN
  - VINBLASTINE SULFATE, VINBLASTINE SULFATE
  - VINORELBINE TARTRATE, VINORELBINE TARTRATE
  - XYLOCAINE 4% PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
  - XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
  - XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
  - XYLOCAINE, LIDOCAINE HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**FRESENIUS MEDCL**

- \* FRESENIUS MEDICAL CARE NORTH AMERICA
  - DELFLEX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DELFLEX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - PHOSLO GELCAPS, CALCIUM ACETATE
  - PHOSLYRA, CALCIUM ACETATE
  - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**FRONTIDA BIOPHARM**

- \* FRONTIDA BIOPHARM INC
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
  - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE

\*\* G \*\*

**G AND W LABS**

- \* G AND W LABORATORIES INC
  - ACEPHEN, ACETAMINOPHEN (OTC)
  - CICLOPIROX, CICLOPIROX
  - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
  - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
  - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
  - INDOMETHACIN, INDOMETHACIN
  - METRONIDAZOLE, METRONIDAZOLE
  - MICONAZOLE 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
  - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
  - MOMETASONE FUROATE, MOMETASONE FUROATE
  - PROCHLORPERAZINE, PROCHLORPERAZINE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**G AND W LABS INC**

- \* G AND W LABORATORIES INC
  - ACYCLOVIR, ACYCLOVIR
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - BETA-VAL, BETAMETHASONE VALERATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* G \*\*

- \* G AND W LABORATORIES INC
  - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
  - CALCIPOTRIENE, CALCIPOTRIENE
  - CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
  - CICLOPIROX, CICLOPIROX
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - DESONIDE, DESONIDE
  - DESOXIMETASONE, DESOXIMETASONE
  - DOXYCYCLINE, DOXYCYCLINE
  - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
  - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
  - FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
  - FLUOCINONIDE, FLUOCINONIDE
  - GENTAMICIN SULFATE, GENTAMICIN SULFATE
  - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
  - IMIQUIMOD, IMIQUIMOD
  - MESALAMINE, MESALAMINE
  - METRONIDAZOLE, METRONIDAZOLE
  - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
  - MYKACET, NYSTATIN
  - NYSTATIN, NYSTATIN
  - PROMETH VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE
  - PROMETH W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
  - QUINIDINE SULFATE, QUINIDINE SULFATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - TAZAROTENE, TAZAROTENE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**GALDERMA LABS**

- \* GALDERMA LABORATORIES INC
  - CLOBEX, CLOBETASOL PROPIONATE
  - EPIDUO FORTE, ADAPALENE

**GALDERMA LABS LP**

- \* GALDERMA LABORATORIES L P
  - CLOBEX, CLOBETASOL PROPIONATE
- \* GALDERMA LABORATORIES LP
  - CAPEX, FLUOCINOLONE ACETONIDE
  - CLOBEX, CLOBETASOL PROPIONATE
  - DESOWEN, DESONIDE
  - DIFFERIN, ADAPALENE
  - DIFFERIN, ADAPALENE (OTC)
  - EPIDUO, ADAPALENE
  - METROCREAM, METRONIDAZOLE
  - METROGEL, METRONIDAZOLE
  - METROLOTION, METRONIDAZOLE
  - MIRVASO, BRIMONIDINE TARTRATE
  - ORACEA, DOXYCYCLINE
  - SOOLANTRA, IVERMECTIN
  - TRI-LUMA, FLUOCINOLONE ACETONIDE
  - VECTICAL, CALCITRIOL

**GALEN SPECIALTY**

- \* GALEN SPECIALTY PHARMA US LLC
  - SYNERA, LIDOCAINE

**GALEN UK**

- \* GALEN LTD
  - ADASUVE, LOXAPINE

**GASTROENTERO**

- \* GASTROENTERO LOGIC LLC
  - OMEPRAZOLE AND CLARITHROMYCIN AND AMOXICILLIN, AMOXICILLIN

**GATE PHARMS**

- \* GATE PHARMACEUTICALS
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - LINEZOLID, LINEZOLID

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* G \*\*

**GATOR PHARMS**

- \* GATOR PHARMACEUTICALS INC  
COLPREP KIT, MAGNESIUM SULFATE

**GAVIS PHARMS**

- \* GAVIS PHARMACEUTICALS LLC  
NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE  
NYSTATIN, NYSTATIN  
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE  
QUINARETIC, HYDROCHLOROTHIAZIDE  
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

**GAVIS PHARMS LLC**

- \* GAVIS PHARMACEUTICALS LLC  
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

**GD SEARLE**

- \* GD SEARLE LLC  
CELEBREX, CELECOXIB  
DAYPRO, OXaprozin

**GD SEARLE LLC**

- \* GD SEARLE LLC  
ALDACTAZIDE, HYDROCHLOROTHIAZIDE  
ALDACTONE, SPIRONOLACTONE  
ARTHROTEC, DICLOFENAC SODIUM  
CALAN, VERAPAMIL HYDROCHLORIDE  
CYTOTEC, MISOPROSTOL  
FLAGYL ER, METRONIDAZOLE  
FLAGYL, METRONIDAZOLE  
INSPRA, EPLERENONE  
LOMOTIL, ATROPINE SULFATE  
NORPACE CR, DISOPYRAMIDE PHOSPHATE  
NORPACE, DISOPYRAMIDE PHOSPHATE  
SYNAREL, NAFARELIN ACETATE

**GE HEALTHCARE**

- \* GE HEALTHCARE  
ADREVIEW, IOBENGUANE SULFATE I-123  
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT  
INDICLOR, INDIUM IN-111 CHLORIDE  
INDIUM IN 111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE  
METASTRON, STRONTIUM CHLORIDE SR-89  
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM  
MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT  
OMNIPAQ 140, IOHEXOL  
OMNIPAQ 180, IOHEXOL  
OMNIPAQ 240, IOHEXOL  
OMNIPAQ 300, IOHEXOL  
OMNIPAQ 350, IOHEXOL  
OMNISCAN, GADODIAMIDE  
OPTISON, ALBUMIN HUMAN  
TECHNETIUM TC 99M GENERATOR, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR  
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201  
VISIPAQ 270, IODIXANOL  
VISIPAQ 320, IODIXANOL  
VIZAMYL, FLUTEMETAMOL F-18

**GE HLTHCARE INC**

- \* GE HEALTHCARE INC  
DATSCAN, IOFLUPANE I-123

**GEDEON RICHTER USA**

- \* GEDEON RICHTER USA INC  
FINASTERIDE, FINASTERIDE

**GEMINI LABS LLC**

- \* GEMINI LABORATORIES LLC  
OXANDRIN, OXANDROLONE  
PRANDIN, REPAGLINIDE

**GENENTECH**

- \* GENENTECH INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* G \*\*

- \* GENENTECH INC
  - ERIVEDGE, VISMODEGIB
  - NUTROPIN AQ NUSPIN, SOMATROPIN RECOMBINANT
  - NUTROPIN AQ PEN, SOMATROPIN RECOMBINANT

**GENENTECH INC**

- \* GENENTECH INC
  - COTELLIC, COBIMETINIB FUMARATE
  - ESBRIET, PIRFENIDONE

**GENUS LIFESCIENCES**

- \* GENUS LIFE SCIENCES INC
  - GOPRELTO, COCAINE HYDROCHLORIDE
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

**GENZYME**

- \* GENZYME CORP
  - CEREZYME, IMIGLUCERASE
  - CLOLAR, CLOFARABINE
  - MOZOBIL, PLERIXAFOR
  - RENAGEL, SEVELAMER HYDROCHLORIDE
  - RENELA, SEVELAMER CARBONATE
  - THYROGEN, THYROTROPIN ALFA

**GENZYME CORP**

- \* GENZYME CORP
  - CAPRELSA, VANDETANIB
  - CERDELGA, ELIGLUSTAT TARTRATE
  - HECTOROL, DOXERCALCIFEROL

**GILEAD**

- \* GILEAD SCIENCES INC
  - ATRIPLA, EFAVIRENZ
  - CAYSTON, AZTREONAM
  - EMTRIVA, EMTRICITABINE
  - HEPSERA, ADEFOVIR DIPIVOXIL
  - LETAIRIS, AMBRISENTAN
  - RANEXA, RANOLAZINE
  - TRUVADA, EMTRICITABINE

**GILEAD SCIENCES INC**

- \* GILEAD SCIENCES INC
  - COMPLERA, EMTRICITABINE
  - DESCOVI, EMTRICITABINE
  - EPCLUSA, SOFOSBUVIR
  - GENVOYA, COBICISTAT
  - HARVONI, LEDIPASVIR
  - ODEFSEY, EMTRICITABINE
  - SOVALDI, SOFOSBUVIR
  - STRIBILD, COBICISTAT
  - TYBOST, COBICISTAT
  - VEMLIDY, TENOFOVIR ALAFENAMIDE FUMARATE
  - VIREAD, TENOFOVIR DISOPROXIL FUMARATE
  - VOSEVI, SOFOSBUVIR
  - ZYDELIG, IDELALISIB

**GLAND PHARMA LTD**

- \* GLAND PHARMA LTD
  - ADENOSINE, ADENOSINE
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
  - AZITHROMYCIN, AZITHROMYCIN
  - CARBOPLATIN, CARBOPLATIN
  - CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
  - CISPLATIN, CISPLATIN
  - DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
  - DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
  - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
  - ETOMIDATE, ETOMIDATE
  - FLUOROURACIL, FLUOROURACIL
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* G \*\*

- \* GLAND PHARMA LTD
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - HEPARIN SODIUM, HEPARIN SODIUM
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
  - LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM
  - MEROPENEM, MEROPENEM
  - MESNA, MESNA
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
  - MILRINONE LACTATE, MILRINONE LACTATE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXALIPLATIN, OXALIPLATIN
  - PACITAXEL, PACLITAXEL
  - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - TRANEXAMIC ACID, TRANEXAMIC ACID
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VECURONIUM BROMIDE, VECURONIUM BROMIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**GLASSHOUSE PHARMS**

- \* GLASSHOUSE PHARMACEUTICALS LTD CANADA
  - FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

**GLAXO GRP ENGLAND**

- \* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
  - INCURSE ELLIPTA, UMECLIDINIUM BROMIDE

**GLAXO GRP LTD**

- \* GLAXO GROUP LTD DBA GLAXOSMITHKLINE
  - FLOVENT HFA, FLUTICASONE PROPIONATE
- \* GLAXO GROUP LTD ENGLAND DBA GLAXOSMITHKLINE
  - ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
  - ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
  - ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
  - ADVAIR HFA, FLUTICASONE PROPIONATE
  - BREO ELLIPTA, FLUTICASONE FUROATE
  - FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
  - FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
  - FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
  - ZANTAC 150, RANITIDINE HYDROCHLORIDE
  - ZANTAC 300, RANITIDINE HYDROCHLORIDE
  - ZANTAC, RANITIDINE HYDROCHLORIDE

**GLAXOSMITHKLINE**

- \* GLAXOSMITHKLINE
  - ABREVA, DOCOSANOL (OTC)
  - AVODART, DUTASTERIDE
  - BACTROBAN, MUPIROCIN CALCIUM
  - BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
  - CEFTIN, CEFUROXIME AXETIL
  - EPIVIR-HBV, LAMIVUDINE
  - IMITREX STATDOSE, SUMATRIPTAN SUCCINATE
  - IMITREX, SUMATRIPTAN
  - IMITREX, SUMATRIPTAN SUCCINATE
  - JALYN, DUTASTERIDE
  - MALARONE PEDIATRIC, ATOVAQUONE
  - MALARONE, ATOVAQUONE
  - NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
  - NICORETTE, NICOTINE POLACRILEX (OTC)
  - RELENZA, ZANAMIVIR
  - VALTREX, VALACYCLOVIR HYDROCHLORIDE
  - WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
  - ZYBAN, BUPROPION HYDROCHLORIDE
- \* GLAXOSMITHKLINE CONSUMER HEALTHCARE HOLDINGS (US) LLC
  - LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
- \* GLAXOSMITHKLINE INTELLECTUAL PROPERTY DEVELOPMENT LTD ENGLAND
  - ANORO ELLIPTA, UMECLIDINIUM BROMIDE
  - ARNUNITY ELLIPTA, FLUTICASONE FUROATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT****\*\* G \*\***

- \* GLAXOSMITHKLINE INTELLECTUAL PROPERTY DEVELOPMENT LTD ENGLAND  
TRELEGY ELLIPTA, FLUTICASONE FUROATE
- \* GLAXOSMITHKLINE INTELLECTUAL PROPERTY LTD ENGLAND  
SEREVENT, SALMETEROL XINAFOATE  
VENTOLIN HFA, ALBUTEROL SULFATE

**GLAXOSMITHKLINE CON**

- \* GLAXOSMITHKLINE CONSUMER HEALTH  
TRANSDERM SCOP, SCOPOLAMINE

**GLAXOSMITHKLINE CONS**

- \* GLAXOSMITHKLINE CONSUMER HEALTHCARE  
ALLI, ORLISTAT (OTC)  
COMMIT, NICOTINE POLACRILEX (OTC)  
EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)  
FLONASE ALLERGY RELIEF, FLUTICASONE PROPIONATE (OTC)  
FLONASE SENSI-MIST ALLERGY RELIEF, FLUTICASONE FUROATE (OTC)  
NICORETTE, NICOTINE POLACRILEX (OTC)  
PREVACID 24 HR, LANSOPRAZOLE (OTC)  
TAVIST-1, CLEMASTINE FUMARATE (OTC)  
VOLTAREN, DICLOFENAC SODIUM

**GLAXOSMITHKLINE LLC**

- \* GLAXOSMITHKLINE LLC  
AMERGE, NARatriptan Hydrochloride  
DYAZIDE, HYDROCHLOROTHIAZIDE  
FLOLAN, EPOPROSTENOL SODIUM  
LAMICTAL CD, LAMOTRIGINE  
LAMICTAL ODT, LAMOTRIGINE  
LAMICTAL XR, LAMOTRIGINE  
LAMICTAL, LAMOTRIGINE  
MEPRON, ATOVAQUONE  
REQUIP XL, ROPINIROLE HYDROCHLORIDE  
REQUIP, ROPINIROLE HYDROCHLORIDE  
RYTHMOL SR, PROPAFENONE HYDROCHLORIDE  
RYTHMOL, PROPAFENONE HYDROCHLORIDE

**GLENMARK GENERICS**

- \* GLENMARK GENERICS INC USA  
ADAPALENE, ADAPALENE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
IMIQUIMOD, IMIQUIMOD  
MOMETASONE FUROATE, MOMETASONE FUROATE  
NIZATIDINE, NIZATIDINE  
ZONISAMIDE, ZONISAMIDE
- \* GLENMARK GENERICS LIMITED  
BRIELLYN, ETHINYL ESTRADIOL  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
- \* GLENMARK GENERICS LTD  
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM  
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE  
ALYACEN 1/35, ETHINYL ESTRADIOL  
ALYACEN 7/7/7, ETHINYL ESTRADIOL  
ASHLYNA, ETHINYL ESTRADIOL  
ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE  
CARVEDILOL, CARVEDILOL  
CICLOPIROX, CICLOPIROX  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
DESOXIMETASONE, DESOXIMETASONE  
ESZOPICLONE, ESZOPICLONE  
FELODIPINE, FELODIPINE  
FLUCONAZOLE, FLUCONAZOLE  
FLUOCINONIDE, FLUOCINONIDE  
HEATHER, NORETHINDRONE  
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE  
LAMOTRIGINE, LAMOTRIGINE  
LEVOFLOXACIN, LEVOFLOXACIN  
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* G \*\*

- \* GLENMARK GENERICS LTD
  - LITHIUM CARBONATE, LITHIUM CARBONATE
  - MARLISSA, ETHINYL ESTRADIOL
  - MELOXICAM, MELOXICAM
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
  - MOMETASONE FUROATE, MOMETASONE FUROATE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - NAPROXEN, NAPROXEN
  - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - NORETHINDRONE, NORETHINDRONE
  - NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - OMEPRAZOLE, OMEPRAZOLE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON, ONDANSETRON
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - THEOPHYLLINE, THEOPHYLLINE
  - TOPIRAMATE, TOPIRAMATE
  - TRANDOLAPRIL AND VERAPAMIL HYDROCHLORIDE, TRANDOLAPRIL
  - TROSPiUM CHLORIDE, TROSPiUM CHLORIDE
  - URSODIOL, URSODIOL
  - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
  - VIORELE, DESOGESTREL
  - ZOLMITRIPTAN, ZOLMITRIPTAN
- \* GLENMARK GENERICS LTD INDIA
  - INDOMETHACIN, INDOMETHACIN
  - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE

**GLENMARK PHARMS**

- \* GLENMARK PHARMACEUTICALS INC USA
  - CICLOPIROX, CICLOPIROX
  - CLOTRIMAZOLE, CLOTRIMAZOLE
  - MUPIROCIN, MUPIROCIN
- \* GLENMARK PHARMACEUTICALS LTD
  - MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
- \* GLENMARK PHARMACEUTICALS SA
  - CALCIPOTRIENE, CALCIPOTRIENE
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
  - DESONIDE, DESONIDE
  - LINEZOLID, LINEZOLID
  - NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**GLENMARK PHARMS INC**

- \* GLENMARK PHARMACEUTICALS INC USA
  - CALCIPOTRIENE, CALCIPOTRIENE
  - LITHIUM CARBONATE, LITHIUM CARBONATE
  - MUPIROCIN, MUPIROCIN CALCIUM
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

**GLENMARK PHARMS LTD**

- \* GLENMARK PHARMACEUTICALS LTD
  - ACYCLOVIR, ACYCLOVIR
  - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
  - APREPITANT, APREPITANT
  - ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
  - BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
  - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
  - DESONIDE, DESONIDE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* G \*\*

- \* GLENMARK PHARMACEUTICALS LTD
  - EZETIMIBE, EZETIMIBE
  - FENOFIBRATE (MICRONIZED), FENOFIBRATE
  - FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
  - GABAPENTIN, GABAPENTIN
  - HAILEY FE 1/20, ETHINYLMESTRADIOL
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - INDOMETHACIN, INDOMETHACIN
  - LAMOTRIGINE, LAMOTRIGINE
  - LEVONORGESTREL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL
  - LEVONORGESTREL, LEVONORGESTREL (OTC)
  - LIDOCAINE, LIDOCAINE
  - NAPROXEN SODIUM, NAPROXEN SODIUM
  - NITROGLYCERIN, NITROGLYCERIN
  - NORETHINDRONE ACETATE AND ETHINYLMESTRADIOL AND FERROUS FUMARATE, ETHINYLMESTRADIOL
  - NORETHINDRONE ACETATE AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - OXCARBAZEPINE, OXCARBAZEPINE
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
  - PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
  - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
  - RILUZOLE, RILUZOLE
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - RUFINAMIDE, RUFINAMIDE
  - TELMISARTAN, TELMISARTAN
  - TRETINOIN, TRETINOIN
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
  - VORICONAZOLE, VORICONAZOLE

**GLOBAL ISOTOPES LLC**

- \* GLOBAL ISOTOPES LLC DBA ZEVACOR MOLECULAR
  - AMMONIA N 13, AMMONIA N-13
  - CHOLINE C-11, CHOLINE C-11
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**GRANULES INDIA**

- \* GRANULES INDIA LTD
  - IBUPROFEN, IBUPROFEN (OTC)
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

**GRANULES INDIA LTD**

- \* GRANULES INDIA LTD
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - IBUPROFEN, IBUPROFEN
  - IBUPROFEN, IBUPROFEN (OTC)

**GUARDIAN PHARMS**

- \* GUARDIAN PHARMACEUTICALS
  - GUAIFENESIN, GUAIFENESIN (OTC)

**GUERBET**

- \* GUERBET LLC
  - DOTAREM, GADOTERATE MEGLUMINE
  - LPIOIDOL, ETHIODIZED OIL

**HANFORD GC**

- \* GC HANFORD MANUFACTURING CO
  - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
  - AMPICILLIN SODIUM, AMPICILLIN SODIUM
  - PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM

**POHL BOSKAMP**

- \* G POHL BOSKAMP GMBH AND CO KG
  - GONITRO, NITROGLYCERIN

\*\* H \*\*

**HAEMONETICS**

- \* HAEMONETICS CORP
  - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

**HALOCARBON PRODS**

- \* HALOCARBON PRODUCTS CORP  
ISOFLURANE, ISOFLURANE  
SEVOFLURANE, SEVOFLURANE

**HALOZYME THERAP**

- \* HALOZYME THERAPEUTICS INC  
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN

**HAMELN PHARMA PLUS**

- \* HAMELN PHARMA PLUS GMBH  
PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM  
PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM

**HAMELN RDS GMBH**

- \* HAMELN RDS GMBH  
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE

**HANDA PHARMS LLC**

- \* HANDA PHARMACEUTICALS LLC  
LAMOTRIGINE, LAMOTRIGINE

**HANSAMED INC**

- \* HANSAMED INC  
ULTACAN FORTE, ARTICAINE HYDROCHLORIDE  
ULTACAN, ARTICAINE HYDROCHLORIDE

**HARRIS PHARM**

- \* HARRIS PHARMACEUTICAL INC  
FLUCONAZOLE, FLUCONAZOLE  
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE

**HAUPT PHARMA**

- \* HAUPT PHARMA INC  
DUTASTERIDE, DUTASTERIDE

**HEC PHARM USA INC**

- \* HEC PHARM USA INC  
CLARITHROMYCIN, CLARITHROMYCIN

**HELSINN HLTHCARE**

- \* HELSINN HEALTHCARE SA  
AKYNZEO, NETUPITANT  
ALOXI, PALONOSETRON HYDROCHLORIDE

**HERCON PHARM**

- \* HERCON PHARMACEUTICAL LLC  
NITROGLYCERIN, NITROGLYCERIN

**HERITAGE LIFE**

- \* HERITAGE LIFE SCIENCES BARBADOS INC  
CLOZARIL, CLOZAPINE

**HERITAGE PHARMA**

- \* HERITAGE PHARMA LABS INC  
ACETAZOLAMIDE, ACETAZOLAMIDE  
ACHROMYCIN V, TETRACYCLINE HYDROCHLORIDE  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE  
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE  
DIFLUNISAL, DIFLUNISAL  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE  
LITHIUM CARBONATE, LITHIUM CARBONATE  
METHIMAZOLE, METHIMAZOLE  
NIFEDIPINE, NIFEDIPINE  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

**HERITAGE PHARMS INC**

- \* HERITAGE PHARMACEUTICALS INC  
ACETAZOLAMIDE, ACETAZOLAMIDE  
ACYCLOVIR, ACYCLOVIR  
ALPRAZOLAM, ALPRAZOLAM  
CALCIUM ACETATE, CALCIUM ACETATE  
CARISOPRODOL AND ASPIRIN, ASPIRIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HERITAGE PHARMACEUTICALS INC
  - DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
  - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
  - DOXYCYCLINE, DOXYCYCLINE
  - ETHOSUXIMIDE, ETHOSUXIMIDE
  - FELODIPINE, FELODIPINE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
  - GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
  - GLYBURIDE, GLYBURIDE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - INDOMETHACIN, INDOMETHACIN
  - KETOPROFEN, KETOPROFEN
  - LEFLUNOMIDE, LEFLUNOMIDE
  - MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
  - NIMODIPINE, NIMODIPINE
  - NYSTATIN, NYSTATIN
  - PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
  - TROSPiUM CHLORIDE, TROSPiUM CHLORIDE
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**HERON THERAPS INC**

- \* HERON THERAPEUTICS INC
  - CINVANTI, APREPITANT
  - SUSTOL, GRANisetron

**HETERO LABS LTD III**

- \* HETERO LABS LTD UNIT III
  - ABACAVIR SULFATE, ABACAVIR SULFATE
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - FENOFIBRATE, FENOFIBRATE
  - FINASTERIDE, FINASTERIDE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - INDOMETHACIN, INDOMETHACIN
  - LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - LITHIUM CARBONATE, LITHIUM CARBONATE
  - METHOCARBAMOL, METHOCARBAMOL
  - NEVIRAPINE, NEVIRAPINE
  - SIMVASTATIN, SIMVASTATIN
  - STAVUDINE, STAVUDINE
  - TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
  - TORSEMIDE, TORSEMIDE
  - ZIDOVUDINE, ZIDOVUDINE

**HETERO LABS LTD V**

- \* HETERO LABS LTD UNIT V
  - ACYCLOVIR, ACYCLOVIR
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - ENTECAVIR, ENTECAVIR
  - FAMCICLOVIR, FAMCICLOVIR
  - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - IRBESARTAN, IRBESARTAN
  - LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
  - LAMIVUDINE, LAMIVUDINE
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LINEZOLID, LINEZOLID
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HETERO LABS LTD UNIT V  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM  
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM  
SILDENAFIL CITRATE, SILDENAFIL CITRATE  
TELMISARTAN, TELMISARTAN  
TETRABENAZINE, TETRABENAZINE  
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE  
VALGANCICLOVIR HYDROCHLORIDE, VALGANCICLOVIR HYDROCHLORIDE  
VALSARTAN, VALSARTAN

**HEYL CHEMISCH**

- \* HEYL CHEMISCH PHARMAZEUTISCHE FABRIK  
RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

**HI TECH PHARMA**

- \* HI TECH PHARMACAL CO INC  
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN  
ACYCLOVIR, ACYCLOVIR  
ALBUTEROL SULFATE, ALBUTEROL SULFATE  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
CALCIPOTRIENE, CALCIPOTRIENE  
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE  
CICLOPIROX, CICLOPIROX  
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE  
CORMAX, CLOBETASOL PROPIONATE  
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE  
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE  
EMBELINE E, CLOBETASOL PROPIONATE  
EMBELINE, CLOBETASOL PROPIONATE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
GABAPENTIN, GABAPENTIN  
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
LACTULOSE, LACTULOSE  
LEVOCARNITINE, LEVOCARNITINE  
LEVOFLOXACIN, LEVOFLOXACIN  
LIDOCAINE AND PRILOCAINE, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)  
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE  
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)  
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)  
NYSTATIN, NYSTATIN  
OFLOXACIN, OFLOXACIN  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE  
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE  
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE  
TIMOLOL MALEATE, TIMOLOL MALEATE  
VOSOL HC, ACETIC ACID, GLACIAL  
VOSOL, ACETIC ACID, GLACIAL

**HI TECH PHARMA CO**

- \* HI TECH PHARMACAL CO INC  
FLUNISOLIDE, FLUNISOLIDE  
PREDNISOLONE, PREDNISOLONE

**HI-TECH PHARMA CO**

- \* HI-TECH PHARMACAL CO INC  
FAMOTIDINE, FAMOTIDINE  
GATIFLOXACIN, GATIFLOXACIN  
LORAZEPAM, LORAZEPAM  
PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE

**HI-TECH PHARMACAL**

- \* HI-TECH PHARMACAL CO INC  
BROMFENAC SODIUM, BROMFENAC SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HI-TECH PHARMACAL CO INC  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
DESONIDE, DESONIDE  
IBUPROFEN, IBUPROFEN  
LEVETIRACETAM, LEVETIRACETAM  
MEGESTROL ACETATE, MEGESTROL ACETATE  
MORPHINE SULFATE, MORPHINE SULFATE  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE  
TRIFLURIDINE, TRIFLURIDINE

**HIGH TECH PHARMA**

- \* HIGH TECHNOLOGY PHARMACAL CO INC  
VALPROIC ACID, VALPROIC ACID

**HIKMA**

- \* HIKMA FARMACEUTICA LDA  
CEFOTAXIME, CEFOTAXIME SODIUM
- \* HIKMA PHARMACEUTICALS  
AMOXICILLIN, AMOXICILLIN  
CEFACLOR, CEFACLOR  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CEPHALEXIN, CEPHALEXIN  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
GLYBURIDE (MICRONIZED), GLYBURIDE

**HIKMA FARMACEUTICA**

- \* HIKMA FARMACEUTICA (PORTUGAL) SA  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE  
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE  
CEFOXITIN, CEFOXITIN SODIUM  
CEFTRIAXONE, CEFTRIAXONE SODIUM  
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN  
CIPROFLOXACIN, CIPROFLOXACIN  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
ENALAPRILAT, ENALAPRILAT  
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUMAZENIL, FLUMAZENIL  
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM  
GLYCOPYRROLATE, GLYCOPYRROLATE  
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
LEVETIRACETAM, LEVETIRACETAM  
LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN  
METOPROLOL TARTRATE, METOPROLOL TARTRATE  
MILRINONE LACTATE IN PLASTIC CONTAINER, MILRINONE LACTATE  
MILRINONE LACTATE, MILRINONE LACTATE  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
PROGESTERONE, PROGESTERONE  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE  
TERBUTALINE SULFATE, TERBUTALINE SULFATE  
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE  
VALPROATE SODIUM, VALPROATE SODIUM
- \* HIKMA FARMACEUTICA PORTUGAL LDA  
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM  
CEFUXIME SODIUM, CEFUXIME SODIUM  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
- \* HIKMA FARMACEUTICA PORTUGAL SA  
CEFOTETAN, CEFOTETAN DISODIUM  
CEFTRIAXONE SODIUM, CEFTRIAXONE SODIUM  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
ETOMIDATE, ETOMIDATE  
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE  
OXYTOCIN, OXYTOCIN  
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HIKMA FARMACEUTICA PORTUGAL SA  
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
- \* HIKMA FARMACEUTICA SA  
ZOLEDRONIC ACID, ZOLEDRONIC ACID

**HIKMA INTL PHARMS**

- \* HIKMA INTERNATIONAL PHARMACEUTICALS LLC  
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN  
CAPTOPRIL, CAPTOPRIL  
CARISOPRODOL, CARISOPRODOL  
CORTISONE ACETATE, CORTISONE ACETATE  
DIGOXIN, DIGOXIN  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
HYDROCORTISONE, HYDROCORTISONE  
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE  
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LISINOPRIL, LISINOPRIL  
METHOCARBAMOL, METHOCARBAMOL  
MITIGARE, COLCHICINE  
PRIMIDONE, PRIMIDONE

**HIKMA PHARM CO LTD**

- \* HIKMA PHARM CO LTD  
ARGATROBAN, ARGATROBAN

**HIKMA PHARMS**

- \* HIKMA PHARMACEUTICALS  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN  
AMOXICILLIN, AMOXICILLIN  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE  
GEMFIBROZIL, GEMFIBROZIL  
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE  
LETROZOLE, LETROZOLE  
MODAFINIL, MODAFINIL  
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM  
RIFAMPIN, RIFAMPIN
- \* HIKMA PHARMACEUTICALS CO LTD  
PARICALCITOL, PARICALCITOL
- \* HIKMA PHARMACEUTICALS LLC  
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM  
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE  
DANTROLENE SODIUM, DANTROLENE SODIUM  
DOXERCALCIFEROL, DOXERCALCIFEROL  
FOLIC ACID, FOLIC ACID  
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE  
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE  
OLANZAPINE, OLANZAPINE  
PIROXICAM, PIROXICAM  
PREDNISONE, PREDNISONE  
ZALEPLON, ZALEPLON

**HILL DERMAC**

- \* HILL DERMACEUTICALS INC  
DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE  
DERMOTIC, FLUOCINOLONE ACETONIDE

**HILL DERMACEUTICALS**

- \* HILL DERMACEUTICALS INC  
TOLAK, FLUOROURACIL

**HISAMITSU PHARM CO**

- \* HISAMITSU PHARMACEUTICAL CO INC  
SALONPAS, MENTHOL (OTC)

**HISUN PHARM HANGZHOU**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HISUN PHARMACEUTICAL (HANGZHOU) CO LTD  
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
- \* HISUN PHARMACEUTICAL HANGZHOU CO LTD  
IRBESARTAN AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE  
IRBESARTAN, IRBESARTAN  
SIMVASTATIN, SIMVASTATIN

**HOFFMANN LA ROCHE**

- \* HOFFMANN LA ROCHE INC  
BONIVA, IBANDRONATE SODIUM  
INVIRASE, SAQUINAVIR MESYLATE  
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE  
XELODA, CAPECITABINE  
ZELBORAF, VEMURAFENIB

**HOFFMANN-LA ROCHE**

- \* HOFFMANN-LA ROCHE INC  
ALECensa, ALECTINIB HYDROCHLORIDE  
INVIRASE, SAQUINAVIR MESYLATE

**HONG KONG**

- \* HONG KONG KING-FRIEND INDUSTRIAL CO LTD  
CYTARABINE, CYTARABINE  
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE

**HOPE PHARMS**

- \* HOPE PHARMACEUTICALS  
NITHIODOTE, SODIUM NITRITE  
SODIUM NITRITE, SODIUM NITRITE  
SODIUM THIOSULFATE, SODIUM THIOSULFATE

**HORIZON PHARMA**

- \* HORIZON PHARMA INC  
DUEXIS, FAMOTIDINE  
RAYOS, PREDNISONE
- \* HORIZON PHARMA IRELAND LTD  
PENNSAID, DICLOFENAC SODIUM
- \* HORIZON PHARMA RHEUMATOLOGY LLC  
MIGERGOT, CAFFEINE

**HORIZON PHARMA INC**

- \* HORIZON PHARMA INC  
BUPHENYL, SODIUM PHENYLBUTYRATE

**HORIZON PHARMA USA**

- \* HORIZON PHARMA USA INC  
PROSYSBI, CYSTEAMINE BITARTRATE  
VIMOVO, ESOMEPRAZOLE MAGNESIUM

**HORIZON THERAPS INC**

- \* HORIZON THERAPEUTICS INC  
RAVICTI, GLYCEROL PHENYLBUTYRATE

**HOSPIRA**

- \* HOSPIRA INC  
A-METHAPRED, METHYLSPREDNISOLONE SODIUM SUCCINATE  
ACETYL CYSTEINE, ACETYL CYSTEINE  
ALFENTANIL, ALFENTANIL HYDROCHLORIDE  
AMIDATE, ETOMIDATE  
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID  
AMINOPHYLLINE, AMINOPHYLLINE  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE  
AQUASOL A, VITAMIN A PALMITATE  
ARTICAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, ARTICAINE HYDROCHLORIDE  
ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE  
ATROPINE SULFATE LIFESHIELD ABBOJECT SYRINGE, ATROPINE SULFATE  
AZITHROMYCIN, AZITHROMYCIN  
BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE  
BUMETANIDE, BUMETANIDE  
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

## \* HOSPIRA INC

BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE  
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE  
CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE  
CARBOPLATIN, CARBOPLATIN  
CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE  
CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE  
CIPROFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN  
CIPROFLOXACIN, CIPROFLOXACIN  
CORLOPAM, FENOLDOPAM MESYLATE  
CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE  
CYTARABINE, CYTARABINE  
DACARBAZINE, DACARBAZINE  
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE  
DEMEROL, MEPERIDINE HYDROCHLORIDE  
DEXTROSE 25%, DEXTROSE  
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 50%, DEXTROSE  
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE  
DIAZEPAM, DIAZEPAM  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE  
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE  
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE  
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE  
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE  
DROPERIDOL, DROPERIDOL  
ENALAPRILAT, ENALAPRILAT  
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE  
ERYTHROGIN, ERYTHROMYCIN LACTOBIONATE  
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE  
FENTANYL CITRATE, FENTANYL CITRATE  
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE  
FUROSEMIDE, FUROSEMIDE  
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN  
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN  
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN  
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM  
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN  
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM  
HEPARIN SODIUM, HEPARIN SODIUM  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
ISOFLURANE, ISOFLURANE  
ISUPREL, ISOPROTERENOL HYDROCHLORIDE  
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
LEVOPHED, NOREPINEPHRINE BITARTRATE  
LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE  
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE  
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
LORAZEPAM, LORAZEPAM  
LTA II KIT, LIDOCAINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

## \* HOSPIRA INC

M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID  
M.V.I. ADULT, ASCORBIC ACID  
M.V.I. PEDIATRIC, ASCORBIC ACID  
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE  
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE  
MAGNESIUM SULFATE, MAGNESIUM SULFATE  
MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE  
MANNITOL 25%, MANNITOL  
MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE  
MARCAINE, BUPIVACAINE HYDROCHLORIDE  
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM  
METHOTREXATE SODIUM, METHOTREXATE SODIUM  
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
METOPROLOL TARTRATE, METOPROLOL TARTRATE  
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE  
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE  
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE  
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE  
MORPHINE SULFATE, MORPHINE SULFATE  
NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE  
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE  
NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
PACLITAXEL, PACLITAXEL  
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM  
PANCURONIUM BROMIDE, PANCURONIUM BROMIDE  
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
POTASSIUM ACETATE, POTASSIUM ACETATE  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
PRECEDEX, DEXMEDETOMIDINE HYDROCHLORIDE  
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE  
PROPOFOL, PROPOFOL  
QUELICIN PRESERVATIVE FREE, SUCCINYLCHOLINE CHLORIDE  
QUELICIN, SUCCINYLCHOLINE CHLORIDE  
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE  
ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE  
SODIUM ACETATE, SODIUM ACETATE  
SODIUM BICARBONATE, SODIUM BICARBONATE  
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE  
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE  
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
SUFENTANIL CITRATE, SUFENTANIL CITRATE  
TALWIN, PENTAZOCINE LACTATE  
TAZICEF, CEFTAZIDIME  
THAM, TROMETHAMINE  
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE  
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE  
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE  
VECURONIUM BROMIDE, VECURONIUM BROMIDE  
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE  
VINORELBINE TARTRATE, VINORELBINE TARTRATE  
VITAMIN K1, PHYTONADIONE  
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE

## \* HOSPIRA WORLDWIDE, INC

DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE  
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HOSPIRA WORLDWIDE, INC
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
  - NITROPRESS, SODIUM NITROPRUSSIDE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE

**HOSPIRA INC**

- \* HOSPIRA INC
  - ADENOSINE, ADENOSINE
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
  - AMPICILLIN SODIUM, AMPICILLIN SODIUM
  - ARGATROBAN, ARGATROBAN
  - ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
  - ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
  - BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE
  - BIVALIRUDIN, BIVALIRUDIN
  - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
  - CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE
  - CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
  - CEFOXITIN, CEFOXITIN SODIUM
  - CEFTRIAXONE, CEFTRIAXONE SODIUM
  - CEFUROXIME SODIUM, CEFUROXIME SODIUM
  - DAPTOMYCIN, DAPTOMYCIN
  - DOCETAXEL, DOCETAXEL
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - HEPARIN SODIUM, HEPARIN SODIUM
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - IMIPENEM AND CILASTATIN, CILASTATIN SODIUM
  - INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
  - LINEZOLID IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, LINEZOLID
  - LINEZOLID, LINEZOLID
  - MAGNESIUM SULFATE, MAGNESIUM SULFATE
  - MAXIPIME, CEFEPIME HYDROCHLORIDE
  - MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
  - MEROPENEM, MEROPENEM
  - MILRINONE LACTATE, MILRINONE LACTATE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - NIPENT, PENTOSTATIN
  - OXACILLIN SODIUM, OXACILLIN SODIUM
  - OXALIPLATIN, OXALIPLATIN
  - PARICALCITOL, PARICALCITOL
  - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
  - SODIUM BICARBONATE, SODIUM BICARBONATE
  - TACROLIMUS, TACROLIMUS
  - THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - VANCOMYCYIN HYDROCHLORIDE, VANCOMYCYIN HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

**HOSPIRA WORLDWIDE**

- \* HOSPIRA WORLDWIDE PTY
  - OXALIPLATIN, OXALIPLATIN

**HOT SHOTS NM LLC**

- \* HOT SHOTS NUCLEAR MEDICINE LLC
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**HOUSTON CYCLOTRON**

- \* HOUSTON CYCLOTRON PARTNERS LP
  - AMMONIA N 13, AMMONIA N-13
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**HQ SPCLT PHARMA**

- \* HQ SPECIALTY PHARMA CORP

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* H \*\*

- \* HQ SPECIALTY PHARMA CORP
  - CISPLATIN, CISPLATIN
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
  - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
  - LINEZOLID, LINEZOLID
  - MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
  - MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
  - TAXOL, PACLITAXEL

**HQ SPECIALTY PHARMA**

- \* HQ SPECIALTY PHARMA LLC
  - LEVETIRACETAM IN SODIUM CHLORIDE, LEVETIRACETAM

**HRA PHARMA**

- \* HRA PHARMA LLC
  - METOPIRONE, METYRAPONE

**HUMANWELL PURACAP**

- \* HUMANWELL PURACAP PHARMACEUTICAL WUHAN CO LTD
  - DUTASTERIDE, DUTASTERIDE
  - IBUPROFEN, IBUPROFEN (OTC)

**ROCHE**

- \* HOFFMANN LA ROCHE INC
  - BONIVA, IBANDRONATE SODIUM
  - COPEGUS, RIBAVIRIN
  - FUZEON, ENFUVIRTIDE
  - KLONOPIN, CLONAZEPAM
  - TAMIFLU, OSELTAMIVIR PHOSPHATE
  - VALIUM, DIAZEPAM

\*\* I \*\*

**IBA MOLECULAR N AM**

- \* IBA MOLECULAR NORTH AMERICA INC
  - AMMONIA N 13, AMMONIA N-13

**ICU MEDICAL INC**

- \* ICU MEDICAL INC
  - ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
  - AMINOSYN 10% (PH6), AMINO ACIDS
  - AMINOSYN 10%, AMINO ACIDS
  - AMINOSYN 3.5% M, AMINO ACIDS
  - AMINOSYN 3.5%, AMINO ACIDS
  - AMINOSYN 5%, AMINO ACIDS
  - AMINOSYN 7% (PH6), AMINO ACIDS
  - AMINOSYN 7% W/ ELECTROLYTES, AMINO ACIDS
  - AMINOSYN 7%, AMINO ACIDS
  - AMINOSYN 8.5% (PH6), AMINO ACIDS
  - AMINOSYN 8.5% W/ ELECTROLYTES, AMINO ACIDS
  - AMINOSYN 8.5%, AMINO ACIDS
  - AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
  - AMINOSYN II 10% W/ ELECTROLYTES, AMINO ACIDS
  - AMINOSYN II 10%, AMINO ACIDS
  - AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
  - AMINOSYN II 7%, AMINO ACIDS
  - AMINOSYN II 8.5% W/ ELECTROLYTES, AMINO ACIDS
  - AMINOSYN II 8.5%, AMINO ACIDS
  - AMINOSYN-HBC 7%, AMINO ACIDS
  - AMINOSYN-HF 8%, AMINO ACIDS
  - AMINOSYN-PF 10%, AMINO ACIDS
  - AMINOSYN-PF 7%, AMINO ACIDS
  - AMINOSYN-RF 5.2%, AMINO ACIDS
  - DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
  - DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
  - DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

## \* ICU MEDICAL INC

DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE  
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE  
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE  
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL  
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL  
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE  
MORPHINE SULFATE, MORPHINE SULFATE  
NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE  
POTASSIUM CHLORIDE 0.149% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, POTASSIUM  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE  
POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE  
POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,  
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,  
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE  
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL  
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
STERILE WATER IN PLASTIC CONTAINER, STERILE WATER FOR IRRIGATION

**IDENTI PHARMS INC**

\* IDENTI PHARMACEUTICALS INC

FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

**IDT AUSTRALIA LTD**

- \* IDT AUSTRALIA LTD
- PINDOLOL, PINDOLOL
- TEMOZOLOMIDE, TEMOZOLOMIDE

**IMPAK LABS**

- \* IMPAX LABORATORIES INC
- ACARBOSE, ACARBOSE
- ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
- BACLOFEN, BACLOFEN
- BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
- CARBIDOPA AND LEVODOPA, CARBIDOPA
- CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
- COLESTIPOL HYDROCHLORIDE, COLESTIPOL HYDROCHLORIDE
- DANTROLENE SODIUM, DANTROLENE SODIUM
- DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
- DIGOXIN, DIGOXIN
- DIPYRIDAMOLE, DIPYRIDAMOLE
- DIVALPROEX SODIUM, DIVALPROEX SODIUM
- FENOFIBRATE (MICRONIZED), FENOFIBRATE
- FENOFIBRATE, FENOFIBRATE
- FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
- METHYLTESTOSTERONE, METHYLTESTOSTERONE
- MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
- NADOLOL AND BENDROFLUMETHIAZIDE, BENDROFLUMETHIAZIDE
- OMEPRAZOLE, OMEPRAZOLE
- OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
- PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
- PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
- PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
- RILUZOLE, RILUZOLE
- RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
- TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
- TERBUTALINE SULFATE, TERBUTALINE SULFATE

**IMPAK LABS INC**

- \* IMPAX LABORATORIES INC
- ACITRETIN, ACITRETIN
- ADRENAClick, EPINEPHRINE
- ALBENZA, ALBENDAZOLE
- ALENDRONATE SODIUM, ALENDRONATE SODIUM
- ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
- BUDESONIDE, BUDESONIDE
- BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
- DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
- DEXEDRINE, DEXTROAMPHETAMINE SULFATE
- DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
- DOXYCYCLINE, DOXYCYCLINE
- EMVERM, MEBENDAZOLE
- EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
- EZETIMIBE AND SIMVASTATIN, EZETIMIBE
- FELBAMATE, FELBAMATE
- FENOFIBRIC ACID, CHOLINE FENOFIBRATE
- GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
- GLYBURIDE, GLYBURIDE
- HYDROCORTISONE, HYDROCORTISONE
- HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
- LAMOTRIGINE, LAMOTRIGINE
- LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
- METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
- METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
- METHYLTESTOSTERONE, METHYLTESTOSTERONE
- METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
- MIRTAZAPINE, MIRTAZAPINE
- MORPHINE SULFATE, MORPHINE SULFATE
- NABUMETONE, NABUMETONE
- NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

- \* IMPAX LABORATORIES INC  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE  
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE  
RYTARY, CARBIDOPA  
SEVELAMER CARBONATE, SEVELAMER CARBONATE  
URSODIOL, URSODIOL

**IMPAK PHARMS**

- \* IMPAX PHARMACEUTICALS  
GEMFIBROZIL, GEMFIBROZIL  
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE  
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

**INCYTE CORP**

- \* INCYTE CORP  
JAKAFI, RUXOLITINIB PHOSPHATE

**INDICUS PHARMA**

- \* INDICUS PHARMA LLC  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
LETROZOLE, LETROZOLE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**INDIVIOR INC**

- \* INDIVIOR INC  
BUPRENEX, BUPRENORPHINE HYDROCHLORIDE  
SUBLOCADE, BUPRENORPHINE  
SUBOXONE, BUPRENORPHINE HYDROCHLORIDE

**INDOCO REMEDIES**

- \* INDOCO REMEDIES LTD  
ALLOPURINOL, ALLOPURINOL  
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE  
GLIMEPIRIDE, GLIMEPIRIDE

**INGENUS PHARMS LLC**

- \* INGENUS PHARMACEUTICALS LLC  
CARBOPLATIN, CARBOPLATIN  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE  
DOCETAXEL, DOCETAXEL  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

**INGENUS PHARMS NJ**

- \* INGENUS PHARMACEUTICALS NJ LLC  
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN  
CARISOPRODOL, CARISOPRODOL  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE  
PRIMAQUINE PHOSPHATE, PRIMAQUINE PHOSPHATE  
PROBENECID AND COLCHICINE, COLCHICINE

**INJECTALIA**

- \* INJECTALIA SRL  
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

**INNOGENIX**

- \* INNOGENIX LLC  
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
METRONIDAZOLE, METRONIDAZOLE  
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE

**INNOPHARMA LICENSING**

- \* INNOPHARMA LICENSING LLC  
BENDAMUSTINE HYDROCHLORIDE, BENDAMUSTINE HYDROCHLORIDE

**INST BIOCHEM**

- \* INSTITUT BIOCHEMIQUE SA  
FLECTOR, DICLOFENAC EPOLAMINE

**INSTITUT BIOCHIMIQUE**

- \* INSTITUT BIOCHIMIQUE SA (IBSA)  
TIROSINT, LEVOTHYROXINE SODIUM

**INSYS DEV CO INC**

- \* INSYS DEVELOPMENT CO INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

- \* INSYS DEVELOPMENT CO INC  
SUBSYS, FENTANYL  
SYNDROS, DRONABINOL

**INTAS PHARMS USA**

- \* INTAS PHARMACEUTICALS USA  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE

**INTELLIPHARMACEUTICS**

- \* INTELLIPHARMACEUTICS CORP  
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE  
LEVETIRACETAM, LEVETIRACETAM  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

**INTERCEPT PHARMS INC**

- \* INTERCEPT PHARMACEUTICALS INC  
OCALIVA, OBETICHOLIC ACID

**INTERGEL PHARM**

- \* INTERGEL PHARMACEUTICAL INC  
NIFEDIPINE, NIFEDIPINE

**INTERGEL PHARMS INC**

- \* INTERGEL PHARMACEUTICALS INC  
DUTASTERIDE, DUTASTERIDE

**INTERPHARMA PRAHA AS**

- \* INTERPHARMA PRAHA AS  
ORALTAG, IOHEXOL

**INTERSECT ENT INC**

- \* INTERSECT ENT INC  
SINUVA, MOMETASONE FUROATE

**INTL MEDICATED**

- \* INTERNATIONAL MEDICATED SYSTEMS LTD  
MILRINONE LACTATE, MILRINONE LACTATE

**INTL MEDICATION**

- \* INTERNATIONAL MEDICATION SYSTEM  
LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
MANNITOL 25%, MANNITOL  
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE  
PHYTONADIONE, PHYTONADIONE  
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
- \* INTERNATIONAL MEDICATION SYSTEMS LTD  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

**INTL MEDICATION SYS**

- \* INTERNATIONAL MEDICATION SYSTEMS LTD  
LORAZEPAM, LORAZEPAM  
SODIUM BICARBONATE, SODIUM BICARBONATE

**INVAGEN PHARMS**

- \* INVAGEN PHARMACEUTICALS INC  
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE  
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
CALCIUM ACETATE, CALCIUM ACETATE  
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
FENOFIBRATE (MICRONIZED), FENOFIBRATE  
FOLIC ACID, FOLIC ACID  
FOSINOPRIL SODIUM AND HYDROCHLORTHIAZIDE, FOSINOPRIL SODIUM  
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM  
GABAPENTIN, GABAPENTIN  
GEMFIBROZIL, GEMFIBROZIL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

- \* INVAGEN PHARMACEUTICALS INC
  - GLIMEPIRIDE, GLIMEPIRIDE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - LEVETIRACETAM, LEVETIRACETAM
  - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LISINOPRIL, LISINOPRIL
  - MEPROBAMATE, MEPROBAMATE
  - NABUMETONE, NABUMETONE
  - NADOLOL, NADOLOL
  - NAPROXEN, NAPROXEN
  - OLANZAPINE, OLANZAPINE
  - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
  - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
  - RAMIPRIL, RAMIPRIL
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SEVELAMER CARBONATE, SEVELAMER CARBONATE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - TOPIRAMATE, TOPIRAMATE
  - TROSPiUM CHLORIDE, TROSPiUM CHLORIDE
  - WARFARIN SODIUM, WARFARIN SODIUM
  - ZOLMITriPTAN, ZOLMITriPTAN
  - ZOLPiDEM TARTRATE, ZOLPiDEM TARTRATE
  - ZONiSAMIDE, ZONiSAMIDE

**INVENTIA HLTHCARE**

- \* INVENTIA HEALTHCARE PRIVATE LTD
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - ILOPERIDONE, ILOPERIDONE
  - LANSOPRAZOLE, LANSOPRAZOLE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - TELMiSARTAN, TELMiSARTAN

**IPCA LABS LTD**

- \* IPCA LABORATORIES LTD
  - ALLOPURiNOL, ALLOPURiNOL
  - ATENOLOL, ATENOLOL
  - CETiRiZiNE HYDROCHLORiDE ALLERGY, CETiRiZiNE HYDROCHLORiDE (OTC)
  - CETiRiZiNE HYDROCHLORiDE HIVES, CETiRiZiNE HYDROCHLORiDE (OTC)
  - CHLOROQUiNE PHOSPHATE, CHLOROQUiNE PHOSPHATE
  - FUROSEMiDE, FUROSEMiDE
  - HYDROCHLOROTHIAZiDE, HYDROCHLOROTHIAZiDE
  - HYDROXYCHLOROQUiNE SULFATE, HYDROXYCHLOROQUiNE SULFATE
  - LOSARTAN POTASSiUM AND HYDROCHLOROTHIAZiDE, HYDROCHLOROTHIAZiDE
  - LOSARTAN POTASSiUM, LOSARTAN POTASSiUM
  - METOCLOPRAMiDE HYDROCHLORiDE, METOCLOPRAMiDE HYDROCHLORiDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - ONDANSETRON HYDROCHLORiDE, ONDANSETRON HYDROCHLORiDE
  - PROPRANOLOL HYDROCHLORiDE, PROPRANOLOL HYDROCHLORiDE
  - TRAMADOL HYDROCHLORiDE, TRAMADOL HYDROCHLORiDE
  - WARFARiN SODiUM, WARFARiN SODiUM

**IPR**

- \* iPR PHARMACEUTICALS INC
  - CRESTOR, ROSUVASTATiN CALCiUM
  - ZOMiG, ZOLMITriPTAN

**IPSEN INC**

- \* iPSEN BIOPHARMACEUTICALS INC
  - INCRELEX, MECASERMIN RECOMBiNANT
  - ONiViDY, IRiNOTECAN HYDROCHLORiDE

**IPSEN PHARMA**

- \* iPSEN PHARMA BiOTECH SAS
  - SOMATULiNE DEPOT, LANREOTiDE ACETATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

**IROKO PHARMS**

- \* IROKO PHARMACEUTICALS LLC  
INDOCIN, INDOMETHACIN

**IROKO PHARMS LLC**

- \* IROKO PHARMACEUTICALS LLC  
TIVORBEX, INDOMETHACIN  
VIVLODEX, MELOXICAM  
ZORVOLEX, DICLOFENAC

**IRONWOOD PHARMS INC**

- \* IRONWOOD PHARMACEUTICALS INC  
DUZALLO, ALLOPURINOL  
ZURAMPIC, LESINURAD

**ISO TEX**

- \* ISO TEX DIAGNOSTICS INC  
JEANATOPE, ALBUMIN IODINATED I-125 SERUM  
MEGATOPE, ALBUMIN IODINATED I-131 SERUM

**ISOTEX**

- \* ISOTEX DIAGNOSTICS  
GLOFIL-125, IOTHALAMATE SODIUM I-125

**ISTITUTO BIO ITA SPA**

- \* ISTITUTO BIOCHIMICO ITALIANO SPA  
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM  
AMPICILLIN SODIUM, AMPICILLIN SODIUM  
NAFCILLIN SODIUM, NAFCILLIN SODIUM  
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM  
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM  
PIPERACILLIN, PIPERACILLIN SODIUM

**IVAX PHARMS**

- \* IVAX PHARMACEUTICALS INC  
VALSARTAN, VALSARTAN

**IVAX PHARMS INC**

- \* IVAX PHARMACEUTICALS INC  
OLANZAPINE, OLANZAPINE

**IVAX SUB TEVA PHARMS**

- \* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA  
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE  
BACLOFEN, BACLOFEN  
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
BUMETANIDE, BUMETANIDE  
CABERGOLINE, CABERGOLINE  
CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CIMETIDINE, CIMETIDINE (OTC)  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
CLOZAPINE, CLOZAPINE  
CYCLOSPORINE, CYCLOSPORINE  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
DIAZEPAM, DIAZEPAM  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM  
FAMOTIDINE, FAMOTIDINE  
FAMOTIDINE, FAMOTIDINE (OTC)  
FLUCONAZOLE, FLUCONAZOLE  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
FUROSEMIDE, FUROSEMIDE  
GABAPENTIN, GABAPENTIN  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
INDOMETHACIN, INDOMETHACIN  
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LISINOPRIL, LISINOPRIL  
METHYLDOPA, METHYLDOPA  
MISOPROSTOL, MISOPROSTOL  
NADOLOL, NADOLOL  
OXAPROZIN, OXAPROZIN  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* I \*\*

- \* IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)  
SIMVASTATIN, SIMVASTATIN  
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE  
TOLTERODINE TARTRATE, TOLTERODINE TARTRATE  
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

\*\* J \*\*

**J AND J CONSUMER INC**

- \* JOHNSON AND JOHNSON CONSUMER INC MCNEIL CONSUMER HEALTHCARE DIVISION  
CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)  
CHILDREN'S MOTRIN, IBUPROFEN (OTC)  
CHILDREN'S ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)  
IMODIUM MULTI-SYMPTOM RELIEF, LOPERAMIDE HYDROCHLORIDE (OTC)  
JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)  
MOTRIN IB, IBUPROFEN (OTC)  
PEPCID AC, FAMOTIDINE (OTC)  
PEPCID AC, FAMOTIDINE (OTC)  
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)  
SINE-AID IB, IBUPROFEN (OTC)  
SUDAFED 24 HOUR, PSEUDOEPHENDRINE HYDROCHLORIDE (OTC)  
TYLENOL, ACETAMINOPHEN (OTC)  
ZYRTEC ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
ZYRTEC HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
ZYRTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE (OTC)

**JACOBUS**

- \* JACOBUS PHARMACEUTICAL CO  
DAPSONE, DAPSONE  
PASER, AMINOSALICYLIC ACID

**JANSSEN BIOTECH**

- \* JANSSEN BIOTECH INC  
ZYTIGA, ABIRATERONE ACETATE

**JANSSEN PHARMS**

- \* JANSSEN PHARMACEUTICALS INC  
AXERT, ALMOTRIPTAN MALATE  
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE  
DITROPAN XL, OXYBUTYNIN CHLORIDE  
DURAGESIC-100, FENTANYL  
DURAGESIC-12, FENTANYL  
DURAGESIC-25, FENTANYL  
DURAGESIC-50, FENTANYL  
DURAGESIC-75, FENTANYL  
ELMIRON, PENTOSAN POLYSULFATE SODIUM  
HALDOL, HALOPERIDOL DECANOATE  
HALDOL, HALOPERIDOL LACTATE  
INVEGA SUSTENNA, PALIPERIDONE PALMITATE  
INVEGA TRINZA, PALIPERIDONE PALMITATE  
INVEGA, PALIPERIDONE  
INVOKAMET XR, CANAGLIFLOZIN  
INVOKAMET, CANAGLIFLOZIN  
INVOKANA, CANAGLIFLOZIN  
LEVAQUIN, LEVOFLOXACIN  
MICRONOR, NORETHINDRONE  
MODICON 28, ETHINYLMESTRADIOL  
NIZORAL, KETOCONAZOLE  
ORTHO CYCLEN-28, ETHINYLMESTRADIOL  
ORTHO TRI-CYCLEN LO, ETHINYLMESTRADIOL  
ORTHO TRI-CYCLEN, ETHINYLMESTRADIOL  
ORTHO-NOVUM 1/35-28, ETHINYLMESTRADIOL  
ORTHO-NOVUM 7/7/7-28, ETHINYLMESTRADIOL  
PANCREAZE, PANCRELIPASE (AMYLASE  
RAZADYNE ER, GALANTAMINE HYDROBROMIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* J \*\*

- \* JANSSEN PHARMACEUTICALS INC
  - RAZADYNE, GALANTAMINE HYDROBROMIDE
  - RISPERDAL CONSTA, RISPERIDONE
  - RISPERDAL, RISPERIDONE
  - SPORANOX, ITRACONAZOLE
  - TERAZOL 3, TERCONAZOLE
  - TERAZOL 7, TERCONAZOLE
  - TOPAMAX, TOPIRAMATE
  - TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
  - TYLENOL W/ CODEINE NO. 4, ACETAMINOPHEN
  - ULTRACET, ACETAMINOPHEN
  - ULTRAM, TRAMADOL HYDROCHLORIDE
  - VERMOX, MEBENDAZOLE
  - XARELTO, RIVAROXABAN

**JANSSEN PRODS**

- \* JANSSEN PRODUCTS LP
  - EDURANT, RILPIVIRINE HYDROCHLORIDE
  - OLYSIO, SIMEPREVIR SODIUM
  - PREZCOBIX, COBICISTAT
  - PREZISTA, DARUNAVIR ETHANOLATE
  - YONDELIS, TRABECTEDIN

**JANSSEN R AND D**

- \* JANSSEN RESEARCH AND DEVELOPMENT LLC
  - INTELENCE, ETRAVIRINE

**JANSSEN RES AND DEV**

- \* JANSSEN RESEARCH AND DEVELOPMENT LLC
  - DOXIL (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE

**JANSSEN THERAP**

- \* JANSSEN THERAPEUTICS DIV JANSSEN PRODUCTS LP
  - SIRTURO, BEDAQUILINE FUMARATE

**JAVELIN PHARMS INC**

- \* JAVELIN PHARMACEUTICALS INC A WHOLLY OWNED SUDSIDIARY OF HOSPIRA INC
  - DYLOJECT, DICLOFENAC SODIUM

**JAZZ PHARMS**

- \* JAZZ PHARMACEUTICALS INC
  - XYREM, SODIUM OXYBATE

**JAZZ PHARMS III**

- \* JAZZ PHARMACEUTICALS III INTERNATIONAL LTD
  - FAZACLO ODT, CLOZAPINE

**JAZZ PHARMS INC**

- \* JAZZ PHARMACEUTICALS INC
  - DEFITELIO, DEFIBROTIDE SODIUM

**JAZZ PHARMS INTL**

- \* JAZZ PHARMACEUTICALS INTERNATIONAL LTD
  - PRIALT, ZICONOTIDE ACETATE

**JIANGSU HANSOH PHARM**

- \* JIANGSU HANSOH PHARMACEUTICAL GROUP CO LTD
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - VINORELBINE TARTRATE, VINORELBINE TARTRATE

**JIANGSU HENGRIUI MED**

- \* JIANGSU HENGRIUI MEDICINE CO LTD
  - CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE
  - CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE
  - CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DOCETAXEL, DOCETAXEL
  - GABAPENTIN, GABAPENTIN
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - LETROZOLE, LETROZOLE
  - OXALIPLATIN, OXALIPLATIN

**JOHNS HOPKINS UNIV**

- \* JOHNS HOPKINS UNIV
  - AMMONIA N 13, AMMONIA N-13

**JOHNSON AND JOHNSON**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* J \*\*

- \* JOHNSON AND JOHNSON CONSUMER INC  
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)  
VISINE-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
- \* JOHNSON AND JOHNSON GROUP CONSUMER COMPANIES  
MEN'S ROGAINE, MOXIDIL (OTC)  
ROGAINE (FOR MEN), MOXIDIL (OTC)  
ROGAINE (FOR WOMEN), MOXIDIL (OTC)  
ROGAINE EXTRA STRENGTH (FOR MEN), MOXIDIL (OTC)  
WOMEN'S ROGAINE, MOXIDIL (OTC)
- \* JOHNSON AND JOHNSON HEALTHCARE PRODUCTS DIV MCNEIL-PPC INC  
NIZORAL A-D, KETOCONAZOLE (OTC)

**JUBILANT CADISTA**

- \* JUBILANT CADISTA PHARMACEUTICALS INC  
ALENDRONATE SODIUM, ALENDRONATE SODIUM  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)  
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LAMOTRIGINE, LAMOTRIGINE  
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE  
METHYLPREDNISOLONE, METHYLPREDNISOLONE  
PREDNISONE, PREDNISONE  
PROCOMP, PROCHLORPERAZINE MALEATE  
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

**JUBILANT DRAXIMAGE**

- \* JUBILANT DRAXIMAGE INC  
DRAX EXAMETAZIME, TECHNETIUM TC-99M EXAMETAZIME KIT  
DRAXIMAGE MDP-25, TECHNETIUM TC-99M MEDRONATE  
HICON, SODIUM IODIDE I-131  
PULMOLITE, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT  
RUBY-FILL, RUBIDIUM CHLORIDE RB-82  
SODIUM IODIDE I-131, SODIUM IODIDE I-131
- \* JUBILANT DRAXIMAGE RADIOPHARMACIES INC  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18
- \* JUBILANT DRAXIMAGE USA INC  
TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT

**JUBILANT GENERICS**

- \* JUBILANT GENERICS LTD  
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
CELECOXIB, CELECOXIB  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
FELODIPINE, FELODIPINE  
INDOMETHACIN, INDOMETHACIN  
IRBESARTAN, IRBESARTAN  
ITRACONAZOLE, ITRACONAZOLE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
LEVOFLOXACIN, LEVOFLOXACIN  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
OLANZAPINE, OLANZAPINE  
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM  
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE  
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE  
RISPERIDONE, RISPERIDONE  
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* J \*\*

- \* JUBILANT GENERICS LTD
  - SPIRONOLACTONE, SPIRONOLACTONE
  - TELMISARTAN, TELMISARTAN
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VALSARTAN, VALSARTAN
  - ZOLMITRIPTAN, ZOLMITRIPTAN

**JUBILANT HOLLISTERSTR**

- \* JUBILANT HOLLISTERSTIER LLC
  - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**STEVENS J**

- \* JEROME STEVENS PHARMACEUTICALS INC
  - BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
  - DIGOXIN, DIGOXIN
  - METHOCARBAMOL AND ASPIRIN, ASPIRIN
  - UNITHROID, LEVOTHYROXINE SODIUM \*\*

\*\* K \*\*

**GRIFFEN**

- \* KW GRIFFEN CO
  - BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

**KADMON PHARMS LLC**

- \* KADMON PHARMACEUTICALS LLC
  - RIBASPHERE, RIBAVIRIN
  - RIBAVIRIN, RIBAVIRIN

**KAI PHARMS INC**

- \* KAI PHARMACEUTICALS INC A WHOLLY OWNED SUBSIDIARY OF AMGEN INC
  - PARSABIV, ETELCALCETIDE

**KALEO INC**

- \* KALEO INC
  - AUVI-Q, EPINEPHRINE
  - EVZIO, NALOXONE HYDROCHLORIDE

**KASTLE THERAPS LLC**

- \* KASTLE THERAPEUTICS LLC
  - KYNAMRO, MIPOMERSEN SODIUM

**KEN LIFESCIENCE**

- \* KEN LIFESCIENCE
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

**KERYX BIOPHARMS**

- \* KERYX BIOPHARMACEUTICALS INC
  - AURYXIA, FERRIC CITRATE

**KETTERING MEDCTR**

- \* KETTERING MEDCTR
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**KINEDEXE UK**

- \* KINEDEXE UK LTD
  - PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE

**KING PHARMS**

- \* KING PHARMACEUTICALS INC
  - SYNERCID, DALFOPRISTIN
- \* KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC
  - CYTOMEL, LIOTHYRONINE SODIUM
  - LEVOXYL, LEVOTHYROXINE SODIUM \*\*
    - TUSSIGON, HOMATROPINE METHYLBROMIDE
- \* KING PHARMACEUTICALS RESEARCH AND DEVELOPMENT LLC A SUB OF PFIZER INC
  - SKELAXIN, METAXALONE

**KING PHARMS LLC**

- \* KING PHARMACEUTICALS LLC
  - ALTACE, RAMIPRIL
  - BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
  - BICILLIN C-R, PENICILLIN G BENZATHINE
  - BICILLIN L-A, PENICILLIN G BENZATHINE
  - CORZIDE, BENDROFLUMETHIAZIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* K \*\*

- \* KING PHARMACEUTICALS LLC
  - PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
  - SILVADENE, SILVER SULFADIAZINE
  - TAPAZOLE, METHIMAZOLE
  - TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE

**KNIGHT THERAPS**

- \* KNIGHT THERAPEUTICS USA INC
  - IMPAVIDO, MILTEFOSINE

**KOWA CO**

- \* KOWA CO LTD
  - LIVALO, PITAVASTATIN CALCIUM

**KREITCHMAN PET CTR**

- \* KREITCHMAN PET CENTER
  - AMMONIA N 13, AMMONIA N-13
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**KREMERS URBAN PHARMS**

- \* KREMERS URBAN PHARMACEUTICALS INC
  - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - GLYCOLAX, POLYETHYLENE GLYCOL 3350
  - GLYCOLAX, POLYETHYLENE GLYCOL 3350 (OTC)
  - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - OMEPRAZOLE, OMEPRAZOLE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - TEMOZOLOMIDE, TEMOZOLOMIDE

**KRKA TOVARNA ZDRAVIL**

- \* KRKA TOVARNA ZDRAVIL DD NOVO MESTO
  - LANSOPRAZOLE, LANSOPRAZOLE

**KVK TECH**

- \* KVK TECH INC
  - BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE
  - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
  - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - KALEXATE, SODIUM POLYSTYRENE SULFONATE
  - PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

**KVK TECH INC**

- \* KVK TECH INC
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

**KYOWA KIRIN**

- \* KYOWA KIRIN INC
  - FARESTON, TOREMIFENE CITRATE
  - SANCUSO, GRANisetron

**KYTHERA BIOPHARMS**

- \* KYTHERA BIOPHARMACEUTICALS INC
  - KYBELLA, DEOXYCHOLIC ACID

\*\* L \*\*

**L PERRIGO CO**

- \* L PERRIGO CO
  - CIMETIDINE, CIMETIDINE (OTC)
  - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
  - IBUPROFEN, IBUPROFEN (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

- \* L PERRIGO CO
  - JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
  - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
  - MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
  - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
  - PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

**LA JOLLA PHARM CO**

- \* LA JOLLA PHARMACEUTICAL CO
  - GIAPREZA, ANGIOTENSIN II

**LAB HRA PHARMA**

- \* LABORATOIRE HRA PHARMA
  - ELLA, ULIPRISTAL ACETATE

**LABORATORIOS GRIFOLS**

- \* LABORATORIOS GRIFOLS SA
  - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**LABORATORIOS SALVAT**

- \* LABORATORIOS SALVAT SA
  - OTOVEL, CIPROFLOXACIN HYDROCHLORIDE

**LABS LEON FARMA**

- \* LABORATORIOS LEON FARMA SA
  - ALTAVERA, ETHINYLMESTRADIOL
  - ELIFEMME, ETHINYLMESTRADIOL
  - ESTARYLLA, ETHINYLMESTRADIOL
  - INTROVALE, ETHINYLMESTRADIOL
  - ISIBLOOM, DESOGESTREL
  - JAIMIES, ETHINYLMESTRADIOL
  - LEVONORGESTREL AND ETHINYLMESTRADIOL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL
  - LORYNA, DROSPIRENONONE
  - SYEDA, DROSPIRENONONE
  - TRI-ESTARYLLA, ETHINYLMESTRADIOL
  - TRI-LO-ESTARYLLA, ETHINYLMESTRADIOL
  - VIENVA, ETHINYLMESTRADIOL
  - VOLNEA, DESOGESTREL

**LABS LICONSA**

- \* LABORATORIOS LICONSA SA
  - LANSOPRAZOLE, LANSOPRAZOLE

**LANDELA PHARM**

- \* LANDELA PHARMACEUTICAL
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

**LANNETT**

- \* LANNETT CO INC
  - ACETAZOLAMIDE, ACETAZOLAMIDE
  - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
  - DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
  - LANIAZID, ISONIAZID
  - LANORINAL, ASPIRIN
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PRIMIDONE, PRIMIDONE
  - PROBALAN, PROBENECID
- \* LANNETT HOLDINGS INC
  - BACLOFEN, BACLOFEN
  - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
  - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
  - DANAZOL, DANAZOL
  - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
  - DOXYCYCLINE, DOXYCYCLINE
  - FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
  - HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
  - RIFAMPIN, RIFAMPIN
  - TERBUTALINE SULFATE, TERBUTALINE SULFATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

- \* LANNETT HOLDINGS INC  
URSODIOL, URSODIOL

**LANNETT CO INC**

- \* LANNETT CO INC  
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE MONOKET, ISOSORBIDE MONONITRATE NIACIN, NIACIN SUMATRIPTAN, SUMATRIPTAN

**LANNETT HOLDINGS INC**

- \* LANNETT HOLDINGS INC  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN CODEINE SULFATE, CODEINE SULFATE DIAZEPAM, DIAZEPAM DIETHYLPROMION HYDROCHLORIDE, DIETHYLPROMION HYDROCHLORIDE HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN LOXAPINE SUCCINATE, LOXAPINE SUCCINATE METAXALONE, METAXALONE MORPHINE SULFATE, MORPHINE SULFATE NEOMYCIN SULFATE, NEOMYCIN SULFATE OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

**LANTHEUS MEDICAL IMAGING INC**

- \* LANTHEUS MEDICAL IMAGING INC  
CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT DEFINITY, PERFLUTREN GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67 NEUROLITE, TECHNETIUM TC-99M BICISATE KIT TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201 XENON XE 133, XENON XE-133

**LANTHEUS MEDICAL**

- \* LANTHEUS MEDICAL IMAGING INC  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18 QUADRAMET, SAMARIUM SM-153 LEXIDRONAM PENTASODIUM

**LARKEN LABS**

- \* LARKEN LABORATORIES INC  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE OFLOXACIN, OFLOXACIN

**LARKEN LABS INC**

- \* LARKEN LABORATORIES INC  
ACETAMINOPHEN, CAFEINE AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN ALLZITAL, ACETAMINOPHEN BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN DEXAMETHASONE, DEXAMETHASONE HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN

**LAVIPHARM LABS**

- \* LAVIPHARM LABORATORIES INC  
FENTANYL-100, FENTANYL FENTANYL-25, FENTANYL FENTANYL-50, FENTANYL FENTANYL-75, FENTANYL

**LEADANT BIOSCI INC**

- \* LEADANT BIOSCIENCES INC  
ABELCET, AMPHOTERICIN B CARNITOR SF, LEVOCARNITINE CARNITOR, LEVOCARNITINE CYSTARAN, CYSTEAMINE HYDROCHLORIDE MATULANE, PROCARBAZINE HYDROCHLORIDE

**LEADING PHARMA LLC**

- \* LEADING PHARMA LLC  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

- \* LEADING PHARMA LLC
  - FOLIC ACID, FOLIC ACID
  - FUROSEMIDE, FUROSEMIDE
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
  - LORAZEPAM, LORAZEPAM

**LEHIGH VALLEY**

- \* LEHIGH VALLEY TECHNOLOGIES INC
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

**LEO LABS**

- \* LEO LABORATORIES LTD
  - PICATO, INGENOL MEBUTATE

**LEO PHARMA AS**

- \* LEO PHARMA AS
  - DOVONEX, CALCIPOTRIENE
  - ENSTILAR, BETAMETHASONE DIPROPIONATE
  - PROTOPIC, TACROLIMUS
  - TACLONEX, BETAMETHASONE DIPROPIONATE

**LEXICON PHARMS INC**

- \* LEXICON PHARMACEUTICALS INC
  - XERMELO, TELOTRISTAT ETIPRATE

**LG CHEM LTD**

- \* LG CHEM LTD
  - FACTIVE, GEMIFLOXACIN MESYLATE
  - VALTROPIN, SOMATROPIN RECOMBINANT

**LIBERTY PHARMA INC**

- \* LIBERTY PHARMA INC
  - PRASUGREL, PRASUGREL HYDROCHLORIDE

**LIEBEL-FLARSHEIM**

- \* LIEBEL-FLARSHEIM CO LLC
  - CONRAY 30, IOTHALAMATE MEGLUMINE
  - CONRAY 43, IOTHALAMATE MEGLUMINE
  - CONRAY, IOTHALAMATE MEGLUMINE
  - CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
  - MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
  - OPTIMARK IN PLASTIC CONTAINER, GADOVERSETAMIDE
  - OPTIMARK, GADOVERSETAMIDE
  - OPTIRAY 240, IOVERSOL
  - OPTIRAY 300, IOVERSOL
  - OPTIRAY 320, IOVERSOL
  - OPTIRAY 350, IOVERSOL
  - SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

**LIFESTAR PHARMA**

- \* LIFESTAR PHARMA LLC
  - RISPERIDONE, RISPERIDONE

**LNK**

- \* LNK INTERNATIONAL INC
  - DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
  - IBUPROFEN, IBUPROFEN (OTC)
  - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

**LNK INTL INC**

- \* LNK INTERNATIONAL INC
  - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)

**LOREAL USA**

- \* LOREAL USA PRODUCTS INC
  - ANTHELIOS 20, AVOBENZONE (OTC)
  - ANTHELIOS 40, AVOBENZONE (OTC)
  - ANTHELIOS SX, AVOBENZONE (OTC)
  - CAPITAL SOLEIL 15, AVOBENZONE (OTC)

**LOTUS PHARM CO LTD**

- \* LOTUS PHARMACEUTICAL CO LTD
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVONORGESTREL, LEVONORGESTREL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

- \* LOTUS PHARMACEUTICAL CO LTD  
LEVONORGESTREL, LEVONORGESTREL (OTC)  
ROWEPPRA, LEVETIRACETAM

**LUITPOLD**

- \* LUITPOLD PHARMACEUTICALS INC  
ACETYLCYSTEINE, ACETYLCYSTEINE  
ADENOSINE, ADENOSINE  
AMINOCAPROIC ACID, AMINOCAPROIC ACID  
AMINOPHYLLINE, AMINOPHYLLINE  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE  
BETAMETHASONE ACETATE AND BETAMETHASONE SODIUM PHOSPHATE, BETAMETHASONE ACETATE  
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
CAFFEINE CITRATE, CAFFEINE CITRATE  
CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
CYANOCOBALAMIN, CYANOCOBALAMIN  
CYCLOSPORINE, CYCLOSPORINE  
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE  
DEXFERRUM, IRON DEXTRAN  
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE  
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE  
DROPERIDOL, DROPERIDOL  
ESTRADIOL VALERATE, ESTRADIOL VALERATE  
ETOMIDATE, ETOMIDATE  
FOMEPIZOLE, FOMEPIZOLE  
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM  
GLYCOPYRROLATE, GLYCOPYRROLATE  
GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE  
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
IBUTILIDE FUMARATE, IBUTILIDE FUMARATE  
LEVETIRACETAM, LEVETIRACETAM  
LEVOCARNITINE, LEVOCARNITINE  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
MANNITOL 25%, MANNITOL  
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE  
METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE  
METOPROLOL TARTRATE, METOPROLOL TARTRATE  
NITROGLYCERIN, NITROGLYCERIN  
OLANZAPINE, OLANZAPINE  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM  
PHENYTOIN SODIUM, PHENYTOIN SODIUM  
PROGESTERONE, PROGESTERONE  
TRANEXAMIC ACID, TRANEXAMIC ACID  
TRIMETHOBENZAMIDE HYDROCHLORIDE PRESERVATIVE FREE, TRIMETHOBENZAMIDE HYDROCHLORIDE  
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE  
VENOFE, IRON SUCROSE  
ZIDOVUDINE, ZIDOVUDINE

**LUITPOLD PHARMS INC**

- \* LUITPOLD PHARMACEUTICALS INC  
BUSULFAN, BUSULFAN  
CALCIUM CHLORIDE 10%, CALCIUM CHLORIDE  
DACTINOMYCIN, DACTINOMYCIN  
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE  
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE  
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE  
GANCICLOVIR, GANCICLOVIR SODIUM  
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE  
INJECTAFER, FERRIC CARBOXYMALTPOSE  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
METHOCARBAMOL, METHOCARBAMOL  
NANDROLONE DECANOATE, NANDROLONE DECANOATE  
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

- \* LUITPOLD PHARMACEUTICALS INC  
OXALIPLATIN, OXALIPLATIN  
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE

**LUKARE MEDICAL LLC**

- \* LUKARE MEDICAL LLC  
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE

**LUNDBECK NA LTD**

- \* LUNDBECK NA LTD  
NORTHERA, DROXIDOPA

**LUNDBECK PHARMS LLC**

- \* LUNDBECK PHARMACEUTICALS LLC  
CARNEXIV, CARBAMAZEPINE  
ONFI, CLOBAZAM  
SABRIL, VIGABATRIN

**LUPIN**

- \* LUPIN INC  
SOLOSEC, SECNIDAZOLE
- \* LUPIN LTD  
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
CARVEDILOL, CARVEDILOL  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CEFDINIR, CEFDINIR  
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM  
CEFPROZIL, CEFPROZIL  
CEFTRIAXONE, CEFTRIAXONE SODIUM  
CEFUROXIME AXETIL, CEFUROXIME AXETIL  
CEPHALEXIN, CEPHALEXIN  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE  
LEVETIRACETAM, LEVETIRACETAM  
LEVOFLOXACIN, LEVOFLOXACIN  
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LISINOPRIL, LISINOPRIL  
LOVASTATIN, LOVASTATIN  
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE  
RAMIPRIL, RAMIPRIL  
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE  
SIMVASTATIN, SIMVASTATIN  
TOPIRAMATE, TOPIRAMATE  
TRANDOLAPRIL, TRANDOLAPRIL

**LUPIN ATLANTIS**

- \* LUPIN ATLANTIS HOLDINGS SA  
ANTARA (MICRONIZED), FENOFIBRATE  
DESOXIMETASONE, DESOXIMETASONE  
MIBELAS 24 FE, ETHINYL ESTRADIOL  
OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE  
TOBRAMYCIN, TOBRAMYCIN  
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**LUPIN LTD**

- \* LUPIN LIMITED  
LEVETIRACETAM, LEVETIRACETAM  
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
- \* LUPIN LTD  
ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE  
ABACAVIR SULFATE, LAMIVUDINE AND ZIDOVUDINE, ABACAVIR SULFATE  
AMABELZ, ESTRADIOL  
AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE  
ARMODAFINIL, ARMODAFINIL  
AZITHROMYCIN, AZITHROMYCIN  
BEKYREE, DESOGESTREL  
BIMATOPROST, BIMATOPROST  
BLISOVI 24 FE, ETHINYL ESTRADIOL  
BLISOVI FE 1.5/30, ETHINYL ESTRADIOL  
BLISOVI FE 1/20, ETHINYL ESTRADIOL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

- \* LUPIN LTD  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
CALCIUM ACETATE, CALCIUM ACETATE  
CELECOXIB, CELECOXIB  
CIPROFLOXACIN, CIPROFLOXACIN  
CLARITHROMYCIN, CLARITHROMYCIN  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
DAYSEE, ETHINYLMESTRADIOL  
DESVENLAFAZINE SUCCINATE, DESVENLAFAZINE SUCCINATE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
DOXYCYCLINE, DOXYCYCLINE  
DROSPIRENONE AND ETHINYLMESTRADIOL, DROSPIRENONE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
ENSKYCE, DESOGESTREL  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
ESZOPICLONE, ESZOPICLONE  
FALLBACK SOLO, LEVONORGESTREL (OTC)  
FAMOTIDINE, FAMOTIDINE  
FAYOSIM, ETHINYLMESTRADIOL  
FENOFLIBRATE, FENOFLIBRATE  
FENOFLIBRIC ACID, CHOLINE FENOFLIBRATE  
FYAVOLV, ETHINYLMESTRADIOL  
GATIFLOXACIN, GATIFLOXACIN  
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE  
IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE  
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
IRBESARTAN, IRBESARTAN  
JENCYCLA, NORETHINDRONE  
KAITLIB FE, ETHINYLMESTRADIOL  
KURVELO, ETHINYLMESTRADIOL  
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE  
LAMIVUDINE, LAMIVUDINE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
LEVONORGESTREL AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL  
LORAZEPAM, LORAZEPAM  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
MEFENAMIC ACID, MEFENAMIC ACID  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
METRONIDAZOLE, METRONIDAZOLE  
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE  
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE  
NABUMETONE, NABUMETONE  
NADOLOL, NADOLOL  
NIACIN, NIACIN  
NIKKI, DROSPIRENONE  
NORETHINDRONE AND ETHINYLMESTRADIOL AND FERROUS FUMARATE, ETHINYLMESTRADIOL  
NORETHINDRONE, NORETHINDRONE  
NORGESTIMATE AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL  
OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL  
OMEPRAZOLE, OMEPRAZOLE  
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE  
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE  
PIRMELLA 1/35, ETHINYLMESTRADIOL  
PIRMELLA 7/7/7, ETHINYLMESTRADIOL  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
QUETIAPINE FUMARATE, QUETIAPINE FUMARATE  
QUININE SULFATE, QUININE SULFATE  
RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM  
REPAGLINIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* L \*\*

\* LUPIN LTD  
 RIFABUTIN, RIFABUTIN  
 ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM  
 SUPRAX, CEFIXIME  
 TELMISARTAN AND AMLODIPIINE, AMLODIPIINE BESYLATE  
 TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
 TESTOSTERONE, TESTOSTERONE  
 TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
 TYDEMY, DROSPIRENONE  
 VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
 VALSARTAN, VALSARTAN  
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE  
 VYFEMLA, ETHINYLMESTRADIOL  
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**LUPIN PHARMS**

\* LUPIN PHARMACEUTICALS INC  
 AMLODIPIINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPIINE BESYLATE  
 DESLORATADINE, DESLORATADINE  
 MELOXICAM, MELOXICAM  
 NORGESTIMATE AND ETHINYLMESTRADIOL, ETHINYLMESTRADIOL  
 PRAVASTATIN SODIUM, PRAVASTATIN SODIUM  
 RIFAMPIN, RIFAMPIN  
 SUPRAX, CEFIXIME  
 ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

**LYMOL MEDCL**

\* LYMOL MEDICAL CORP  
 SCLEROSOL, TALC  
 TALC, TALC

**LYNE**

\* LYNE LABORATORIES INC  
 CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE  
 CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE  
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
 DEXAMETHASONE, DEXAMETHASONE  
 ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
 FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE  
 LEVOCARNITINE, LEVOCARNITINE  
 NYSTATIN, NYSTATIN  
 TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**PERRIGO**

\* L PERRIGO CO  
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)  
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)  
 CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)  
 CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)  
 DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)  
 FAMOTIDINE, FAMOTIDINE (OTC)  
 IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)  
 IBUPROFEN, IBUPROFEN (OTC)  
 LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)  
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)  
 LORATADINE, LORATADINE (OTC)  
 MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)  
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)  
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)  
 NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)  
 NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)  
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)  
 TAB-PROFEN, IBUPROFEN (OTC)  
 TIOCONAZOLE, TIOCONAZOLE (OTC)

\*\* M \*\*

**MA GENERAL HOSP**

\* MASSACHUSETTS GENERAL HOSP  
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

**MACLEODS PHARMS LTD**

- \* MACLEODS PHARMACEUTICALS LTD
  - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
  - CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
  - CELECOXIB, CELECOXIB
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - DARIFENACIN, DARIFENACIN HYDROBROMIDE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ENTACAPONE, ENTACAPONE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ESZOPICLONE, ESZOPICLONE
  - FAMCICLOVIR, FAMCICLOVIR
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IRBESARTAN, IRBESARTAN
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - NEVIRAPINE, NEVIRAPINE
  - OLANZAPINE, OLANZAPINE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RISEDRONATE SODIUM, RISEDRONATE SODIUM
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - VALSARTAN, VALSARTAN
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
  - ZOLMITRIPTAN, ZOLMITRIPTAN

**MAGNA PHARMS**

- \* MAGNA PHARMACEUTICALS INC
  - ZOLPIMIST, ZOLPIDEM TARTRATE

**MAIA PHARMS INC**

- \* MAIA PHARMACEUTICALS INC
  - SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

**MALLINCKRODT ARD**

- \* MALLINCKRODT ARD INC
  - H.P. ACTHAR GEL, CORTICOTROPIN

**MALLINCKRODT HOSP**

- \* MALLINCKRODT HOSPITAL PRODUCTS IP LTD
  - INOMAX, NITRIC OXIDE
  - UVADEX, METHOXSALEN

**MALLINCKRODT IP**

- \* MALLINCKRODT IP

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MALLINCKRODT IP  
OFIRMEV, ACETAMINOPHEN

**MALLINCKRODT NUCLEAR**

- \* MALLINCKRODT NUCLEAR MEDICINE LLC  
GALLIUM CITRATE GA 67, GALLIUM CITRATE GA-67  
INDIUM IN 111 CHLORIDE, INDIUM IN-111 CHLORIDE  
OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT  
SODIUM IODIDE I-123, SODIUM IODIDE I-123  
SODIUM IODIDE I-131, SODIUM IODIDE I-131  
TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT  
TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT  
TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT  
TECHNETIUM TC-99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT  
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE TL-201  
ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR  
ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT  
XENON XE 133, XENON XE-133

**MANNKIND**

- \* MANNKIND CORP  
AFREZZA, INSULIN RECOMBINANT HUMAN

**MARINA BIOTECH**

- \* MARINA BIOTECH INC  
PRESTALIA, AMLODIPINE BESYLATE

**MARKSANS PHARMA**

- \* MARKSANS PHARMA LTD  
DUTASTERIDE, DUTASTERIDE  
GABAPENTIN, GABAPENTIN  
IBUPROFEN, IBUPROFEN  
IBUPROFEN, IBUPROFEN (OTC)  
LORATADINE, LORATADINE (OTC)  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)  
NAPROXEN, NAPROXEN  
PARICALCITOL, PARICALCITOL

**MARNEL PHARMS**

- \* MARNEL PHARMACEUTICALS LLC  
CROTAN, CROTAMITON

**MAYER LABS INC**

- \* MAYER LABORATORIES INC  
TODAY, NONOXYNOL-9 (OTC)

**MAYNE PHARMA**

- \* MAYNE PHARMA INTERNATIONAL PTY LTD  
DORYX MPC, DOXYCYCLINE HYCLATE  
DORYX, DOXYCYCLINE HYCLATE  
ERYC, ERYTHROMYCIN
- \* MAYNE PHARMA LLC  
BUDESONIDE, BUDESONIDE  
CAMILA, NORETHINDRONE  
CARBIDOPA AND LEVODOPA, CARBIDOPA  
CLARITHROMYCIN, CLARITHROMYCIN  
CLONIDINE, CLONIDINE  
CLOZAPINE, CLOZAPINE  
CYCLOSPORINE, CYCLOSPORINE  
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
DIAZEPAM, DIAZEPAM  
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE  
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE  
ERRIN, NORETHINDRONE  
ESTAZOLAM, ESTAZOLAM  
ESTRADIOL, ESTRADIOL  
FABIOR, TAZAROTENE  
FENTANYL-100, FENTANYL  
FENTANYL-25, FENTANYL  
FENTANYL-50, FENTANYL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

\* MAYNE PHARMA LLC  
 FENTANYL-75, FENTANYL  
 LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
 LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL  
 LOW-OGESTREL-28, ETHINYL ESTRADIOL  
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
 MICROGESTIN 1.5/30, ETHINYL ESTRADIOL  
 MICROGESTIN 1/20, ETHINYL ESTRADIOL  
 MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL  
 MICROGESTIN FE 1/20, ETHINYL ESTRADIOL  
 NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL  
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE  
 SORILUX, CALCIPOTRIENE  
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE  
 TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL  
 TRIMETHOPRIM, TRIMETHOPRIM  
 TRIVORA-28, ETHINYL ESTRADIOL  
 ZOVIA 1/35E-28, ETHINYL ESTRADIOL

**MAYNE PHARMA INC**

\* MAYNE PHARMA INC  
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
 BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
 BUTALBITAL, ASPIRIN, CAFFEINE AND CODEINE PHOSPHATE, ASPIRIN  
 DOFETILIDE, DOFETILIDE  
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
 DOXYCYCLINE, DOXYCYCLINE  
 HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE, HYDROCODONE BITARTRATE  
 HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE,  
 LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM  
 METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE  
 MORPHINE SULFATE, MORPHINE SULFATE  
 NYSTATIN, NYSTATIN  
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
 OXYCODONE AND ASPIRIN, ASPIRIN  
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

**MCGUFF**

\* MCGUFF PHARMACEUTICALS INC  
 ASCOR, ASCORBIC ACID

**MCNEIL**

\* MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC  
 IBUPROFEN, IBUPROFEN (OTC)

**MCNEIL CONS**

\* MCNEIL CONSUMER HEALTHCARE  
 SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

**MCPRF**

\* MAYO CLINIC PET RADIOCHEMISTRY FACILITY  
 AMMONIA N 13, AMMONIA N-13  
 CHOLINE C-11, CHOLINE C-11  
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**MEDAC PHARMA INC**

\* MEDAC PHARMA INC  
 RASUVO, METHOTREXATE

**MEDEFIL INC**

\* MEDEFIL INC  
 SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

**MEDICINES360**

\* MEDICINES360  
 LILETTA, LEVONORGESTREL

**MEDICIS**

\* MEDICIS PHARMACEUTICAL CORP  
 ALDARA, IMIQUIMOD  
 AMMONUL, SODIUM BENZOATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MEDICIS PHARMACEUTICAL CORP
  - CALCIUM DISODIUM VERSENATE, EDETALE CALCIUM DISODIUM
  - LOPROX, CICLOPIROX
  - LUZU, LULICONAZOLE
  - METROGEL-VAGINAL, METRONIDAZOLE
  - MINITRAN, NITROGLYCERIN
  - SOLODYN, MINOCYCLINE HYDROCHLORIDE
  - VANOS, FLUOCINONIDE
  - ZIANA, CLINDAMYCIN PHOSPHATE
  - ZYCLARA, IMIQUIMOD

**MEDICURE**

- \* MEDICURE INTERNATIONAL INC
  - AGRASTAT, TIROFIBAN HYDROCHLORIDE

**MEDIGENE AG**

- \* MEDIGENE AG
  - VEREGEN, SINECATECHINS

**MEDIMETRIKS PHARMS**

- \* MEDIMETRIKS PHARMACEUTICALS INC
  - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
  - LOPROX, CICLOPIROX
  - NEO-SYNALAR, FLUOCINOLONE ACETONIDE
  - SYNALAR, FLUOCINOLONE ACETONIDE

**MEDTECH PRODUCTS**

- \* MEDTECH PRODUCTS INC
  - MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
  - MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
  - MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
  - MONISTAT 3, MICONAZOLE NITRATE
  - MONISTAT 3, MICONAZOLE NITRATE (OTC)
  - MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
  - MONISTAT 7, MICONAZOLE NITRATE (OTC)
  - NIX, PERMETHRIN (OTC)
  - TAGAMET HB, CIMETIDINE (OTC)

**MELINTA**

- \* MELINTA SUBSIDIARY CORP
  - BAXDELA, DELAFLOXACIN MEGLUMINE

**MEM SLOAN-KETTERING**

- \* MEMORIAL SLOAN-KETTERING CANCER CENTER
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**MERCK**

- \* MERCK AND CO INC
  - CANCIDAS, CASPOFUNGIN ACETATE
  - EMEND, APREPITANT
  - FOSAMAX PLUS D, ALENDRONATE SODIUM
  - MAXALT, RIZATRIPTAN BENZOATE
  - MAXALT-MLT, RIZATRIPTAN BENZOATE
  - PRIMAXIN, CILASTATIN SODIUM
  - PROSCAR, FINASTERIDE
  - ZOLINZA, VORINOSTAT
- \* MERCK RESEARCH LABORATORIES DIV MERCK CO INC
  - PRINIVIL, LISINOPRIL
  - PROPECIA, FINASTERIDE
  - SINGULAIR, MONTELUKAST SODIUM
  - TRUSOPT, DORZOLAMIDE HYDROCHLORIDE

**MERCK AND CO INC**

- \* MERCK AND CO INC
  - EMEND, FOSAPREPITANT DIMEGLUMINE
  - FOSAMAX, ALENDRONATE SODIUM

**MERCK SHARP DOHME**

- \* MERCK SHARP AND DOHME CORP
  - ASMANEX HFA, MOMETASONE FUROATE
  - ASMANEX TWISTHALER, MOMETASONE FUROATE
  - BELSOMRA, SUVOREXANT
  - CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
  - CLARINEX D 24 HOUR, DESLORATADINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

\* MERCK SHARP AND DOHME CORP  
 CLARINEX, DESLORATADINE  
 CLARINEX-D 12 HOUR, DESLORATADINE  
 COZAAR, LOSARTAN POTASSIUM  
 CRIXIVAN, INDINAVIR SULFATE  
 DIPROLENE AF, BETAMETHASONE DIPROPIONATE  
 DIPROLENE, BETAMETHASONE DIPROPIONATE  
 DULEREA, FORMOTEROL FUMARATE  
 ELOCON, MOMETASONE FUROATE  
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE  
 HYZAAR, HYDROCHLOROTHIAZIDE  
 INVANZ, ERTAPENEM SODIUM  
 ISENTRESS HD, Raltegravir POTASSIUM  
 ISENTRESS, Raltegravir POTASSIUM  
 JANUMET XR, METFORMIN HYDROCHLORIDE  
 JANUMET, METFORMIN HYDROCHLORIDE  
 JANUVIA, SITAGLIPTIN PHOSPHATE  
 LOTRISONE, BETAMETHASONE DIPROPIONATE  
 NASONEX, MOMETASONE FUROATE  
 NOXAFIL, POSACONAZOLE  
 PREVYMIS, LETERMOVIR  
 REBETOL, RIBAVIRIN  
 SEGLUROMET, ERTUGLIFLOZIN  
 SINEMET CR, CARBIDOPA  
 SINEMET, CARBIDOPA  
 STEGLATRO, ERTUGLIFLOZIN  
 STELUJAN, ERTUGLIFLOZIN  
 STROMECTOL, IVERMECTIN  
 TEMODAR, TEMOZOLOMIDE  
 ZEPATIER, ELBASVIR

**MERIDIAN MEDCL**

\* MERIDIAN MEDICAL TECHNOLOGIES INC  
 DUODOTE, ATROPINE

**MERIDIAN MEDCL TECHN**

\* MERIDIAN MEDICAL TECHNOLOGIES INC  
 ATROPEN, ATROPINE  
 MORPHINE SULFATE, MORPHINE SULFATE  
 PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE

**MERR PHARM**

\* MERR PHARMACEUTICAL CO LTD  
 IBUPROFEN, IBUPROFEN (OTC)

**MERZ PHARMS**

\* MERZ PHARMACEUTICALS LLC  
 CUVPOSA, GLYCOPYRROLATE

**METHAPHARM**

\* METHAPHARM INC  
 PROVOCHOLINE, METHACHOLINE CHLORIDE

**METUCHEN PHARMS**

\* METUCHEN PHARMACEUTICALS LLC  
 STENDRA, AVANAFIL

**MICRO LABS**

\* MICRO LABS LTD  
 AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE  
 CAFFEINE CITRATE, CAFFEINE CITRATE  
 PIROXICAM, PIROXICAM  
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE  
 TELMISARTAN, TELMISARTAN

**MICRO LABS LTD**

\* MICRO LABS LTD  
 NEVIRAPINE, NEVIRAPINE

**MICRO LABS LTD INDIA**

\* MICRO LABS LTD INDIA  
 AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN  
 CROMOLYN SODIUM, CROMOLYN SODIUM  
 LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MICRO LABS LTD INDIA  
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN

**MIDATECH PHARMA US**

- \* MIDATECH PHARMA US INC  
ORAVIG, MICONAZOLE  
SOLTAMOX, TAMOXIFEN CITRATE  
ZUPLENZ, ONDANSETRON

**MIDWEST MEDCL**

- \* MIDWEST MEDICAL ISOTOPES LLC CYCLOTRON DIV  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**MIKART**

- \* MIKART INC  
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
BENZONATATE, BENZONATATE  
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE  
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
BUTAPAP, ACETAMINOPHEN  
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE  
CHLORZOXAZONE, CHLORZOXAZONE  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE  
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE  
ETHOSUXIMIDE, ETHOSUXIMIDE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
ISONIAZID, ISONIAZID  
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE  
METHAZOLAMIDE, METHAZOLAMIDE  
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE  
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

**MIKART INC**

- \* MIKART INC  
BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN  
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE  
CHLORZOXAZONE, CHLORZOXAZONE

**MILLENNIUM PHARMS**

- \* MILLENNIUM PHARMACEUTICALS INC  
NINLARO, IXAZOMIB CITRATE  
VELCADE, BORTEZOMIB

**MIPS CRF**

- \* MIPS CYCLOTRON AND RADIOCHEMISTRY FACILITY  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**MISSION PHARMA**

- \* MISSION PHARMACAL CO  
BINOSTO, ALENDRONATE SODIUM  
LITHOSTAT, ACETOHYDROXAMIC ACID  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE  
TEXACORT, HYDROCORTISONE  
THIOLA, TIOPRONIN  
TINDAMAX, TINIDAZOLE  
UROCIT-K, POTASSIUM CITRATE

**MISSION PHARMACAL CO**

- \* MISSION PHARMACAL CO  
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE  
FLOWTUSS, GUAIFENESIN  
HYCOFENIX, GUAIFENESIN  
POTASSIUM IODIDE, POTASSIUM IODIDE (OTC)

**MIST PHARMS LLC**

- \* MIST PHARMACEUTICALS LLC  
NITROMIST, NITROGLYCERIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

**MITSUBISHI TANABE**

- \* MITSUBISHI TANABE PHARMA CORP  
RADICAVA, EDARAVONE

**MOBERG PHARMA NORTH**

- \* MOBERG PHARMA NORTH AMERICA LLC  
CHILDREN'S ELIXSURE, IBUPROFEN (OTC)

**MOBIUS THERAP**

- \* MOBIUS THERAPEUTICS LLC  
MITOSOL, MITOMYCIN

**MOLNLYCKE HLTH**

- \* MOLNLYCKE HEALTH CARE  
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)  
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

**MONARCH PHARMS**

- \* MONARCH PHARMACEUTICALS LLC  
CORTISPORIN, BACITRACIN ZINC  
CORTISPORIN, HYDROCORTISONE ACETATE  
CORTISPORIN, HYDROCORTISONE  
MENEST, ESTROGENS, ESTERIFIED  
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE  
NEOSPORIN, GRAMICIDIN  
SEPTRA DS, SULFAMETHOXAZOLE  
SEPTRA, SULFAMETHOXAZOLE  
VIROPTIC, TRIFLURIDINE

**MONTEREY PHARMS LLC**

- \* MONTEREY PHARMACEUTICALS LLC  
METHOCARBAMOL, METHOCARBAMOL

**MSD INTL**

- \* MSD INTERNATIONAL GMBH  
VYTORIN, EZETIMIBE

**MSD INTL GMBH**

- \* MSD INTERNATIONAL GMBH  
ZETIA, EZETIMIBE

**MSD MERCK CO**

- \* MERCK SHARP AND DOHME CORP A SUB OF MERCK AND CO INC  
EMEND, APREPITANT  
SINGULAIR, MONTELUKAST SODIUM  
ZOCOR, SIMVASTATIN

**MSN LABS PVT LTD**

- \* MSN LABORATORIES PRIVATE LTD  
CLOFARABINE, CLOFARABINE  
MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE  
ROUVASTATIN CALCIUM, ROUVASTATIN CALCIUM

**MURTY PHARMS**

- \* MURTY PHARMACEUTICALS INC  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
DIPYRIDAMOLE, DIPYRIDAMOLE

**MUSTAFA NEVZAT ILAC**

- \* MUSTAFA NEVZAT ILAC SANAYII AS (MN PHARMACEUTICALS)  
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM  
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

**MUTUAL PHARM**

- \* MUTUAL PHARMACEUTICAL CO INC  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
MINOXIDIL, MINOXIDIL  
PREDNISONE, PREDNISONE

**MYLAN**

- \* MYLAN PHARMACEUTICALS  
FENOFLIBRATE, FENOFLIBRATE  
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
TRANEXAMIC ACID, TRANEXAMIC ACID
- \* MYLAN PHARMACEUTICALS INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN PHARMACEUTICALS INC
  - ACARBOSE, ACARBOSE
  - ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - ALENDRONATE SODIUM, ALENDRONATE SODIUM
  - ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
  - ALLOPURINOL, ALLOPURINOL
  - ALPRAZOLAM, ALPRAZOLAM
  - AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
  - AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
  - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
  - ANASTROZOLE, ANASTROZOLE
  - ATENOLOL AND CHLORTHALIDONE, ATENOLOL
  - ATENOLOL, ATENOLOL
  - AVITA, TRETINOIN
  - AZATHIOPRINE, AZATHIOPRINE
  - AZITHROMYCIN, AZITHROMYCIN
  - BACLOFEN, BACLOFEN
  - BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
  - BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
  - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
  - BICALUTAMIDE, BICALUTAMIDE
  - BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
  - BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
  - BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
  - BUDESONIDE, BUDESONIDE
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
  - BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
  - CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
  - CARBIDOPA AND LEVODOPA, CARBIDOPA
  - CARVEDILOL, CARVEDILOL
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHLOROTHIAZIDE, CHLOROTHIAZIDE
  - CHLORPROPAMIDE, CHLORPROPAMIDE
  - CHLORTHALIDONE, CHLORTHALIDONE
  - CIMETIDINE, CIMETIDINE
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
  - CLONAZEPAM, CLONAZEPAM
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
  - CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
  - CLORPRES, CHLORTHALIDONE
  - CLOZAPINE, CLOZAPINE
  - CYSTAGON, CYSTEAMINE BITARTRATE
  - DIAZEPAM, DIAZEPAM
  - DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
  - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
  - DOXYCYCLINE, DOXYCYCLINE
  - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
  - ENALAPRIL MALEATE, ENALAPRIL MALEATE
  - ESTRADIOL, ESTRADIOL
  - ESTROPIPATE, ESTROPIPATE
  - ETIDRONATE DISODIUM, ETIDRONATE DISODIUM
  - ETOPOSIDE, ETOPOSIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN PHARMACEUTICALS INC
  - EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
  - FAMCICLOVIR, FAMCICLOVIR
  - FAMOTIDINE, FAMOTIDINE
  - FAMOTIDINE, FAMOTIDINE (OTC)
  - FELODIPINE, FELODIPINE
  - FENOFLIBRATE, FENOFLIBRATE
  - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
  - FINASTERIDE, FINASTERIDE
  - FLUCONAZOLE, FLUCONAZOLE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
  - FLURBIPROFEN, FLURBIPROFEN
  - FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
  - FUROSEMIDE, FUROSEMIDE
  - GABAPENTIN, GABAPENTIN
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - GLIMEPIRIDE, GLIMEPIRIDE
  - GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
  - GLIPIZIDE, GLIPIZIDE
  - GLYBURIDE (MICRONIZED), GLYBURIDE
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
  - HALOPERIDOL, HALOPERIDOL
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - INDAPAMIDE, INDAPAMIDE
  - INDOMETHACIN, INDOMETHACIN
  - KETOCONAZOLE, KETOCONAZOLE
  - KETOPROFEN, KETOPROFEN
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - LAMOTRIGINE, LAMOTRIGINE
  - LATANOPROST, LATANOPROST
  - LETROZOLE, LETROZOLE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOOTHYROXINE SODIUM, LEVOOTHYROXINE SODIUM \*\*
  - LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
  - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LISINOPRIL, LISINOPRIL
  - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
  - LORATADINE, LORATADINE (OTC)
  - LORAZEPAM, LORAZEPAM
  - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - LOVASTATIN, LOVASTATIN
  - LOXPINE SUCCINATE, LOXPINE SUCCINATE
  - MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
  - MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
  - MELOXICAM, MELOXICAM
  - MENTAX, BUTENAFINE HYDROCHLORIDE
  - MERCAPTOPURINE, MERCAPTOPURINE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHIMAZOLE, METHIMAZOLE
  - METHOTREXATE SODIUM, METHOTREXATE SODIUM
  - METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - METHYLDOPA, METHYLDOPA
  - METOLAZONE, METOLAZONE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MIRTAZAPINE, MIRTAZAPINE
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
  - NADOLOL, NADOLOL
  - NAPROXEN, NAPROXEN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN PHARMACEUTICALS INC
  - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
  - NIFEDIPINE, NIFEDIPINE
  - NISOLDIPINE, NISOLDIPINE
  - NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
  - NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
  - OMEPRAZOLE, OMEPRAZOLE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON, ONDANSETRON
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
  - PENTOXIFYLLINE, PENTOXIFYLLINE
  - PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
  - PHENYTEK, PHENYTOIN SODIUM
  - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
  - PROBENECID, PROBENECID
  - PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
  - PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
  - QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
  - RISPERIDONE, RISPERIDONE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
  - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
  - SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - SPIRONOLACTONE, SPIRONOLACTONE
  - STAVUDINE, STAVUDINE
  - SULINDAC, SULINDAC
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TACROLIMUS, TACROLIMUS
  - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TEMAZEPAM, TEMAZEPAM
  - THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
  - THIOTHIXENE, THIOTHIXENE
  - TIMOLOL MALEATE, TIMOLOL MALEATE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TOLMETIN SODIUM, TOLMETIN SODIUM
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
  - URSODIOL, URSODIOL
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
  - ZONISAMIDE, ZONISAMIDE

**MYLAN ASI**

- \* MYLAN ASI LLC
  - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE

**MYLAN INSTITUTIONAL**

- \* MYLAN INSTITUTIONAL INC
  - SULFAMYLYN, MAFENIDE ACETATE
- \* MYLAN INSTITUTIONAL LLC
  - ACETYLcysteine, ACETYLcysteine
  - ALOPRIM, ALLOPURINOL SODIUM
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - ARGATROBAN, ARGATROBAN
  - AZACITIDINE, AZACITIDINE
  - CARBOPLATIN, CARBOPLATIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN INSTITUTIONAL LLC
  - CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
  - CIDOFOVIR, CIDOFOVIR
  - COSYNTROPIN, COSYNTROPIN
  - DANTROLENE SODIUM, DANTROLENE SODIUM
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
  - DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
  - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
  - DURACLON, CLONIDINE HYDROCHLORIDE
  - ENILON, EDROPHONIUM CHLORIDE
  - ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
  - ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
  - FOMEPIZOLE, FOMEPIZOLE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - IBUTILIDE FUMARATE, IBUTILIDE FUMARATE
  - ISOSULFAN BLUE, ISOSULFAN BLUE
  - KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
  - MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
  - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
  - MESNA, MESNA
  - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
  - METHOCARBAMOL, METHOCARBAMOL
  - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
  - NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
  - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
  - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
  - PENTOSTATIN, PENTOSTATIN
  - PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
  - RIMSO-50, DIMETHYL SULFOXIDE
  - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
  - SOTRADECOL, SODIUM TETRADECYL SULFATE
  - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
  - TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
  - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
  - TRANEXAMIC ACID, TRANEXAMIC ACID
  - ULTIVA, REMIFENTANIL HYDROCHLORIDE

**MYLAN IRELAND LTD**

- \* MYLAN IRELAND LTD
  - ARIIXTRA, FONDAPARINUX SODIUM
  - CARBAMAZEPINE, CARBAMAZEPINE
  - MIACALCIN, CALCITONIN SALMON
  - PIROXICAM, PIROXICAM
  - SUCRALFATE, SUCRALFATE
  - THEOPHYLLINE, THEOPHYLLINE

**MYLAN LABS**

- \* MYLAN LABORATORIES LTD
  - NEVIRAPINE, NEVIRAPINE

**MYLAN LABS LTD**

- \* MYLAN LABORATORIES LTD
  - ADENOSINE, ADENOSINE
  - AMIFOSTINE, AMIFOSTINE
  - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
  - AMPICILLIN SODIUM, AMPICILLIN SODIUM
  - ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
  - ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
  - AZITHROMYCIN, AZITHROMYCIN
  - CAPREOMYCIN SULFATE, CAPREOMYCIN SULFATE
  - CARBOPLATIN, CARBOPLATIN
  - CASPOFUNGIN ACETATE, CASPOFUNGIN ACETATE
  - CISPLATIN, CISPLATIN
  - CLADRIBINE, CLADRIBINE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CLOFARABINE, CLOFARABINE
  - CYANOCOBALAMIN, CYANOCOBALAMIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN LABORATORIES LTD
  - CYTARABINE, CYTARABINE
  - DACTINOMYCIN, DACTINOMYCIN
  - DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
  - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
  - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
  - DOXYCYCLINE, DOXYCYCLINE HYCLATE
  - DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
  - EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE
  - ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
  - ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
  - ETHYNODIOL DIACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - ETOMIDATE, ETOMIDATE
  - ETOPOSIDE, ETOPOSIDE
  - FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
  - FAMOTIDINE, FAMOTIDINE
  - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
  - FLUMAZENIL, FLUMAZENIL
  - FLUOROURACIL, FLUOROURACIL
  - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
  - GANCICLOVIR, GANCICLOVIR SODIUM
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - HEPARIN SODIUM, HEPARIN SODIUM
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
  - IFOSFAMIDE, IFOSFAMIDE
  - LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LEVONORGESTREL AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - LEVONORGESTREL, LEVONORGESTREL
  - LEVONORGESTREL, LEVONORGESTREL (OTC)
  - LINEZOLID, LINEZOLID
  - METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
  - METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
  - MITOMYCIN, MITOMYCIN
  - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
  - MOXIFLOXACIN HYDROCHLORIDE IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN
  - MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
  - NAFCILLIN SODIUM, NAFCILLIN SODIUM
  - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
  - NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
  - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
  - NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - NORETHINDRONE, NORETHINDRONE
  - NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXALIPLATIN, OXALIPLATIN
  - PACLITAXEL, PACLITAXEL
  - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
  - PARICALCITOL, PARICALCITOL
  - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RIFAMPIN, RIFAMPIN
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN LABORATORIES LTD
  - VECURONIUM BROMIDE, VECURONIUM BROMIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID
- MYLAN PHARMS INC**
  - \* MYLAN PHARMACEUTICALS INC
    - ABACAVIR SULFATE, ABACAVIR SULFATE
    - ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM
    - ACITRETIN, ACITRETIN
    - ACYCLOVIR, ACYCLOVIR
    - ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
    - AMLODIPINE BESYLATE AND ATORVASTATIN CALCIUM, AMLODIPINE BESYLATE
    - AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
    - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
    - AMNESTEEM, ISOTRETINOIN
    - ARMODAFINIL, ARMODAFINIL
    - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
    - ATOVAQUONE AND PROGUANIL HYDROCHLORIDE, ATOVAQUONE
    - AVITA, TRETINOIN
    - BACLOFEN, BACLOFEN
    - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
    - CABERGOLINE, CABERGOLINE
    - CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL
    - CAPECITABINE, CAPECITABINE
    - CAPTOPRIL, CAPTOPRIL
    - CELECOXIB, CELECOXIB
    - CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
    - CIPROFLOXACIN EXTENDED RELEASE, CIPROFLOXACIN
    - CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
    - CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
    - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
    - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
    - CLOZAPINE, CLOZAPINE
    - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
    - CYTARABINE, CYTARABINE
    - DENAVIR, PENCICLOVIR
    - DESLORATADINE, DESLORATADINE
    - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
    - DESVENLAFAKINE SUCCINATE, DESVENLAFAKINE SUCCINATE
    - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
    - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
    - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
    - DICLOFENAC SODIUM, DICLOFENAC SODIUM
    - DIGOXIN, DIGOXIN
    - DISULFIRAM, DISULFIRAM
    - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
    - DOXE PIN HYDROCHLORIDE, DOXE PIN HYDROCHLORIDE
    - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
    - DOXYCYCLINE, DOXYCYCLINE
    - DUTASTERIDE, DUTASTERIDE
    - EFAVIRENZ, EFAVIRENZ
    - ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
    - ELIMITE, PERMETHRIN
    - EPLERENONE, EPLERENONE
    - EPROSARTAN MESYLATE, EPROSARTAN MESYLATE
    - ERYGEL, ERYTHRHYOMYCIN
    - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
    - ESTRADIOL, ESTRADIOL
    - ESZOPICLONE, ESZOPICLONE
    - EOCLIN, CLINDAMYCIN PHOSPHATE
    - EXEMESTANE, EXEMESTANE
    - EXTINA, KETOCONAZOLE
    - FENOFIBRATE (MICRONIZED), FENOFIBRATE
    - FENOFIBRATE, FENOFIBRATE
    - FENOFIBRIC ACID, CHOLINE FENOFIBRATE
    - FINASTERIDE, FINASTERIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN PHARMACEUTICALS INC
  - FLUOROURACIL, FLUOROURACIL
  - FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
  - FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
  - FOSAMPRENAVIR CALCIUM, FOSAMPRENAVIR CALCIUM
  - FROVATRIPTAN SUCCINATE, FROVATRIPTAN SUCCINATE
  - GABAPENTIN, GABAPENTIN
  - GLATIRAMER ACETATE, GLATIRAMER ACETATE
  - GLIPIZIDE, GLIPIZIDE
  - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IMATINIB MESYLATE, IMATINIB MESYLATE
  - INDOMETHACIN, INDOMETHACIN
  - IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - ITRACONAZOLE, ITRACONAZOLE
  - LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
  - LAMIVUDINE, LAMIVUDINE
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
  - LINEZOLID, LINEZOLID
  - LITHIUM CARBONATE, LITHIUM CARBONATE
  - LUXIQ, BETAMETHASONE VALERATE
  - MAXZIDE, HYDROCHLOROTHIAZIDE
  - MAXZIDE-25, HYDROCHLOROTHIAZIDE
  - MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - MESALAMINE, MESALAMINE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHYCLOTHIAZIDE, METHYCLOTHIAZIDE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
  - MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
  - MODAFINIL, MODAFINIL
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MORPHINE SULFATE, MORPHINE SULFATE
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
  - NABUMETONE, NABUMETONE
  - NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
  - NEVIRAPINE, NEVIRAPINE
  - OLANZAPINE, OLANZAPINE
  - OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - OLUX E, CLOBETASOL PROPIONATE
  - OLUX, CLOBETASOL PROPIONATE
  - PALIPERIDONE, PALIPERIDONE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PERPHENAZINE, PERPHENAZINE
  - PHENYTOIN, PHENYTOIN
  - PINDOLOL, PINDOLOL
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
  - PRASUGREL, PRASUGREL HYDROCHLORIDE
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
  - PREDNISONE, PREDNISONE
  - PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
  - QUININE SULFATE, QUININE SULFATE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - RASAGILINE MESYLATE, RASAGILINE MESYLATE
  - REPAGLINIDE, REPAGLINIDE
  - RILUZOLE, RILUZOLE
  - RISEDRONATE SODIUM, RISEDRONATE SODIUM
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

- \* MYLAN PHARMACEUTICALS INC
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - RUFINAMIDE, RUFINAMIDE
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - TELMISARTAN AND AMLODIPIINE, AMLODIPIINE BESYLATE
  - TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TELMISARTAN, TELMISARTAN
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TOBRAMYCIN, TOBRAMYCIN
  - TOLAZAMIDE, TOLAZAMIDE
  - TOLBUTAMIDE, TOLBUTAMIDE
  - TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - TRAVOPROST, TRAVOPROST
  - TRETINOIN, TRETINOIN
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
  - TRIAZOLAM, TRIAZOLAM
  - TRILYTE, POLYETHYLENE GLYCOL 3350
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - VALSARTAN, VALSARTAN
  - VORICONAZOLE, VORICONAZOLE
  - VUSION, MICONAZOLE NITRATE
  - ZIDOVUDINE, ZIDOVUDINE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
  - ZOLMITRIPTAN, ZOLMITRIPTAN
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
  - ZONALON, DOXEPIN HYDROCHLORIDE
  - ZOVIRAX, ACYCLOVIR
- \* MYLAN PHARMACEUTICALS INC.
  - FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
  - NIZATIDINE, NIZATIDINE
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

**MYLAN SPECIALTY LP**

- \* MYLAN SPECIALTY LP
  - ACCUNEB, ALBUTEROL SULFATE
  - AEROSPAN HFA, FLUNISOLIDE
  - ANADROL-50, OXYMETHOLONE
  - ASTELIN, AZELASTINE HYDROCHLORIDE
  - ASTEPRO, AZELASTINE HYDROCHLORIDE
  - AVC, SULFANILAMIDE
  - BUTISOL SODIUM, BUTABARBITAL SODIUM
  - CESAMET, NABILONE
  - COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
  - CORTIFOAM, HYDROCORTISONE ACETATE
  - CROMOLYN SODIUM, CROMOLYN SODIUM
  - DEMADEX, TORSEMIDE
  - DEPEN, PENICILLAMINE
  - DIPENTUM, OLSALAZINE SODIUM
  - DYMISTA, AZELASTINE HYDROCHLORIDE
  - EDLUAR, ZOLPIDEM TARTRATE
  - ELESTRIN, ESTRADIOL
  - EPIFOAM, HYDROCORTISONE ACETATE
  - EPIPEN JR., EPINEPHRINE
  - EPIPEN, EPINEPHRINE
  - FELBATOL, FELBAMATE
  - GASTROCROM, CROMOLYN SODIUM
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
  - LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
  - MUSE, ALPROSTADIL
  - PROCTOFOAM HC, HYDROCORTISONE ACETATE
  - ROWASA, MESALAMINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* M \*\*

\* MYLAN SPECIALTY LP  
 SFROWASA, MESALAMINE  
 SOMA, CARISOPRODOL

**MYLAN SPECLT**

\* MYLAN SPECIALTY LP  
 PERFOROMIST, FORMOTEROL FUMARATE

**MYLAN TECHNOLOGIES**

\* MYLAN TECHNOLOGIES INC  
 CLONIDINE, CLONIDINE  
 ESTRADIOL, ESTRADIOL  
 FENTANYL-100, FENTANYL  
 FENTANYL-12, FENTANYL  
 FENTANYL-25, FENTANYL  
 FENTANYL-37, FENTANYL  
 FENTANYL-50, FENTANYL  
 FENTANYL-62, FENTANYL  
 FENTANYL-75, FENTANYL  
 FENTANYL-87, FENTANYL  
 LIDOCAINE, LIDOCAINE  
 NITROGLYCERIN, NITROGLYCERIN  
 XULANE, ETHINYL ESTRADIOL

**MYLAN TEORANTA**

\* MYLAN TEORANTA  
 LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM

\*\* N \*\*

**NAMIGEN LLC**

\* NAMIGEN LLC  
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

**NANG KUANG PHARM CO**

\* NANG KUANG PHARMACEUTICAL CO LTD  
 LINEZOLID, LINEZOLID

**NANJING KING-FRIEND**

\* NANJING KING-FRIEND BIOCHEMICAL PHARMACEUTICAL CO LTD  
 ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE  
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE  
 CARBOPLATIN, CARBOPLATIN

**NAPO PHARMS INC**

\* NAPO PHARMACEUTICALS INC  
 FULYZAQ, CROFELEMER

**NATCO PHARMA**

\* NATCO PHARMA LTD  
 GRANisetron Hydrochloride, GRANisetron Hydrochloride

**NATCO PHARMA LTD**

\* NATCO PHARMA LIMITED  
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE  
 \* NATCO PHARMA LTD  
 ALPRAZOLAM, ALPRAZOLAM  
 ANASTROZOLE, ANASTROZOLE  
 ARMODAFINIL, ARMODAFINIL  
 AZACITIDINE, AZACITIDINE  
 CARISOPRODOL, CARISOPRODOL  
 CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE  
 GLYCOPYRROLATE, GLYCOPYRROLATE  
 LANSOPRAZOLE, LANSOPRAZOLE  
 LANSOPRAZOLE, LANSOPRAZOLE (OTC)  
 LANTHANUM CARBONATE, LANTHANUM CARBONATE  
 LETROZOLE, LETROZOLE  
 ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
 OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE  
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

**NAVINTA LLC**

\* NAVINTA LLC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* N \*\*

- \* NAVINTA LLC
  - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
  - FAMOTIDINE, FAMOTIDINE
  - FOMEPIZOLE, FOMEPIZOLE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - INDOMETHACIN SODIUM, INDOMETHACIN SODIUM
  - METHOCARBAMOL, METHOCARBAMOL
  - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
  - RIBAVIRIN, RIBAVIRIN
  - ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE
  - SODIUM PHENYLACETATE AND SODIUM BENZOATE, SODIUM BENZOATE

**NCM USA BRONX LLC**

- \* NCM USA BRONX LLC
  - AMMONIA N 13, AMMONIA N-13
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**NEOS THERAP INC**

- \* NEOS THERAPEUTICS INC
  - HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX

**NEOS THERAPS**

- \* NEOS THERAPEUTICS
  - ADZENYS XR-ODT, AMPHETAMINE

**NEOS THERAPS INC**

- \* NEOS THERAPEUTICS INC
  - ADZENYS ER, AMPHETAMINE
  - COTEMPLA XR-ODT, METHYLPHENIDATE

**NEPHRON**

- \* NEPHRON CORP
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
- \* NEPHRON PHARMACEUTICALS CORP
  - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE

**NESHER PHARMS**

- \* NESHER PHARMACEUTICALS USA LLC
  - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
  - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
  - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
  - MICRO-K 10, POTASSIUM CHLORIDE
  - MICRO-K, POTASSIUM CHLORIDE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - NYSTATIN, NYSTATIN
  - OSELTAMIVIR PHOSPHATE, OSELTAMIVIR PHOSPHATE
  - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

**NEUROCRINE**

- \* NEUROCRINE BIOSCIENCES INC
  - INGREZZA, VALBENAZINE TOSYLATE

**NEW HAVEN PHARMS**

- \* NEW HAVEN PHARMACEUTICALS INC
  - DURLAZA, ASPIRIN

**NEW RIVER**

- \* NEW RIVER PHARMACEUTICALS INC
  - PROFERDEX, IRON DEXTRAN

**NEWGEN PHARMS LLC**

- \* NEWGEN PHARMACEUTICALS LLC
  - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE

**NEXGEN PHARMA**

- \* NEXGEN PHARMA INC
  - BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
  - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
  - CHENODIOL, CHENODIOL
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - MECAMYLAMINE HYDROCHLORIDE, MECAMYLMINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* N \*\*

- \* NEXGEN PHARMA INC  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)

**NEXGEN PHARMA INC**

- \* NEXGEN PHARMA INC  
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350

**NEXTWAVE PHARMS**

- \* NEXTWAVE PHARMACEUTICALS INC  
QUILLIVANT XR, METHYLPHENIDATE HYDROCHLORIDE

**NEXUS PHARMS**

- \* NEXUS PHARMACEUTICALS INC  
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE

**NIAGARA PHARMS**

- \* NIAGARA PHARMACEUTICALS INC  
PUR-WASH, PURIFIED WATER (OTC)

**NODEN PHARMA**

- \* NODEN PHARMA DAC  
TEKTURNA HCT, ALISKIREN HEMIFUMARATE  
TEKTURNA, ALISKIREN HEMIFUMARATE

**NORTEC DEV ASSOC**

- \* NORTEC DEVELOPMENT ASSOC INC  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

**NORTHSTAR HLTHCARE**

- \* NORTHSTAR HEALTHCARE HOLDINGS LTD  
ALLOPURINOL, ALLOPURINOL  
BACLOFEN, BACLOFEN  
GEMFIBROZIL, GEMFIBROZIL  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

**NORTON WATERFORD**

- \* NORTON WATERFORD LTD  
QVAR REDIHALER, BECLOMETHASONE DIPROPIONATE

**NOSTRUM LABS INC**

- \* NOSTRUMLABORATORIES INC  
ACETAZOLAMIDE, ACETAZOLAMIDE  
CALCIUM ACETATE, CALCIUM ACETATE  
CARISOPRODOL, CARISOPRODOL  
DAPSONE, DAPSONE  
ELIXOPHYLLIN, THEOPHYLLINE  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
MORPHINE SULFATE, MORPHINE SULFATE  
NITROFURANTOIN, NITROFURANTOIN  
PINDOLOL, PINDOLOL  
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
VENLAFAKINE HYDROCHLORIDE, VENLAFAKINE HYDROCHLORIDE

**NOSTRUMLPHARMS LLC**

- \* NOSTRUMLPHARMA INC  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
THEOCHRON, THEOPHYLLINE

**NOVA LABS LTD**

- \* NOVA LABORATORIES LTD  
PURIXAN, MERCAPTOPURINE

**NOVARTIS**

- \* NOVARTIS CONSUMER HEALTH INC  
LAMISIL AT, TERBINAFINE (OTC)  
LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)  
THRIVE, NICOTINE POLACRILEX (OTC)
- \* NOVARTIS PHARMACEUTICALS CORP  
AFINITOR, EVEROLIMUS

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* N \*\*

\* NOVARTIS PHARMACEUTICALS CORP  
COARTEM, ARTEMETHER  
DESFERAL, DEFEROXAMINE MESYLATE  
DIOVAN HCT, HYDROCHLOROTHIAZIDE  
DIOVAN, VALSARTAN  
EXELON, RIVASTIGMINE  
EXELON, RIVASTIGMINE TARTRATE  
EXFORGE HCT, AMLODIPINE BESYLATE  
EXFORGE, AMLODIPINE BESYLATE  
EXJADE, DEFERASIROX  
FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE  
FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE  
GILENYA, FINGOLIMOD  
GLEEVEC, IMATINIB MESYLATE  
LAMISIL, TERBINAFINE HYDROCHLORIDE  
LESCOL XL, FLUVASTATIN SODIUM  
LOPRESSOR, METOPROLOL TARTRATE  
LOTREL, AMLODIPINE BESYLATE  
MYFORTIC, MYCOPHENOLIC ACID  
NEORAL, CYCLOSPORINE  
RECLAST, ZOLEDRONIC ACID  
RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE  
RITALIN, METHYLPHENIDATE HYDROCHLORIDE  
RITALIN-SR, METHYLPHENIDATE HYDROCHLORIDE  
SANDIMMUNE, CYCLOSPORINE  
SANDOSTATIN LAR, OCTREOTIDE ACETATE  
SANDOSTATIN, OCTREOTIDE ACETATE  
SIGNIFOR, PASIREOTIDE DIASPARTATE  
STARLIX, NATEGLINIDE  
TASIGNA, NILOTINIB HYDROCHLORIDE MONOHYDRATE  
TEGRETOL, CARBAMAZEPINE  
TEGRETOL-XR, CARBAMAZEPINE  
TOBI PODHALER, TOBRAMYCIN  
TRILEPTAL, OXCARBAZEPINE  
VIVELLE-DOT, ESTRADIOL  
VOLTAREN, DICLOFENAC SODIUM  
ZOMETA, ZOLEDRONIC ACID  
ZORTRESS, EVEROLIMUS

**NOVARTIS PHARM**

\* NOVARTIS PHARMACEUTICAL CORP  
AFINITOR DISPERZ, EVEROLIMUS

**NOVARTIS PHARMS**

\* NOVARTIS PHARMACEUTICALS CORP  
FEMARA, LETROZOLE  
TOBI, TOBRAMYCIN

**NOVARTIS PHARMS CORP**

\* NOVARTIS PHARMACEUTICALS CORP  
ALCAINE, PROPARACAIN HYDROCHLORIDE  
ALOMIDE, LODOXAMIDE TROMETHAMINE  
ARGATROBAN, ARGATROBAN  
ARRANON, NELARABINE  
AZOPT, BRINZOLAMIDE  
BETOPTIC S, BETAXOLOL HYDROCHLORIDE  
CILOXAN, CIPROFLOXACIN HYDROCHLORIDE  
CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE  
CIPRODEX, CIPROFLOXACIN  
CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE  
CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE  
DUREZOL, DIFLUPREDNATE  
EMADINE, EMEDASTINE DIFUMARATE  
ENTRESTO, SACUBITRIL  
FARYDAK, PANOBINOSTAT LACTATE  
FLAREX, FLUOROMETHOLONE ACETATE  
FLUORESCITE, FLUORESCIN SODIUM  
HYCAMTIN, TOPOTECAN HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* N \*\*

\* NOVARTIS PHARMACEUTICALS CORP  
 ILEVRO, NEPAFENAC  
 IOPIDINE, APRACLONIDINE HYDROCHLORIDE  
 ISOPTO ATROPINE, ATROPINE SULFATE  
 ISOPTO CARPINE, PILOCARPINE HYDROCHLORIDE  
 JADENU SPRINKLE, DEFERASIROX  
 JADENU, DEFERASIROX  
 KISQALI FEMARA CO-PACK (COPACKAGED), LETROZOLE  
 KISQALI, RIBOCICLIB SUCCINATE  
 MAXIDEX, DEXAMETHASONE  
 MAXITROL, DEXAMETHASONE  
 MEKINIST, TRAMETINIB DIMETHYL SULFOXIDE  
 MOXEZA, MOXIFLOXACIN HYDROCHLORIDE  
 MYDRIACYL, TROPICAMIDE  
 NATACYN, NATAMYCIN  
 NEVANAC, NEPAFENAC  
 OMNIPRED, PREDNISOLONE ACETATE  
 PATADAY, OLOPATADINE HYDROCHLORIDE  
 PATANASE, OLOPATADINE HYDROCHLORIDE  
 PATANOL, OLOPATADINE HYDROCHLORIDE  
 PAZEO, OLOPATADINE HYDROCHLORIDE  
 PROMACTA, ELTROMBOPAG OLAMINE  
 RYDAPT, MIDOSTAURIN  
 SIGNIFOR LAR, PASIREOTIDE PAMOATE  
 SIMBRINZA, BRIMONIDINE TARTRATE  
 TAFINLAR, DABRAFENIB MESYLATE  
 TETRACAINE HYDROCHLORIDE, TETRACAINE HYDROCHLORIDE  
 TOBRADEX ST, DEXAMETHASONE  
 TOBRADEX, DEXAMETHASONE  
 TOBREX, TOBRAMYCIN  
 TRAVATAN Z, TRAVOPROST  
 TRIESENCE, TRIAMCINOLONE ACETONIDE  
 TYKERB, LAPATINIB DITOSYLATE  
 VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE  
 VOTRIENT, PAZOPANIB HYDROCHLORIDE  
 XTORO, FINAFLOXACIN  
 ZOFRAN ODT, ONDANSETRON  
 ZOFRAN, ONDANSETRON HYDROCHLORIDE  
 ZYKADIA, CERITINIB

**NOVAST LABS**

\* NOVAST LABORATORIES CHINA LTD  
 NORETHINDRONE, NORETHINDRONE

**NOVAST LABS LTD**

\* NOVAST LABORATORIES LTD  
 ACETAZOLAMIDE, ACETAZOLAMIDE  
 CARISOPRODOL AND ASPIRIN, ASPIRIN  
 DASETTA 1/35, ETHINYL ESTRADIOL  
 DASETTA 7/7/7, ETHINYL ESTRADIOL  
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL  
 ELINEST, ETHINYL ESTRADIOL  
 FALMINA, ETHINYL ESTRADIOL  
 HER STYLE, LEVONORGESTREL (OTC)  
 INDOMETHACIN, INDOMETHACIN  
 LARIN 1.5/30, ETHINYL ESTRADIOL  
 LARIN 1/20, ETHINYL ESTRADIOL  
 LARIN 24 FE, ETHINYL ESTRADIOL  
 LARIN FE 1.5/30, ETHINYL ESTRADIOL  
 LARIN FE 1/20, ETHINYL ESTRADIOL  
 LERIBANE, ETHINYL ESTRADIOL  
 LEVONEST, ETHINYL ESTRADIOL  
 MAFENIDE ACETATE, MAFENIDE ACETATE  
 MELAMISA, DROSPIRENONE  
 MONO-LINYAH, ETHINYL ESTRADIOL  
 NIFEDIPINE, NIFEDIPINE  
 NORETHINDRONE, NORETHINDRONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* N \*\*

\* NOVAST LABORATORIES LTD  
 PHILITH, ETHINYL ESTRADIOL  
 PIMTREA, DESOGESTREL  
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE  
 SETLAKIN, ETHINYL ESTRADIOL  
 TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE  
 TRI-LINYAH, ETHINYL ESTRADIOL  
 WERA, ETHINYL ESTRADIOL  
 YAELA, DROSPIRENONE

**NOVATECH SA**

\* NOVATECH SA  
 STERITALC, TALC

**NOVEL LABS INC**

\* NOVEL LABORATORIES INC  
 AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE  
 CALCIPOTRIENE, CALCIPOTRIENE  
 CARBIDOPA, CARBIDOPA  
 CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE  
 CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
 CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
 DESOXIMETASONE, DESOXIMETASONE  
 DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE  
 DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
 DICLOFENAC SODIUM, DICLOFENAC SODIUM  
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
 FAMOTIDINE, FAMOTIDINE  
 FLUCYTOSINE, FLUCYTOSINE  
 FLUOCINONIDE, FLUOCINONIDE  
 HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE  
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE  
 LEVONORGESTREL, LEVONORGESTREL (OTC)  
 LINEZOLID, LINEZOLID  
 METHYLERGONOVINE MALEATE, METHYLERGONOVINE MALEATE  
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
 MISOPROSTOL, MISOPROSTOL  
 MORPHINE SULFATE, MORPHINE SULFATE  
 MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE  
 NITROFURANTOIN, NITROFURANTOIN  
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
 PEG 3350 AND ELECTROLYTES, POLYETHYLENE GLYCOL 3350  
 PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL  
 PEG-3350, SODIUM CHLORIDE, SODIUM BICARBONATE, POTASSIUM CHLORIDE AND BISACODYL,  
 PHENELZINE SULFATE, PHENELZINE SULFATE  
 POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)  
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
 SODIUM SULFATE, POTASSIUM SULFATE AND MAGNESIUM SULFATE, MAGNESIUM SULFATE  
 TEMAZEPAM, TEMAZEPAM  
 TINIDAZOLE, TINIDAZOLE  
 TRIMETHOPRIM, TRIMETHOPRIM  
 VORICONAZOLE, VORICONAZOLE  
 ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**NOVEN**

\* NOVEN PHARMACEUTICALS INC  
 MINIVELLE, ESTRADIOL

**NOVEN PHARMS INC**

\* NOVEN PHARMACEUTICALS INC  
 COMBIPATCH, ESTRADIOL  
 DAYTRAN, METHYLPHENIDATE

**NOVITIUM PHARMA**

\* NOVITIUM PHARMA LLC  
 DAPSONE, DAPSONE  
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* N \*\*

**NOVO NORDISK**

- \* NOVO NORDISK PHARMACEUTICALS INC  
GLUCAGEN, GLUCAGON HYDROCHLORIDE RECOMBINANT
- NOVO NORDISK INC**
- \* NOVO NORDISK INC  
FIASP FLEXTOUCH, INSULIN ASPART  
FIASP, INSULIN ASPART  
LEVEMIR FLEXTOUCH, INSULIN DETEMIR RECOMBINANT  
LEVEMIR, INSULIN DETEMIR RECOMBINANT  
NORDITROPIN FLEXPRO, SOMATROPIN RECOMBINANT  
NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)  
NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)  
NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)  
NOVOLOG FLEXPEN, INSULIN ASPART RECOMBINANT  
NOVOLOG FLEXTOUCH, INSULIN ASPART RECOMBINANT  
NOVOLOG MIX 70/30 FLEXPEN, INSULIN ASPART PROTAMINE RECOMBINANT  
NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT  
NOVOLOG PENFILL, INSULIN ASPART RECOMBINANT  
NOVOLOG, INSULIN ASPART RECOMBINANT  
OZEMPIC, SEMAGLUTIDE  
RYZODEG 70/30, INSULIN ASPART  
SAXENDA, LIRAGLUTIDE RECOMBINANT  
TRESIBA, INSULIN DEGLUDEC  
VAGIFEM, ESTRADIOL  
VICTOZA, LIRAGLUTIDE RECOMBINANT  
XULTOPHY 100/3.6, INSULIN DEGLUDEC

**NPS PHARMS INC**

- \* NPS PHARMACEUTICALS INC  
GATTEX KIT, TEDUGLUTIDE RECOMBINANT

**NU PHARM**

- \* NU PHARM INC  
DIVALPROEX SODIUM, DIVALPROEX SODIUM

**NUVO PHARM**

- \* NUVO PHARMACEUTICAL INC  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE

**NUVO PHARM INC**

- \* NUVO PHARMACEUTICAL INC  
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
FOLIC ACID, FOLIC ACID  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE

**NXDC**

- \* NX DEVELOPMENT CORP  
GLEOLAN, AMINOLEVULINIC ACID HYDROCHLORIDE

**NYCOMED US**

- \* NYCOMED US INC  
TERCONAZOLE, TERCONAZOLE

\*\* O \*\*

**OAK PHARMS**

- \* OAK PHARMACEUTICALS INC  
NEMBUTAL SODIUM, PENTOBARBITAL SODIUM  
XYLOCAINE, LIDOCAINE HYDROCHLORIDE

**OAK PHARMS AKORN**

- \* OAK PHARMACEUTICALS INC SUB AKORN INC  
COGETIN, BENZTROPINE MESYLATE  
DIURIL, CHLOROTHIAZIDE SODIUM

**OAK PHARMS INC**

- \* OAK PHARMACEUTICALS INC  
ZIOPTAN, TAFLUPROST
- \* OAK PHARMACEUTICALS INC SUBSIDIARY OF AKORN INC  
AZASITE, AZITHROMYCIN  
BETIMOL, TIMOLOL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* O \*\*

- \* OAK PHARMACEUTICALS INC SUBSIDIARY OF AKORN INC  
COSOPT PF, DORZOLAMIDE HYDROCHLORIDE  
COSOPT, DORZOLAMIDE HYDROCHLORIDE  
XOPENEX, LEVALBUTEROL HYDROCHLORIDE

**OC PHARMA**

- \* OC PHARMA LLC  
LEVONORGESTREL, LEVONORGESTREL (OTC)  
NORGESTIMATE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL

**ODYSSEY PHARMS**

- \* ODYSSEY PHARMACEUTICALS INC  
ANTABUSE, DISULFIRAM  
NYSTATIN, NYSTATIN  
SURMONTIL, TRIMIPRAMINE MALEATE  
URECHOLINE, BETHANECHOL CHLORIDE  
VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE

**OHM**

- \* OHM CORP  
IBUPROFEN, IBUPROFEN (OTC)

**OHM LABS**

- \* OHM LABORATORIES INC  
ACETAMINOPHEN, ACETAMINOPHEN (OTC)  
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)  
IBUPROHM, IBUPROFEN (OTC)  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

**OHM LABS INC**

- \* OHM LABORATORIES INC  
EZETIMIBE, EZETIMIBE  
VALSARTAN, VALSARTAN

**OLTA PHARMS**

- \* OLTA PHARMACEUTICALS CORP  
LINDANE, LINDANE

**OMEROS**

- \* OMEROS CORP  
OMIDRIA, KETOROLAC TROMETHAMINE

**ONY**

- \* ONY INC  
INFASURF PRESERVATIVE FREE, CALFACTANT

**ONYX THERAP**

- \* ONYX THERAPEUTICS INC A WHOLLY OWNED SUB OF AMGEN INC  
KYPROLIS, CARFILZOMIB

**OPKO IRELAND GLOBAL**

- \* OPKO IRELAND GLOBAL HOLDINGS LTD  
RAYALDEE, CALCIFEDIOL

**OPTNOSE US**

- \* OPTNOSE US INC  
XHANCE, FLUTICASONE PROPIONATE

**ORAPHARMA**

- \* ORAPHARMA INC  
ARESTIN, MINOCYCLINE HYDROCHLORIDE

**ORCHID HLTHCARE**

- \* ORCHID HEALTHCARE  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
ARIPIPRAZOLE, ARIPIPRAZOLE  
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE  
CEFDINIR, CEFDINIR  
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL  
CEFPROZIL, CEFPROZIL  
CEFUROXIME AXETIL, CEFUROXIME AXETIL  
CEPHALEXIN, CEPHALEXIN  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)  
DESLORATADINE, DESLORATADINE  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
ESZOPICLONE, ESZOPICLONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* O \*\*

- \* ORCHID HEALTHCARE
  - FELODIPINE, FELODIPINE
  - GEMIFLOXACIN MESYLATE, GEMIFLOXACIN MESYLATE
  - GRANisetron Hydrochloride, GRANisetron Hydrochloride
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN, LEVOFLOXACIN
  - MODAFINIL, MODAFINIL
  - NARatriptan, NARatriptan Hydrochloride
  - OLANzapine, OLANzapine
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - RASAGILINE MESYLATE, RASAGILINE MESYLATE
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - VENLAFAxINE HYDROCHLORIDE, VENLAFAxINE HYDROCHLORIDE
  - ZALEPLON, ZALEPLON

**OREXIGEN**

- \* OREXIGEN THERAPEUTICS INC
  - CONTRAVE, BUPROPION HYDROCHLORIDE

**OREXO US INC**

- \* OREXO US INC
  - ZUBSOLV, BUPRENORPHINE HYDROCHLORIDE

**ORGANON SUB MERCK**

- \* ORGANON USA INC A SUB OF MERCK AND CO INC
  - BRIDION, SUGAMMADEX SODIUM
  - NUVARING, ETHINYl ESTRADIOL

**ORGANON USA INC**

- \* ORGANON USA INC
  - DESOGEN, DESOGESTREL
  - FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
  - GANIRELIX ACETATE, GANIRELIX ACETATE
  - NEXPLANON, ETONOGESTREL
  - PREGNYL, GONADOTROPIN, CHORIONIC
  - REMERON SOLTAB, MIRTAZAPINE
  - REMERON, MIRTAZAPINE

**ORIENT PHARMA CO LTD**

- \* ORIENT PHARMA CO LTD
  - CARISOPRODOL, CARISOPRODOL
  - MIGLITOL, MIGLITOL
  - PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

**ORION CORP ORION**

- \* ORION CORP ORION PHARMA
  - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE

**ORION PHARMA**

- \* ORION PHARMA
  - COMTAN, ENTACAPONE
  - STALEVO 100, CARBIDOPA
  - STALEVO 125, CARBIDOPA
  - STALEVO 150, CARBIDOPA
  - STALEVO 200, CARBIDOPA
  - STALEVO 50, CARBIDOPA
  - STALEVO 75, CARBIDOPA

**ORIT LABS LLC**

- \* ORIT LABORATORIES LLC
  - BENZONATATE, BENZONATATE
  - BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
  - CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE
  - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
  - ERGOCALCIFEROL, ERGOCALCIFEROL
  - LEVETIRACETAM, LEVETIRACETAM
  - METRONIDAZOLE, METRONIDAZOLE

**ORPHAN EUROPE**

- \* ORPHAN EUROPE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* O \*\*

- \* ORPHAN EUROPE  
CARBAGLU, CARGLUMIC ACID
- \* ORPHAN EUROPE SARL  
CYSTADANE, BETAINE HYDROCHLORIDE

**OSI PHARMS**

- \* OSI PHARMACEUTICALS INC  
TARCEVA, ERLOTINIB HYDROCHLORIDE

**OSMOTICA**

- \* OSMOTICA KERESKEDELMI ES SZOLGALTATO KFT  
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE  
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE

**OSMOTICA PHARM**

- \* OSMOTICA PHARMACEUTICAL CORP  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

**OSMOTICA PHARM CORP**

- \* OSMOTICA PHARMACEUTICAL CORP  
KHEDEZLA, DESVENLAFAXINE

**OSMOTICA PHARM US**

- \* OSMOTICA PHARMACEUTICAL US LLC  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
NIFEDIPINE, NIFEDIPINE  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

**OTONOMY INC**

- \* OTONOMY INC  
OTIPRIO, CIPROFLOXACIN

**OTSUKA**

- \* OTSUKA PHARMACEUTICAL CO LTD  
ABILIFY, ARIPIPRAZOLE

**OTSUKA AMERICA PHARM**

- \* OTSUKA AMERICA PHARMACEUTICAL INC  
SAMSCA, TOLVAPTAN

**OTSUKA PHARM**

- \* OTSUKA PHARMACEUTICAL CO LTD  
BUSULFEX, BUSULFAN

**OTSUKA PHARM CO LTD**

- \* OTSUKA PHARMACEUTICAL CO LTD  
ABILIFY MAINTENA KIT, ARIPIPRAZOLE  
ABILIFY MYCITE KIT, ARIPIPRAZOLE  
DACOGEN, DECITABINE  
REXULTI, BREXIPIPRAZOLE

**OUTLOOK PHARMS**

- \* OUTLOOK PHARMACEUTICALS INC  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

**OXFORD PHARMS**

- \* OXFORD PHARMACEUTICALS LLC  
BACLOFEN, BACLOFEN  
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE  
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE  
CARISOPRODOL AND ASPIRIN, ASPIRIN  
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE  
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE  
RIMACTANE, RIFAMPIN  
RISPERIDONE, RISPERIDONE  
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE  
SIMVASTATIN, SIMVASTATIN  
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

\*\* P \*\*

**P AND L DEV LLC**

- \* P AND L DEVELOPMENT LLC DBA PLD DEVELOPMENTS LLC  
IBUPROFEN, IBUPROFEN (OTC)

**PACIFIC PHARMA**

- \* PACIFIC PHARMA

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PACIFIC PHARMA  
TIMOLOL MALEATE, TIMOLOL MALEATE
- \* PACIFIC PHARMA INC  
TIMOLOL MALEATE, TIMOLOL MALEATE

**PACIRA PHARMS INC**

- \* PACIRA PHARMACEUTICALS INC  
DEPOCYT, CYTARABINE  
EXPAREL, BUPIVACAINE

**PADDOCK LLC**

- \* PADDOCK LABORATORIES LLC  
ATOVAQUONE, ATOVAQUONE  
BROMFENAC SODIUM, BROMFENAC SODIUM  
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE  
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN  
CALCIUM ACETATE, CALCIUM ACETATE  
CICLOPIROX, CICLOPIROX  
CLINDA-DERM, CLINDAMYCIN PHOSPHATE  
CLINDAMYCIN PALMITATE HYDROCHLORIDE, CLINDAMYCIN PALMITATE HYDROCHLORIDE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLOTRIMAZOLE, CLOTRIMAZOLE  
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM  
COLOCORT, HYDROCORTISONE  
COMPRO, PROCHLORPERAZINE  
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE  
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE  
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE  
HYDROCODONE BITARTRATE AND PSEUDOEPHEDRINE HYDROCHLORIDE, HYDROCODONE BITARTRATE  
HYDROCODONE BITARTRATE, CHLORPHENIRAMINE MALEATE AND PSEUDOEPHEDRINE HYDROCHLORIDE ,  
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE  
KIONEX, SODIUM POLYSTYRENE SULFONATE  
LAX-LYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350  
MIDAMOR, AMILORIDE HYDROCHLORIDE  
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE  
MORPHINE SULFATE, MORPHINE SULFATE  
NARatriptan, NARatriptan HYDROCHLORIDE  
NYSTOP, NYSTATIN  
PODOFILOX, PODOFILOX  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE  
REPAGLINIDE, REPAGLINIDE  
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE  
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE  
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE  
TROSPiUM CHLORIDE, TROSPiUM CHLORIDE

**PANACEA BIOTEC LTD**

- \* PANACEA BIOTEC LTD  
PRASUGREL, PRASUGREL HYDROCHLORIDE  
RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
TACROLIMUS, TACROLIMUS

**PAR FORM**

- \* PAR FORMULATIONS PRIVATE LTD  
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
MAFENIDE ACETATE, MAFENIDE ACETATE  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**PAR PHARM**

- \* PAR PHARMACEUTICAL  
OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE  
PROPafenone HYDROCHLORIDE, PROPafenone HYDROCHLORIDE  
TESTOSTERONE, TESTOSTERONE
- \* PAR PHARMACEUTICAL INC  
ALPRAZOLAM, ALPRAZOLAM  
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE  
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PAR PHARMACEUTICAL INC
  - AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
  - CABERGOLINE, CABERGOLINE
  - CALCITONIN-SALMON, CALCITONIN SALMON
  - CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
  - CHOLESTYRAMINE, CHOLESTYRAMINE
  - CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
  - CLONAZEPAM, CLONAZEPAM
  - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
  - DEXAMETHASONE, DEXAMETHASONE
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
  - DOXYCYCLINE, DOXYCYCLINE
  - ESTAZOLAM, ESTAZOLAM
  - FENTANYL CITRATE, FENTANYL CITRATE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUTAMIDE, FLUTAMIDE
  - GLIPIZIDE, GLIPIZIDE
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - HYDROXYUREA, HYDROXYUREA
  - IBUPROFEN, IBUPROFEN (OTC)
  - IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
  - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
  - LAMOTRIGINE, LAMOTRIGINE
  - MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
  - MEGESTROL ACETATE, MEGESTROL ACETATE
  - METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
  - METRONIDAZOLE, METRONIDAZOLE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MINOXIDIL, MINOXIDIL
  - NATEGLINIDE, NATEGLINIDE
  - NIFEDIPINE, NIFEDIPINE
  - OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - OLANZAPINE, OLANZAPINE
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
  - OXANDROLONE, OXANDROLONE
  - PIMOZIDE, PIMOZIDE
  - POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RISPERIDONE, RISPERIDONE
  - SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TORSEMIDE, TORSEMIDE
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRANLYLCYPROMINE SULFATE, TRANLYLCYPROMINE SULFATE
  - TRAVOPROST, TRAVOPROST
  - URSODIOL, URSDIOL
  - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

**PAR PHARM INC**

- \* PAR PHARMACEUTICAL INC
  - ACCOLATE, ZAFIRLUKAST
  - AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
  - ANTIZOL, FOMEPIZOLE
  - ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
  - BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
  - DEXLANSOPRAZOLE, DEXLANSOPRAZOLE
  - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
  - DOXYLAMINE SUCCINATE AND PYRIDOXINE HYDROCHLORIDE, DOXYLAMINE SUCCINATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PAR PHARMACEUTICAL INC
  - ENTECAVIR, ENTECAVIR
  - ETHACRYNIC ACID, ETHACRYNIC ACID
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - ITRACONAZOLE, ITRACONAZOLE
  - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
  - OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
  - PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
  - PRAZIQUANTEL, PRAZIQUANTEL
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TOLCAPONE, TOLCAPONE
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - VIGABATRIN, VIGABATRIN
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**PAR STERILE PRODUCTS**

- \* PAR STERILE PRODUCTS LLC
  - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
  - ADRENALIN, EPINEPHRINE
  - ARGATROBAN, ARGATROBAN
  - BREVITAL SODIUM, METHOHEXITAL SODIUM
  - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - COLY-MYCIN M, COLISTIMETHATE SODIUM
  - CORPHEDRA, EPHEDRINE SULFATE
  - DANTRIUM, DANTROLENE SODIUM
  - DELESTROGEN, ESTRADIOL VALERATE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
  - ETOMIDATE, ETOMIDATE
  - FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
  - GANCICLOVIR, GANCICLOVIR SODIUM
  - KETALAR, KETAMINE HYDROCHLORIDE
  - LEVOOTHYROXINE SODIUM, LEVOOTHYROXINE SODIUM
  - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
  - MEROPENEM, MEROPENEM
  - MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
  - NEOSTIGMINE METHYLSULFATE, NEOSTIGMINE METHYLSULFATE
  - PITOCIN, OXYTOCIN
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
  - TRIOSTAT, LIOTHYRONINE SODIUM
  - VASOSTRICT, VASOPRESSIN

**PARAGON BIOTECK**

- \* PARAGON BIOTECK INC
  - PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE

**PARAPRO LLC**

- \* PARAPRO LLC
  - NATROBA, SPINOSAD

**PARKE DAVIS**

- \* PARKE DAVIS DIV WARNER LAMBERT CO
  - CELONTIN, MENTSUXIMIDE
  - CEREBYX, FOSPHENYTOIN SODIUM
  - DILANTIN-125, PHENYTOIN
  - NARDIL, PHENELZINE SULFATE
  - NEURONTIN, GABAPENTIN
  - ZARONTIN, ETHOSUXIMIDE

**PARKE-DAVIS**

- \* PARKE-DAVIS DIVISION OF PFIZER INC
  - DILANTIN, PHENYTOIN SODIUM
  - ZARONTIN, ETHOSUXIMIDE

**PERNIX IRELAND LTD**

- \* PERNIX IRELAND LTD
  - TREXIMET, NAPROXEN SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

**PERNIX IRELAND PAIN**

- \* PERNIX IRELAND PAIN LIMITED  
ZOHYDRO ER, HYDROCODONE BITARTRATE

**PERNIX THERAPS LLC**

- \* PERNIX THERAPEUTICS LLC  
SILENOR, DOXEPIN HYDROCHLORIDE

**PERRIGO**

- \* PERRIGO CO  
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)

**PERRIGO CO**

- \* PERRIGO CO OF TENNESSEE INC  
CICLOPIROX, CICLOPIROX  
CLINDETS, CLINDAMYCIN PHOSPHATE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
ERYTHROMYCIN, ERYTHROMYCIN  
STIE-CORT, HYDROCORTISONE

**PERRIGO CO TENNESSEE**

- \* PERRIGO CO TENNESSEE INC  
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC  
BACITRACIN, BACITRACIN  
BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN  
ERYTHROMYCIN, ERYTHROMYCIN  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC  
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE  
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM

**PERRIGO ISRAEL**

- \* PERRIGO ISRAEL PHARMACEUTICALS LTD  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
DESOXIMETASONE, DESOXIMETASONE  
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE  
FLUOCINONIDE, FLUOCINONIDE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
GYNIAZOLE-1, BUTOCONAZOLE NITRATE  
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE  
IMIQUIMOD, IMIQUIMOD  
KETOCONAZOLE, KETOCONAZOLE  
MESALAMINE, MESALAMINE  
MINOXIDIL, MINOXIDIL (OTC)  
MOMETASONE FUROATE, MOMETASONE FUROATE  
NITROGLYCERIN, NITROGLYCERIN  
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE  
TESTOSTERONE, TESTOSTERONE  
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE (OTC)

**PERRIGO NEW YORK**

- \* PERRIGO NEW YORK INC  
ACETAMINOPHEN, ACETAMINOPHEN (OTC)  
AMMONIUM LACTATE, AMMONIUM LACTATE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CENTANY, MUPIROCIN  
CICLOPIROX, CICLOPIROX  
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
DESONIDE, DESONIDE  
DESOXIMETASONE, DESOXIMETASONE  
ECONAZOLE NITRATE, ECONAZOLE NITRATE  
ERYTHROMYCIN, ERYTHROMYCIN  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE  
HYDROCORTISONE, HYDROCORTISONE  
KETOCONAZOLE, KETOCONAZOLE  
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PERRIGO NEW YORK INC
  - MOMETASONE FUROATE, MOMETASONE FUROATE
  - MUPIROCIN, MUPIROCIN
  - NYSTATIN, NYSTATIN
  - PERMETHRIN, PERMETHRIN
  - PERMETHRIN, PERMETHRIN (OTC)
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - SELENIUM SULFIDE, SELENIUM SULFIDE
  - TERCONAZOLE, TERCONAZOLE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**PERRIGO PHARMA INTL**

- \* PERRIGO PHARMA INTERNATIONAL DAC
  - CLINDESSE, CLINDAMYCIN PHOSPHATE
  - ENTOCORT EC, BUDESONIDE
  - EVAMIST, ESTRADIOL
  - LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
  - LORATADINE, LORATADINE (OTC)
  - PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
  - TRETINOIN, TRETINOIN

**PERRIGO PHARMS CO**

- \* PERRIGO PHARMACEUTICALS CO
  - SCOPOLAMINE, SCOPOLAMINE

**PERRIGO R AND D**

- \* PERRIGO R AND D CO
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
  - DESLORATADINE, DESLORATADINE
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM (OTC)
  - FAMOTIDINE, CALCIUM CARBONATE, AND MAGNESIUM HYDROXIDE, CALCIUM CARBONATE (OTC)
  - FAMOTIDINE, FAMOTIDINE
  - FAMOTIDINE, FAMOTIDINE (OTC)
  - GUAIFENESIN, GUAIFENESIN (OTC)
  - IBUPROFEN AND DIPHENHYDRAMINE CITRATE, DIPHENHYDRAMINE CITRATE (OTC)
  - IBUPROFEN AND PHENYLEPHRINE HYDROCHLORIDE, IBUPROFEN (OTC)
  - IBUPROFEN SODIUM, IBUPROFEN SODIUM (OTC)
  - IBUPROFEN, IBUPROFEN
  - IBUPROFEN, IBUPROFEN (OTC)
  - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
  - LEVONORGESTREL, LEVONORGESTREL
  - LEVONORGESTREL, LEVONORGESTREL (OTC)
  - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
  - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - NAPROXEN, NAPROXEN
  - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE (OTC)
  - OMEPRAZOLE MAGNESIUM, OMEPRAZOLE MAGNESIUM (OTC)
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)

**PERRIGO UK FINCO**

- \* PERRIGO UK FINCO LTD PARTNERSHIP
  - BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - FLURANDRENOLIDE, FLURANDRENOLIDE
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - TESTOSTERONE, TESTOSTERONE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**PETNET**

- \* PETNET SOLUTIONS INC
  - AMMONIA N 13, AMMONIA N-13
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT****\*\* P \*\***

\* PETNET SOLUTIONS INC  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**PF PRISM CV**

\* PF PRISM CV  
BOSULIF, BOSUTINIB MONOHYDRATE  
INLYTA, AXITINIB  
LYRICA, PREGABALIN  
RAPAMUNE, SIROLIMUS  
TORISEL, TEMSIROLIMUS  
TYGACIL, TIGECYCLINE  
VFEND, VORICONAZOLE  
XALKORI, CRIZOTINIB  
XELJANZ, TOFACITINIB CITRATE  
ZMAX, AZITHROMYCIN

**PFIZER**

\* PFIZER CENTRAL RESEARCH  
DIFLUCAN, FLUCONAZOLE  
ZITHROMAX, AZITHROMYCIN

\* PFIZER CHEMICALS DIV PFIZER INC  
DIFLUCAN, FLUCONAZOLE  
ZITHROMAX, AZITHROMYCIN

\* PFIZER INC  
ADVIL ALLERGY AND CONGESTION RELIEF, CHLORPHENIRAMINE MALEATE (OTC)  
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)  
ADVIL COLD AND SINUS, IBUPROFEN (OTC)  
ADVIL CONGESTION RELIEF, IBUPROFEN (OTC)  
ADVIL LIQUI-GELS, IBUPROFEN (OTC)  
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN (OTC)  
ADVIL MULTI-SYMPTOM COLD & FLU, CHLORPHENIRAMINE MALEATE (OTC)  
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)  
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)  
ADVIL, IBUPROFEN (OTC)  
ALAVERT, LORATADINE (OTC)  
AXID AR, NIZATIDINE (OTC)  
CADUET, AMLODIPINE BESYLATE  
CALAN SR, VERAPAMIL HYDROCHLORIDE  
CARDURA XL, DOXAZOSIN MESYLATE  
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)  
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)  
CHILDREN'S ADVIL, IBUPROFEN (OTC)  
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)  
ELELYSO, TALIGLUCERASE ALFA  
GEODON, ZIPRASIDONE HYDROCHLORIDE  
GEODON, ZIPRASIDONE MESYLATE  
GLUCOTROL XL, GLIPIZIDE  
GLUCOTROL, GLIPIZIDE  
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM  
HEPARIN SODIUM, HEPARIN SODIUM  
JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)  
LIPITOR, ATORVASTATIN CALCIUM  
LORATADINE, LORATADINE (OTC)  
MERREM, MEROPENEM  
NORVASC, AMLODIPINE BESYLATE  
PEDIATRIC ADVIL, IBUPROFEN (OTC)  
PROCARDIA, NIFEDIPINE  
REVATIO, SILDENAFIL CITRATE  
SONATA, ZALEPLON  
TESSALON, BENZONATATE  
TOVIAZ, FESOTERODINE FUMARATE  
UNASYN, AMPICILLIN SODIUM  
ZITHROMAX, AZITHROMYCIN

\* PFIZER LABORATORIES DIV PFIZER INC  
CARDURA, DOXAZOSIN MESYLATE  
DIABINESE, CHLORPROPAMIDE  
FELDENE, PIROXICAM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PFIZER LABORATORIES DIV PFIZER INC  
MINIPRESS, PRAZOSIN HYDROCHLORIDE  
PFIZERPEN, PENICILLIN G POTASSIUM  
PROCARDIA XL, NIFEDIPIINE  
UNASYN, AMPICILLIN SODIUM  
VIBRAMYCIN, DOXYCYCLINE  
VIBRAMYCIN, DOXYCYCLINE CALCIUM  
VIBRAMYCIN, DOXYCYCLINE HYCLATE  
VISTARIL, HYDROXYZINE PAMOATE
- \* PFIZER PHARMACEUTICALS INC  
DILANTIN, PHENYTOIN  
ZOLOFT, SERTRALINE HYDROCHLORIDE
- \* PFIZER PHARMACEUTICALS PRODUCTION CORP LTD  
TIKOSYN, DOFETILIDE

**PFIZER CONS HLTHCARE**

- \* PFIZER CONSUMER HEALTHCARE  
ADVIL, IBUPROFEN SODIUM (OTC)

**PFIZER INC**

- \* PFIZER INC  
CAMPTOSAR, IRINOTECAN HYDROCHLORIDE  
CHANTIX, VARENICLINE TARTRATE  
ELLENCE, EPIRUBICIN HYDROCHLORIDE  
FRAGMIN, DALTEPARIN SODIUM  
IBRANCE, PALBOCICLIB  
LYRICA CR, PREGABALIN  
NICOTROL, NICOTINE  
QUILLICHEW ER, METHYLPHENIDATE HYDROCHLORIDE  
VIAGRA, SILDENAFIL CITRATE  
XELJANZ XR, TOFACITINIB CITRATE

**PFIZER IRELAND**

- \* PFIZER IRELAND PHARMACEUTICALS  
RELPAX, ELETRIPTAN HYDROBROMIDE

**PFIZER PHARMS**

- \* PFIZER PHARMACEUTICALS LTD  
ACCUPRIL, QUINAPRIL HYDROCHLORIDE  
ACCURETIC, HYDROCHLOROTHIAZIDE  
LOPID, GEMFIBROZIL  
NEURONTIN, GABAPENTIN  
NITROSTAT, NITROGLYCERIN

**PHARM ASSOC**

- \* PHARMACEUTICAL ASSOC INC  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
LORAZEPAM, LORAZEPAM  
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
- \* PHARMACEUTICAL ASSOCIATES INC  
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE  
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE  
ETHOSUXIMIDE, ETHOSUXIMIDE  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE  
HALOPERIDOL, HALOPERIDOL LACTATE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
LACTULOSE, LACTULOSE  
LEVETIRACETAM, LEVETIRACETAM  
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
MORPHINE SULFATE, MORPHINE SULFATE  
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE  
NYSTATIN, NYSTATIN  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
PREDNISOLONE, PREDNISOLONE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PHARMACEUTICAL ASSOCIATES INC  
THEOPHYLLINE, THEOPHYLLINE  
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE  
VALPROIC ACID, VALPROIC ACID

**PHARMA RES SOFTWARE**

- \* PHARMA RESEARCH SOFTWARE SOLUTION LLC  
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

**PHARMACARE**

- \* PHARMACARE LTD  
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE

**PHARMACHEMIE BV**

- \* PHARMACHEMIE BV  
CARBOPLATIN, CARBOPLATIN  
CISPLATIN, CISPLATIN  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM

**PHARMACIA AND UPJOHN**

- \* PHARMACIA AND UPJOHN  
XANAX XR, ALPRAZOLAM
- \* PHARMACIA AND UPJOHN CO  
AROMASIN, EXEMESTANE  
AZULFIDINE EN-TABS, SULFASALAZINE  
AZULFIDINE, SULFASALAZINE  
BACITRACIN, BACITRACIN  
CAVERJECT IMPULSE, ALPROSTADIL  
CAVERJECT, ALPROSTADIL  
CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE  
CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
CLEOCIN T, CLINDAMYCIN PHOSPHATE  
CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE  
CLEOCIN, CLINDAMYCIN PHOSPHATE  
CORTEF, HYDROCORTISONE  
CORVERT, IBUTILIDE FUMARATE  
CYKLOKAPRON, TRANEXAMIC ACID  
DEPO-ESTRADIOL, ESTRADIOL CYPIONATE  
DEPO-MEDROL, METHYLPREDNISOLONE ACETATE  
DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE  
DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE  
DETROL LA, TOLTERODINE TARTRATE  
DETROL, TOLTERODINE TARTRATE  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM  
ESTRING, ESTRADIOL  
GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT  
GENOTROPIN, SOMATROPIN RECOMBINANT  
GLYNASE, GLYBURIDE  
GLYSET, MIGLITOL  
HALCION, TRIAZOLAM  
HEMABATE, CARBOPROST TROMETHAMINE  
IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE  
LINCOCIN, LINCOMYCIN HYDROCHLORIDE  
MEDROL, METHYLPREDNISOLONE  
MYCOBUTIN, RIFABUTIN  
NICOTROL, NICOTINE  
OGEN 5, ESTROPIPATE  
PREPIDIL, DINOPROSTONE  
PROSTIN E2, DINOPROSTONE  
PROSTIN VR PEDIATRIC, ALPROSTADIL  
PROVERA, MEDROXYPROGESTERONE ACETATE  
R-GENE 10, ARGININE HYDROCHLORIDE  
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE  
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE  
SOMAVERT, PEGVISOMANT  
XALATAN, LATANOPROST

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PHARMACIA AND UPJOHN CO
  - XANAX, ALPRAZOLAM
  - ZINECARD, DEXRAZOXANE HYDROCHLORIDE
  - ZYVOX, LINEZOLID
- \* PHARMACIA AND UPJOHN SUB PFIZER INC
  - DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE

**PHARMACIA UPJOHN**

- \* PHARMACIA AND UPJOHN CO A SUB OF PFIZER INC
  - COLESTID, COLESTIPOL HYDROCHLORIDE
  - FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE

**PHARMACYCLICS INC**

- \* PHARMACYCLICS INC
  - IMBRUVICA, IBRUTINIB

**PHARMADAX INC**

- \* PHARMADAX INC
  - GLYBURIDE, GLYBURIDE
  - LEVETIRACETAM, LEVETIRACETAM
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE

**PHARMAFORCE**

- \* PHARMAFORCE INC
  - FLOXURIDINE, FLOXURIDINE

**PHARMA LUCENCE**

- \* PHARMA LUCENCE INC
  - AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
  - CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
  - CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
  - HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
  - TECHNETIUM TC 99M SESTAMIBI, TECHNETIUM TC-99M SESTAMIBI KIT
  - TECHNETIUM TC-99M MEBOFENIN, TECHNETIUM TC-99M MEBOFENIN KIT

**PHARMASCIENCE INC**

- \* PHARMASCIENCE INC
  - BUSULFAN, BUSULFAN
  - DECITABINE, DECITABINE
  - GANCICLOVIR SODIUM, GANCICLOVIR SODIUM

**PHARMTAK INC**

- \* PHARMTAK INC
  - LEVETIRACETAM, LEVETIRACETAM

**PHOTOCURE ASA**

- \* PHOTOCURE ASA
  - CYSVIEW KIT, HEXAMINOLEVULINATE HYDROCHLORIDE

**PIERRE FABRE**

- \* PIERRE FABRE MEDICAMENT
  - NAVELBINE, VINORELBINE TARTRATE

**PIERRE FABRE DERMA**

- \* PIERRE FABRE DERMATOLOGIE
  - HEMANGEOL, PROPRANOLOL HYDROCHLORIDE

**PIERREL**

- \* PIERREL S.P.A.
  - ORABLOC, ARTICAINE HYDROCHLORIDE

**PII**

- \* PHARMACEUTICS INTERNATIONAL INC
  - BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
  - DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE
  - HYDROCORTISONE, HYDROCORTISONE
  - PIROXICAM, PIROXICAM
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

**PIRAMAL CRITICAL**

- \* PIRAMAL CRITICAL CARE INC
  - ISOFLURANE, ISOFLURANE
  - SOJOURN, SEVOFLURANE
- \* PIRAMAL CRITICAL CARE LTD
  - GABLOFEN, BACLOFEN

**PIRAMAL ENT**

- \* PIRAMAL ENTERPRISES LTD

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PIRAMAL ENTERPRISES LTD  
ISOFLURANE, ISOFLURANE

**PIRAMAL IMAGING**

- \* PIRAMAL IMAGING SA  
NEURACEQ, FLORBETABEN F-18

**PLD ACQUISITIONS LLC**

- \* PLD ACQUISITIONS LLC  
ZOLMITRIPTAN, ZOLMITRIPTAN

**PLIVA**

- \* PLIVA INC  
AZITHROMYCIN, AZITHROMYCIN  
BENZTROPIINE MESYLATE, BENZTROPIINE MESYLATE  
CIMETIDINE, CIMETIDINE  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE  
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
METRONIDAZOLE, METRONIDAZOLE  
NAPROXEN, NAPROXEN  
THEOPHYLLINE, THEOPHYLLINE  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE  
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
WARFARIN SODIUM, WARFARIN SODIUM

**PLIVA HRVATSKA DOO**

- \* PLIVA HRVATSKA DOO  
ADAPALENE, ADAPALENE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
RISPERIDONE, RISPERIDONE

**PLIVA LACHEMA**

- \* PLIVA LACHEMA AS  
CARBOPLATIN, CARBOPLATIN  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

**PLIVA PHARM IND**

- \* PLIVA PHARMACEUTICAL INDUSTRY INC  
TORSEMIDE, TORSEMIDE

**PLX PHARMA**

- \* PLX PHARMA INC  
ASPIRIN, ASPIRIN (OTC)

**POHL BOSKAMP**

- \* POHL BOSKAMP  
NITROLINGUAL PUMPSpray, NITROGLYCERIN

**POLYGEN PHARMS**

- \* POLYGEN PHARMACEUTICALS INC  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

**POLYMEDICA**

- \* POLYMEDICA INDUSTRIES INC  
NEOPAP, ACETAMINOPHEN (OTC)

**PORTOLA PHARMS INC**

- \* PORTOLA PHARMACEUTICALS INC  
BEVYXXA, BETRIXABAN

**POWDER PHARMS**

- \* POWDER PHARMACEUTICALS INC  
ZINGO, LIDOCAINE HYDROCHLORIDE

**PRAGMA PHARMS LLC**

- \* PRAGMA PHARMACEUTICALS LLC  
KEFLEX, CEPHALEXIN

**PRECISION DERMAT**

- \* PRECISION DERMATOLOGY INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

- \* PRECISION DERMATOLOGY INC  
CLINDAGEL, CLINDAMYCIN PHOSPHATE  
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE  
LOCOID, HYDROCORTISONE BUTYRATE  
MINOCIN, MINOCYCLINE HYDROCHLORIDE

**PRECISION DOSE**

- \* PRECISION DOSE INC  
RISPERIDONE, RISPERIDONE

**PRECISION DOSE INC**

- \* PRECISION DOSE INC  
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE

**PRECISION NUCLEAR**

- \* PRECISION NUCLEAR LLC  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**PRINSTON INC**

- \* PRINSTON PHARMACEUTICAL INC  
ARIPIPRAZOLE, ARIPIPRAZOLE  
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
CAPTOPRIL, CAPTOPRIL  
CLONAZEPAM, CLONAZEPAM  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE  
DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE  
ENTECAVIR, ENTECAVIR  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM  
FUROSEMIDE, FUROSEMIDE  
GLIMEPIRIDE, GLIMEPIRIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE  
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
IRBESARTAN, IRBESARTAN  
LEVETIRACETAM, LEVETIRACETAM  
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LISINOPRIL, LISINOPRIL  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
METHOCARBAMOL, METHOCARBAMOL  
NEVIRAPINE, NEVIRAPINE  
OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
PAROXETINE MESYLATE, PAROXETINE MESYLATE  
PAROXETINE, PAROXETINE HYDROCHLORIDE  
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE  
RISPERIDONE, RISPERIDONE  
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE  
TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
TELMISARTAN, TELMISARTAN  
TEMAZEPAM, TEMAZEPAM  
VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
VALSARTAN, VALSARTAN  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE  
VORICONAZOLE, VORICONAZOLE

**PROF DSPLS**

- \* PROFESSIONAL DISPOSABLES INC  
PREVANTICS MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)  
PREVANTICS SWAB, CHLORHEXIDINE GLUCONATE (OTC)  
PREVANTICS SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* P \*\*

**PROMIUS PHARMA**

- \* PROMIUS PHARMA LLC  
SECTRAL, ACEBUTOLOL HYDROCHLORIDE  
TENEX, GUANFACINE HYDROCHLORIDE

**PROMIUS PHARMA LLC**

- \* PROMIUS PHARMA LLC  
CLODERM, CLOCORTOLONE PIVALATE  
IMPOYZ, CLOBETASOL PROPIONATE  
SERNIVO, BETAMETHASONE DIPROPIONATE

**PROVENESIS**

- \* PROVENESIS LTD  
VARITHENA, POLIDOCANOL

**PROVEPHARM SAS**

- \* PROVEPHARM SAS  
PROVAYBLUE, METHYLENE BLUE

**PROVIDENT PHARM**

- \* PROVIDENT PHARMACEUTICAL INC  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**PTC THERAP**

- \* PTC THERAPEUTICS INC  
EMFLAZA, DEFLAZACORT

**PULMOFLOW INC**

- \* PULMOFLOW INC  
KITABIS PAK, TOBRAMYCIN

**PUMA BIOTECH**

- \* PUMA BIOTECHNOLOGY INC  
NERLYNX, NERATINIB MALEATE

**PURACAP PHARM**

- \* PURACAP PHARMACEUTICAL LLC  
MELOXICAM, MELOXICAM

**PURACAP PHARM LLC**

- \* PURACAP PHARMACEUTICAL LLC  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE

**PURDUE GMP**

- \* PURDUE GMP CENTER LLC DBA THE CHAO CENTER INDUSTRIAL PHARMACY  
SEROMYCIN, CYCLOSERINE

**PURDUE PHARMA**

- \* PURDUE PHARMA PRODUCTS LP  
INTERMEZZO, ZOLPIDEM TARTRATE

**PURDUE PHARMA LP**

- \* PURDUE PHARMA LP  
BUTRANS, BUPRENORPHINE  
HYSINGLA, HYDROCODONE BITARTRATE  
MS CONTIN, MORPHINE SULFATE  
OXYCONTIN, OXYCODONE HYDROCHLORIDE

\*\* Q \*\*

**QILU PHARM CO LTD**

- \* QILU PHARMACEUTICAL CO LTD  
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM  
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE  
CEFTRIAXONE, CEFTRIAXONE SODIUM  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
OLANZAPINE, OLANZAPINE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
OXALIPLATIN, OXALIPLATIN

**QOL MEDCL**

- \* QOL MEDICAL LLC  
ETHAMOLIN, ETHANOLAMINE OLEATE  
SUCRAID, SACROSIDASE

**QUEEN HAMAMATSU PET**

- \* QUEEN HAMAMATSU PET IMAGING CENTER  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* R \*\*

**R-PHARM US LLC**

- \* R-PHARM US LLC  
IXEMPRA KIT, IXABEPILONE

**R2 PHARMA LLC**

- \* R2 PHARMA LLC  
ESOMEPRAZOLE STRONTIUM, ESOMEPRAZOLE STRONTIUM

**RADIUS HEALTH INC**

- \* RADIUS HEALTH INC  
TYMLOS, ABALOPARATIDE

**RANBAXY**

- \* RANBAXY LABORATORIES INC  
EURAX, CROTAMITON  
HALOG, HALCINONIDE  
ULTRAVATE, HALOBETASOL PROPIONATE

**RANBAXY LABS LTD**

- \* RANBAXY LABORATORIES LTD  
KENALOG, TRIAMCINOLONE ACETONIDE

**RECIP**

- \* RECIP AB  
THYROSafe, POTASSIUM IODIDE (OTC)

**RECKITT BENCKISER**

- \* RECKITT BENCKISER LLC  
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)  
LEVONORGESTREL, LEVONORGESTREL (OTC)  
MUCINEX D, GUAIFENESIN (OTC)  
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)  
MUCINEX, GUAIFENESIN (OTC)

**RECORDATI RARE**

- \* RECORDATI RARE DISEASES INC  
CHEMET, SUCCIMER  
COSMEGEN, DACTINOMYCIN  
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE  
INDOCIN, INDOMETHACIN SODIUM  
MUSTARGEN, MECHLORETHAMINE HYDROCHLORIDE  
NEOPROFEN, IBUPROFEN LYSINE  
PEGANONE, ETHOTOIN  
TRANXENE, CLORAZEPATE DIPOTASSIUM

**RECRO GAINESVILLE**

- \* RECRO GAINESVILLE LLC  
VERELAN PM, VERAPAMIL HYDROCHLORIDE  
VERELAN, VERAPAMIL HYDROCHLORIDE

**RELYPSA INC**

- \* RELYPSA INC  
VELTASSA, PATIROMER SORBITEX CALCIUM

**REMPEX PHARMS INC**

- \* REMPEX PHARMACEUTICALS INC  
MINOCIN, MINOCYCLINE HYDROCHLORIDE

**REMPEX PHARMS MEDCNS**

- \* REMPEX PHARMACEUTICALS A WHOLLY OWNED SUB OF THE MEDICINES CO  
VABOMERE, MEROPENEM

**RENAISSANCE SSA LLC**

- \* RENAISSANCE SSA LLC  
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE  
METHOCARBAMOL, METHOCARBAMOL  
MILRINONE LACTATE IN DEXTROSE 5%, MILRINONE LACTATE  
OXACILLIN SODIUM, OXACILLIN SODIUM  
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM

**RHODES PHARMS**

- \* RHODES PHARMACEUTICALS LP  
APTENSIO XR, METHYLPHENIDATE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE  
DILAUDID, HYDROMORPHONE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* R \*\*

- \* RHODES PHARMACEUTICALS LP
  - FENOFIBRATE (MICRONIZED), FENOFIBRATE
  - FENOFIBRATE, FENOFIBRATE
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - MORPHINE SULFATE, MORPHINE SULFATE
  - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - THEOPHYLLINE, THEOPHYLLINE

**RICONPHARMA LLC**

- \* RICONPHARMA LLC
  - BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
  - CHLORTHALIDONE, CHLORTHALIDONE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
  - LIDOCAINE, LIDOCAINE
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - QUININE SULFATE, QUININE SULFATE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**RISING PHARMS INC**

- \* RISING PHARMACEUTICALS INC
  - CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - CROMOLYN SODIUM, CROMOLYN SODIUM
  - DESOXIMETASONE, DESOXIMETASONE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DOXERCALCIFEROL, DOXERCALCIFEROL
  - DUTASTERIDE, DUTASTERIDE
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - HYDROCORTISONE, HYDROCORTISONE
  - LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN
  - LEVOCARNITINE, LEVOCARNITINE
  - LEVOFLOXACIN, LEVOFLOXACIN
  - METHIMAZOLE, METHIMAZOLE
  - PARICALCITOL, PARICALCITOL
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - ZILEUTON, ZILEUTON

**ROCHE PALO**

- \* ROCHE PALO ALTO LLC
  - CELLCEPT, MYCOPHENOLATE MOFETIL
  - CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
  - CYTOVENE, GANCICLOVIR SODIUM

**ROCKWELL MEDICAL INC**

- \* ROCKWELL MEDICAL INC
  - TRIFERIC, FERRIC PYROPHOSPHATE CITRATE
  - TRIFERIC, FERRIC PYROPHOSPHATE CITRATE

**ROMARK**

- \* ROMARK LABORATORIES
  - ALINIA, NITAZOXANIDE

**ROUSES POINT PHARMS**

- \* ROUSES POINT PHARMACEUTICALS LLC
  - LEVETIRACETAM, LEVETIRACETAM

**RP SCHERER**

- \* RP SCHERER TECHNOLOGIES LLC
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

**RTRX**

- \* RETROPHIN INC
  - CHOLBAM, CHOLIC ACID

**RUBICON RES PVT LTD**

- \* RUBICON RESEARCH PVT LTD
  - BACLOFEN, BACLOFEN
  - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE

\*\* S \*\*

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

**SAGE PRODS**

- \* SAGE PRODUCTS INC  
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

**SAGENT AGILA**

- \* SAGENT AGILA LLC  
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM  
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE  
BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE  
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

**SAGENT AGILA LLC**

- \* SAGENT AGILA LLC  
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
LEVOFLOXACIN, LEVOFLOXACIN  
MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE, MIDAZOLAM HYDROCHLORIDE  
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE

**SAGENT PHARMS**

- \* SAGENT PHARMACEUTICALS INC  
ACETYLCYSTEINE, ACETYLCYSTEINE  
AMIKACIN SULFATE, AMIKACIN SULFATE  
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM  
AMPICILLIN SODIUM, AMPICILLIN SODIUM  
CAFFEINE CITRATE, CAFFEINE CITRATE  
CEFEPIME HYDROCHLORIDE, CEFEPIME HYDROCHLORIDE  
CEFTRIAZONE SODIUM, CEFTRIAZONE SODIUM  
CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM  
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE  
COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM  
DAPTOMYCIN, DAPTOMYCIN  
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE  
FLUMAZENIL, FLUMAZENIL  
FLUOROURACIL, FLUOROURACIL  
GLYDO, LIDOCAINE HYDROCHLORIDE  
HALOPERIDOL, HALOPERIDOL LACTATE  
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM  
HEPARIN SODIUM, HEPARIN SODIUM  
IBANDRONATE SODIUM, IBANDRONATE SODIUM  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM  
LEVETIRACETAM, LEVETIRACETAM  
LINEZOLID, LINEZOLID  
MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE  
MESNA, MESNA  
METHOCARBAMOL, METHOCARBAMOL  
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE  
NAFCILLIN SODIUM, NAFCILLIN SODIUM  
OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE  
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE  
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE  
OXACILLIN SODIUM, OXACILLIN SODIUM  
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM  
PENTOBARBITAL SODIUM, PENTOBARBITAL SODIUM  
PROPOFOL, PROPOFOL  
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE  
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE  
TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE  
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE  
VECURONIUM BROMIDE, VECURONIUM BROMIDE  
ZOLEDRONIC ACID, ZOLEDRONIC ACID

**SAGENT STRIDES**

- \* SAGENT STRIDES LLC  
ADENOSINE, ADENOSINE  
AZITHROMYCIN, AZITHROMYCIN  
BACITRACIN, BACITRACIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SAGENT STRIDES LLC
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MIDOZALAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
  - POLYMYCIN B SULFATE, POLYMYXIN B SULFATE
  - ROPIVACAINE HYDROCHLORIDE, ROPIVACAINE HYDROCHLORIDE

**SALIX PHARMS**

- \* SALIX PHARMACEUTICALS INC
  - ANUSOL HC, HYDROCORTISONE
  - DIURIL, CHLOROTHIAZIDE
  - METOZOLV ODT, METOCLOPRAMIDE HYDROCHLORIDE
  - MOVIPREP, ASCORBIC ACID
  - OSMOPREP, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
  - PEPCID, FAMOTIDINE
  - RELISTOR, METHYLNALTREXONE BROMIDE
  - XIFAXAN, RIFAXIMIN

**SALIX PHARMS INC**

- \* SALIX PHARMACEUTICALS INC
  - RELISTOR, METHYLNALTREXONE BROMIDE

**SAMSON MEDCL**

- \* SAMSON MEDICAL TECHNOLOGIES LLC
  - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
  - CEFOXITIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
  - CEFTRIAXONE, CEFTRIAXONE SODIUM
  - VANCOMYCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE

**SANDOZ**

- \* SANDOZ
  - DOCETAXEL, DOCETAXEL
- \* SANDOZ INC
  - ALPRAZOLAM, ALPRAZOLAM
  - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
  - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
  - AMOXICILLIN, AMOXICILLIN
  - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
  - AMPICILLIN SODIUM, AMPICILLIN SODIUM
  - AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
  - APREPITANT, APREPITANT
  - ARGATROBAN IN SODIUM CHLORIDE, ARGATROBAN
  - ATENOLOL, ATENOLOL
  - AZITHROMYCIN, AZITHROMYCIN
  - BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
  - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
  - BICALUTAMIDE, BICALUTAMIDE
  - BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
  - BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
  - BUMETANIDE, BUMETANIDE
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CAFERGOT, CAFFEINE
  - CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL
  - CARISOPRODOL AND ASPIRIN, ASPIRIN
  - CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
  - CARVEDILOL, CARVEDILOL
  - CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
  - CEFDINIR, CEFDINIR
  - CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
  - CEFPROZIL, CEFPROZIL
  - CEFTRIAXONE, CEFTRIAXONE SODIUM
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
  - CHOLESTYRAMINE, CHOLESTYRAMINE
  - CILOSTAZOL, CILOSTAZOL

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SANDOZ INC
  - CLARITHROMYCIN, CLARITHROMYCIN
  - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
  - CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
  - CLONAZEPAM, CLONAZEPAM
  - COSYNTROPIN, COSYNTROPIN
  - CYCLOSPORINE, CYCLOSPORINE
  - DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
  - DESLORATADINE, DESLORATADINE
  - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
  - DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
  - DICLOFENAC SODIUM AND MISOPROSTOL, DICLOFENAC SODIUM
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
  - ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM
  - EPLERENONE, EPLERENONE
  - ETODOLAC, ETODOLAC
  - FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
  - FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
  - FUROSEMIDE, FUROSEMIDE
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - GLIPIZIDE, GLIPIZIDE
  - HALOPERIDOL, HALOPERIDOL
  - HEPARIN SODIUM, HEPARIN SODIUM
  - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
  - HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
  - IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
  - INDOMETHACIN, INDOMETHACIN
  - IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - IRBESARTAN, IRBESARTAN
  - ISONIAZID, ISONIAZID
  - ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
  - ITRACONAZOLE, ITRACONAZOLE
  - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LISINOPRIL, LISINOPRIL
  - LORATADINE, LORATADINE (OTC)
  - LORAZEPAM, LORAZEPAM
  - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - LOVASTATIN, LOVASTATIN
  - MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
  - METAXALONE, METAXALONE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
  - METHAZOLAMIDE, METHAZOLAMIDE
  - METHIMAZOLE, METHIMAZOLE
  - METHYLSPREDNISOLONE, Methylprednisolone
  - METOLAZONE, METOLAZONE
  - MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
  - NABUMETONE, NABUMETONE
  - NADOLOL, NADOLOL
  - NAFCILLIN SODIUM, NAFCILLIN SODIUM
  - NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SANDOZ INC
  - NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
  - NIZATIDINE, NIZATIDINE
  - OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - OMEPRAZOLE, OMEPRAZOLE
  - OMNITROPE, SOMATROPIN RECOMBINANT
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON, ONDANSETRON
  - ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
  - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
  - OXACILLIN SODIUM, OXACILLIN SODIUM
  - OXALIPLATIN, OXALIPLATIN
  - OXaprozin, OXaprozin
  - Oxazepam, Oxazepam
  - PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
  - PENICILLIN G SODIUM, PENICILLIN G SODIUM
  - PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
  - PERPHENAZINE, PERPHENAZINE
  - PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PIOGLITAZONE HYDROCHLORIDE AND GLIMEPIRIDE, GLIMEPIRIDE
  - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - QUINIDINE SULFATE, QUINIDINE SULFATE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RIBAVIRIN, RIBAVIRIN
  - RIFAMPIN, RIFAMPIN
  - RISPERIDONE, RISPERIDONE
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - SPIRONOLACTONE, SPIRONOLACTONE
  - SULFADIAZINE, SULFADIAZINE
  - TACROLIMUS, TACROLIMUS
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TEMAZEPAM, TEMAZEPAM
  - TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
  - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**SANDOZ INC**

- \* SANDOZ INC
  - ACETAMINOPHEN, ACETAMINOPHEN
  - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
  - ANECTINE, SUCCINYLCHOLINE CHLORIDE
  - ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
  - ASPIRIN AND DIPYRIDAMOLE, ASPIRIN
  - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
  - BIMATOPROST, BIMATOPROST
  - BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
  - BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
  - BUDESONIDE, BUDESONIDE
  - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - BUSULFAN, BUSULFAN
  - CARBOPLATIN, CARBOPLATIN
  - CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
  - CEFIXIME, CEFIXIME
  - CEFTRIAXONE, CEFTRIAXONE SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SANDOZ INC  
CISATRACURIUM BESYLATE PRESERVATIVE FREE, CISATRACURIUM BESYLATE  
CISATRACURIUM BESYLATE, CISATRACURIUM BESYLATE  
CLINDAMYCIN PHOSPHATE IN 5% DEXTROSE IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE  
DECITABINE, DECITABINE  
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE  
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE  
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DIGOXIN, DIGOXIN  
DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE  
DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE  
DOXERCALCIFEROL, DOXERCALCIFEROL  
ENALAPRIL MALEATE, ENALAPRIL MALEATE  
ENOXAPARIN SODIUM, ENOXAPARIN SODIUM  
ENTECAVIR, ENTECAVIR  
EPHEDRINE SULFATE, EPHEDRINE SULFATE  
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE  
EZETIMIBE, EZETIMIBE  
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE  
FLUMAZENIL, FLUMAZENIL  
GATIFLOXACIN, GATIFLOXACIN  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
GLATOPA, GLATIRAMER ACETATE  
GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE  
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE  
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE  
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE  
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID  
INFUVITE PEDIATRIC, ASCORBIC ACID  
ISONIAZID, ISONIAZID  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYCIN, AMOXICILLIN  
LATANOPROST, LATANOPROST  
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE  
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM  
LINEZOLID, LINEZOLID  
MAXITROL, DEXAMETHASONE  
METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM  
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE  
METIPRANOL, METIPRANOL HYDROCHLORIDE  
METOPROLOL TARTRATE, METOPROLOL TARTRATE  
MONTELUKAST SODIUM, MONTELUKAST SODIUM  
MYDRIACYL, TROPICAMIDE  
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE  
NEVIRAPINE, NEVIRAPINE  
OFLOXACIN, OFLOXACIN  
OLANZAPINE, OLANZAPINE  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
PACLITAXEL, PACLITAXEL  
PALONOSETRON HYDROCHLORIDE, PALONOSETRON HYDROCHLORIDE  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM  
PARICALCITOL, PARICALCITOL  
PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM  
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE  
PROGESTERONE, PROGESTERONE  
QOLIANA, BRIMONIDINE TARTRATE  
REGONOL, PYRIDOSTIGMINE BROMIDE  
REPAGLINIDE, REPAGLINIDE  
RIBAVIRIN, RIBAVIRIN  
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE  
ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE  
ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM  
SILODOSIN, SILODOSIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SANDOZ INC
  - SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
  - SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
  - TELMISARTAN, TELMISARTAN
  - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
  - TIGECYCLINE, TIGECYCLINE
  - TIMOLOL MALEATE, TIMOLOL MALEATE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TOBREX, TOBRAMYCIN
  - TREPROSTINIL, TREPROSTINIL
  - TRIFLURIDINE, TRIFLURIDINE
  - TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VORICONAZOLE, VORICONAZOLE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**SANJA PHARMS CO**

- \* SANJA PHARMACEUTICALS CO
  - CARBOPLATIN, CARBOPLATIN
  - OXALIPLATIN, OXALIPLATIN

**SANOCHEMIA CORP USA**

- \* SANOCHEMIA CORP USA
  - SCANLUX-300, IOPAMIDOL
  - SCANLUX-370, IOPAMIDOL

**SANOFI AVENTIS US**

- \* SANOFI AVENTIS US INC
  - JEVTANA KIT, CABAZITAXEL
- \* SANOFI AVENTIS US LLC
  - ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - ALLEGRA, FEXOFENADINE HYDROCHLORIDE
  - ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
  - ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION, FEXOFENADINE HYDROCHLORIDE (OTC)
  - AMARYL, GLIMEPIRIDE
  - AMBIEN CR, ZOLPIDEM TARTRATE
  - AMBIEN, ZOLPIDEM TARTRATE
  - APIDRA SOLOSTAR, INSULIN GLULISINE RECOMBINANT
  - APIDRA, INSULIN GLULISINE RECOMBINANT
  - ARAVA, LEFLUNOMIDE
  - AUBAGIO, TERIFLUNOMIDE
  - avalide, HYDROCHLOROTHIAZIDE
  - AVapro, IRBESARTAN
  - CHILDREN'S ALLEGRA ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHILDREN'S ALLEGRA HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CLomid, CLOMIPHENE CITRATE
  - DIABETA, GLYBURIDE
  - ELOXATIN, OXALIPLATIN
  - FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
  - GAVISCON, ALUMINUM HYDROXIDE (OTC)
  - LANTUS SOLOSTAR, INSULIN GLARGINE RECOMBINANT
  - LANTUS, INSULIN GLARGINE RECOMBINANT
  - LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
  - LOVENOX, ENOXAPARIN SODIUM
  - MULTAQ, DRONEDARONE HYDROCHLORIDE
  - NASACORT ALLERGY 24 HOUR, TRIAMCINOLONE ACETONIDE (OTC)
  - NICODERM CQ, NICOTINE (OTC)
  - PLAVIX, CLOPIDOGREL BISULFATE
  - PRIFTIN, RIFAPENTINE
  - PRIMAQUINE, PRIMAQUINE PHOSPHATE
  - RIFADIN, RIFAMPIN
  - RIFAMATE, ISONIAZID
  - RIFATER, ISONIAZID
  - TAXOTERE, DOCETAXEL

**SANOFI US**

- \* SANOFI US

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SANOFI US
  - ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
  - ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)

**SANOFI US SERVICES**

- \* SANOFI US SERVICES INC
  - TOUJEOL SOLOSTAR, INSULIN GLARGINE RECOMBINANT

**SANOFI-AVENTIS US**

- \* SANOFI-AVENTIS US LLC
  - ADLYXIN, LIXISENATIDE
  - ADMELOG SOLOSTAR, INSULIN LISPRO
  - ADMELOG, INSULIN LISPRO
  - SOLIQUA 100/33, INSULIN GLARGINE
  - XYZAL ALLERGY 24HR, LEVOCETIRIZINE DIHYDROCHLORIDE (OTC)
  - XYZAL, LEVOCETIRIZINE DIHYDROCHLORIDE

**SANTARUS INC**

- \* SANTARUS INC
  - FENOGLIDE, FENOFIBRATE
  - GLUMETZA, METFORMIN HYDROCHLORIDE
  - ZEGERID, OMEPRAZOLE

**SANTOS BIOTECH**

- \* SANTOS BIOTECH INDUSTRIES INC
  - ANASTROZOLE, ANASTROZOLE
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - BICALUTAMIDE, BICALUTAMIDE
  - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
  - GLYCOPYRROLATE, GLYCOPYRROLATE

**SAOL THERAPS RES LTD**

- \* SAOL THERAPEUTICS RESEARCH LTD
  - LIORESAL, BACLOFEN

**SAPTALIS PHARMS**

- \* SAPTALIS PHARMACEUTICALS LLC
  - LORAZEPAM, LORAZEPAM

**SAREPTA THERAPS INC**

- \* SAREPTA THERAPEUTICS INC
  - EXONDYS 51, ETEPLIRSEN

**SAVIOR LIFETEC CORP**

- \* SAVIOR LIFETEC CORP
  - MEROPENEM, MEROPENEM

**SAWAI USA**

- \* SAWAI USA INC
  - PITAVASTATIN CALCIUM, PITAVASTATIN CALCIUM

**SB PHARMCO**

- \* SB PHARMCO PUERTO RICO INC
  - AVANDIA, ROSIGLITAZONE MALEATE

**SCHERING**

- \* SCHERING CORP
  - INTEGRILIN, EPTIFIBATIDE
  - NOXAFL, POSACONAZOLE
  - REBETOL, RIBAVIRIN

**SCHERING PLOUGH**

- \* SCHERING PLOUGH HEALTHCARE PRODUCTS INC
  - AFRINOL, PSEUDOEPHEDRINE SULFATE (OTC)

**SCIECURE PHARMA INC**

- \* SCIECURE PHARMA INC
  - BUDESONIDE, BUDESONIDE

**SCIEGEN PHARMS INC**

- \* SCIEGEN PHARMACEUTICALS INC
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CARISOPRODOL, CARISOPRODOL
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SCIEGEN PHARMACEUTICALS INC
  - FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - GABAPENTIN, GABAPENTIN
  - IRBESARTAN, IRBESARTAN
  - LAMOTRIGINE, LAMOTRIGINE
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - METAXALONE, METAXALONE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - OMEPRAZOLE AND SODIUM BICARBONATE, OMEPRAZOLE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE

**SCIOS LLC**

- \* SCIOS LLC
  - NATRECOR, NESIRITIDE RECOMBINANT

**SEBELA IRELAND LTD**

- \* SEBELA IRELAND LTD
  - BRISDELLE, PAROXETINE MESYLATE
  - IMURAN, AZATHIOPRINE
  - LOTRONEX, ALOSETRON HYDROCHLORIDE
  - MICORT-HC, HYDROCORTISONE ACETATE
  - MOTOFEN, ATROPINE SULFATE
  - NAFTIN, NAFTIFINE HYDROCHLORIDE
  - ONMEL, ITRACONAZOLE
  - PEXEVA, PAROXETINE MESYLATE
  - PRAMOSONE, HYDROCORTISONE ACETATE
  - RIDAURA, AURANOFIN

**SECAN PHARMS**

- \* SECAN PHARMACEUTICALS INC
  - LEVETIRACETAM, LEVETIRACETAM

**SENTYNL THERAPS INC**

- \* SENTYNL THERAPEUTICS INC
  - ABSTRAL, FENTANYL CITRATE
  - LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE

**SEPTODONT**

- \* SEPTODONT INC
  - BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
  - OCTOCAINE, EPINEPHRINE

**SEPTODONT HOLDING**

- \* SEPTODONT HOLDING SAS
  - ORaverse, PHENTOLAMINE MESYLATE

**SEPTODONT INC**

- \* SEPTODONT INC
  - ISOCAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
  - LIDOCAINE, LIDOCAINE
  - PRILOCAINE HYDROCHLORIDE AND EPINEPHRINE BITARTRATE, EPINEPHRINE BITARTRATE
  - PRILOCAINE HYDROCHLORIDE, PRILOCAINE HYDROCHLORIDE

**SERB SA**

- \* SERB SA
  - CYANOKIT, HYDROXOCOBALAMIN

**SETON PHARM**

- \* SETON PHARMACEUTICAL LLC
  - PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE

**SETON PHARMS**

- \* SETON PHARMACEUTICALS LLC
  - PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

**SHANGHAI HENGRI**

- \* SHANGHAI HENGRI PHARMACEUTICAL CO LTD
  - SEVOFLURANE, SEVOFLURANE

**SHENZHEN TECHDOW**

- \* SHENZHEN TECHDOW PHARMACEUTICAL CO LTD
  - HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
  - HEPARIN SODIUM, HEPARIN SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

**SHERTECH LABS LLC**

- \* SHERTECH LABORATORIES LLC  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**SHILPA MEDICARE**

- \* SHILPA MEDICARE LTD  
AZACITIDINE, AZACITIDINE

**SHILPA MEDICARE LTD**

- \* SHILPA MEDICARE LTD  
CAPECITABINE, CAPECITABINE

**SHIONOGI INC**

- \* SHIONOGI INC  
PONSTEL, MEFENAMIC ACID  
SYMPROIC, NALDEMEDINE TOSYLATE  
ULESFIA, BENZYL ALCOHOL

**SHIRE**

- \* SHIRE DEVELOPMENT INC  
ADDERALL XR 10, AMPHETAMINE ASPARTATE  
ADDERALL XR 15, AMPHETAMINE ASPARTATE  
ADDERALL XR 20, AMPHETAMINE ASPARTATE  
ADDERALL XR 25, AMPHETAMINE ASPARTATE  
ADDERALL XR 30, AMPHETAMINE ASPARTATE  
ADDERALL XR 5, AMPHETAMINE ASPARTATE  
CARBATROL, CARBAMAZEPINE  
INTUNIV, GUANFACINE HYDROCHLORIDE  
LIALDA, MESALAMINE  
PENTASA, MESALAMINE

**SHIRE DEV LLC**

- \* SHIRE DEVELOPMENT LLC  
FOSRENOL, LANTHANUM CARBONATE  
MYDAYIS, AMPHETAMINE ASPARTATE  
VYVANSE, LISDEXAMFETAMINE Dimesylate  
XIIDRA, LIFITEGRAST

**SHIRE DEVELOPMENT**

- \* SHIRE DEVELOPMENT INC  
VYVANSE, LISDEXAMFETAMINE Dimesylate

**SHIRE HUMAN GENETIC**

- \* SHIRE HUMAN GENETIC THERAPIES INC  
VPRIV, VELAGLUCERASE ALFA

**SHIRE LLC**

- \* SHIRE DEVELOPMENT LLC  
AGRYLIN, ANAGRELIDE HYDROCHLORIDE  
FOSRENOL, LANTHANUM CARBONATE

**SHIRE ORPHAN THERAP**

- \* SHIRE ORPHAN THERAPIES INC  
FIRAZYR, ICATIBANT ACETATE

**SIDMAK LABS INDIA**

- \* SIDMAK LABORATORIES INDIA PVT LTD  
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

**SIGMA TAU**

- \* SIGMA TAU PHARMACEUTICALS INC  
ADAGEN, PEGADEMASE BOVINE

**SIGMAPHARM LABS LLC**

- \* SIGMAPHARM LABORATORIES LLC  
ACITRETIN, ACITRETIN  
ADEFOVIR DIPIVOXIL, ADEFUVIR DIPIVOXIL  
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE  
DISULFIRAM, DISULFIRAM  
ERGOCALCIFEROL, ERGOCALCIFEROL  
FLUCYTOSINE, FLUCYTOSINE  
GRISEOFULVIN, GRISEOFULVIN, MICROSIZE  
GRISEOFULVIN, ULTRAMICROSIZE, GRISEOFULVIN, ULTRAMICROSIZE  
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SIGMAPHARM LABORATORIES LLC
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
  - PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
  - SODIUM PHENYLBUTYRATE, SODIUM PHENYLBUTYRATE

**SILARX**

- \* SILARX PHARMACEUTICALS INC
  - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - DOXEPEPIN HYDROCHLORIDE, DOXEPEPIN HYDROCHLORIDE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - LEVETIRACETAM, LEVETIRACETAM
  - LORATADINE, LORATADINE (OTC)
  - METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - THEOPHYLLINE, THEOPHYLLINE

**SILARX PHARMS INC**

- \* SILARX PHARMACEUTICALS INC
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - LAMIVUDINE, LAMIVUDINE
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - LOPINAVIR AND RITONAVIR, LOPINAVIR
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE

**SILVERGATE PHARMS**

- \* SILVERGATE PHARMACEUTICALS INC
  - EPANED KIT, ENALAPRIL MALEATE
  - EPANED, ENALAPRIL MALEATE
  - QBRELIS, LISINOPRIL
  - XATMEP, METHOTREXATE SODIUM

**SINOOTHERAPEUTICS INC**

- \* SINOOTHERAPEUTICS INC
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE

**SKINMEDICA**

- \* SKINMEDICA INC
  - VANIQA, EFLORNITHINE HYDROCHLORIDE

**SKYEPHARMA AG**

- \* SKYEPHARMA AG
  - TRIGLIDE, FENOGLIBRATE

**SMITHKLINE BEECHAM**

- \* SMITHKLINE BEECHAM
  - LOVAZA, OMEGA-3-ACID ETHYL ESTERS
- \* SMITHKLINE BEECHAM (CORK) LTD IRELAND
  - COREG CR, CARVEDILOL PHOSPHATE
  - COREG, CARVEDILOL

**SOAPCO**

- \* SOAPCO INC
  - BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)

**SOFGEN PHARMS**

- \* SOFGEN PHARMACEUTICALS
  - NIMODIPINE, NIMODIPINE
- \* SOFGEN PHARMACEUTICALS LLC
  - IBUPROFEN, IBUPROFEN (OTC)
  - PROGESTERONE, PROGESTERONE

**SOFIE**

- \* SOFIE CO DBA SOFIE (FKA ZEVACOR PHARMA INC)
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**SOMERSET**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SOMERSET PHARMACEUTICALS INC  
ELDEPRYL, SELEGILINE HYDROCHLORIDE  
EMSAM, SELEGILINE

**SOMERSET THERAPS LLC**

- \* SOMERSET THERAPEUTICS LLC  
CYANOCOBALAMIN, CYANOCOBALAMIN  
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE  
GLYCOPYRROLATE, GLYCOPYRROLATE  
METHOCARBAMOL, METHOCARBAMOL  
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE  
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE  
TOBRAMYCIN, TOBRAMYCIN

**SPEAR PHARMS**

- \* SPEAR PHARMACEUTICALS INC  
FLUOROURACIL, FLUOROURACIL

**SPECGX LLC**

- \* SPECGX LLC  
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN  
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE  
ANEXSIA 5/325, ACETAMINOPHEN  
ANEXSIA 7.5/325, ACETAMINOPHEN  
BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE  
BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE  
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE  
EXALGO, HYDROMORPHONE HYDROCHLORIDE  
FENTANYL CITRATE, FENTANYL CITRATE  
FENTANYL-100, FENTANYL  
FENTANYL-12, FENTANYL  
FENTANYL-25, FENTANYL  
FENTANYL-50, FENTANYL  
FENTANYL-75, FENTANYL  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE  
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE  
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE  
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE  
METHADOSE, METHADONE HYDROCHLORIDE  
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE  
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE  
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
MORPHINE SULFATE, MORPHINE SULFATE  
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE  
OXYCET, ACETAMINOPHEN  
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
OXYMORPHONE HYDROCHLORIDE, OXMORPHONE HYDROCHLORIDE  
PAMELOR, NORTRIPTYLINE HYDROCHLORIDE  
RESTORIL, TEMAZEPAM  
ROXICODONE, OXYCODONE HYDROCHLORIDE  
TOFRANIL, IMIPRAMINE HYDROCHLORIDE  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

**SPECTRA MDCL DEVICES**

- \* SPECTRA MEDICAL DEVICES INC  
SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE

**SPECTRON MRC LLC**

- \* SPECTRON MRC LLC  
AMMONIA N 13, AMMONIA N-13  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**SPECTRUM PHARMS**

- \* SPECTRUM PHARMACEUTICALS INC  
BELEODAQ, BELINOSTAT

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SPECTRUM PHARMACEUTICALS INC  
EVOMELA, MELPHALAN HYDROCHLORIDE  
FUSILEV, LEVOLEUCOVORIN CALCIUM

**SPROUT PHARMS**

- \* SPROUT PHARMACEUTICALS INC  
ADDYI, FLIBANSERIN

**ST RENATUS**

- \* ST RENATUS LLC  
KOVANAZE, OXYMETAZOLINE HYDROCHLORIDE

**STAND HOMEOPATH**

- \* STANDARD HOMEOPATHIC CO  
IVY BLOCK, BENTOQUATAM (OTC)

**STASON PHARMS**

- \* STASON PHARMACEUTICALS INC  
PURINETHOL, MERCAPTOPURINE

**STI PHARMA LLC**

- \* STI PHARMA LLC  
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE  
DEXAMETHASONE, DEXAMETHASONE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE  
SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE  
TRIACIN-C, CODEINE PHOSPHATE

**STIEFEL**

- \* STIEFEL LABORATORIES INC  
DUAC, BENZOYL PEROXIDE

**STIEFEL LABS INC**

- \* STIEFEL LABORATORIES INC  
SORIATANE, ACITRETIN

**STRIDES PHARMA**

- \* STRIDES PHARMA GLOBAL PTE LTD  
ABACAVIR SULFATE, ABACAVIR SULFATE  
ACARBOSE, ACARBOSE  
ACETAZOLAMIDE, ACETAZOLAMIDE  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
BENZONATATE, BENZONATATE  
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE  
CALCITRIOL, CALCITRIOL  
CARISOPRODOL, CARISOPRODOL  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
DUTASTERIDE, DUTASTERIDE  
ERGOCALCIFEROL, ERGOCALCIFEROL  
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE  
IBUPROFEN AND DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)  
IBUPROFEN, IBUPROFEN  
IBUPROFEN, IBUPROFEN (OTC)  
IMIQUIMOD, IMIQUIMOD  
LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE  
MELOXICAM, MELOXICAM  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
METHOXSALEN, METHOXSALEN  
METRONIDAZOLE, METRONIDAZOLE  
MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL  
NEVIRAPINE, NEVIRAPINE  
OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS  
PEG-3350, POTASSIUM CHLORIDE, SODIUM BICARBONATE, SODIUM CHLORIDE, POLYETHYLENE GLYCOL  
POLYETHYLENE GLYCOL 3350, POLYETHYLENE GLYCOL 3350 (OTC)  
POTASSIUM CITRATE, POTASSIUM CITRATE  
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)  
TACROLIMUS, TACROLIMUS  
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

**STRONGBRIDGE US**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* STRONGBRIDGE US INC  
KEVEYIS, DICHLORPHENAMIDE

**SUCAMPO PHARMA LLC**

- \* SUCAMPO PHARMA AMERICAS LLC  
AMITIZA, LUBIPROSTONE

**SUN PHARM INDS**

- \* SUN PHARMACEUTICAL INDUSTRIES LTD  
CARBIDOPA AND LEVODOPA, CARBIDOPA  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
DESLORATADINE, DESLORATADINE  
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
EPINASTINE HYDROCHLORIDE, EPINASTINE HYDROCHLORIDE  
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM  
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE  
OLANZAPINE, OLANZAPINE  
ONDANSETRON, ONDANSETRON  
OXCARBAZEPINE, OXCARBAZEPINE  
SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE  
TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

**SUN PHARM INDS (IN)**

- \* SUN PHARMACEUTICAL INDUSTRIES LTD  
CEPHALEXIN, CEPHALEXIN  
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
ZONISAMIDE, ZONISAMIDE

**SUN PHARM INDS INC**

- \* SUN PHARMACEUTICAL INDUSTRIES INC  
ABSORICA, ISOTRETINOIN  
ALLOPURINOL, ALLOPURINOL  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
ATENOLOL, ATENOLOL  
BACLOFEN, BACLOFEN  
BENZONATATE, BENZONATATE  
CARVEDILOL, CARVEDILOL  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
CLONAZEPAM, CLONAZEPAM  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
CLOZAPINE, CLOZAPINE  
DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE  
DIGOXIN, DIGOXIN  
ERGOCALCIFEROL, ERGOCALCIFEROL  
EXELDERM, SULCONAZOLE NITRATE  
FLUMADINE, RIMANTADINE HYDROCHLORIDE  
FLURBIPROFEN, FLURBIPROFEN  
GEMFIBROZIL, GEMFIBROZIL  
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE  
GLIPIZIDE, GLIPIZIDE  
HALOG, HALCINONIDE  
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE  
INDOMETHACIN, INDOMETHACIN  
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE  
LITHIUM CARBONATE, LITHIUM CARBONATE  
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SUN PHARMACEUTICAL INDUSTRIES INC
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHIMAZOLE, METHIMAZOLE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MIRTAZAPINE, MIRTAZAPINE
  - NIMODIPINE, NIMODIPINE
  - OXaprozin, OXaprozin
  - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
  - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - REPAGLINIDE, REPAGLINIDE
  - RISPERIDONE, RISPERIDONE
  - TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - VENLAFAKINE HYDROCHLORIDE, VENLAFAKINE HYDROCHLORIDE
  - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**SUN PHARM INDS LTD**

- \* SUN PHARMACEUTICAL INDUSTRIES LTD
  - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - ACETAMINOPHEN, ACETAMINOPHEN (OTC)
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - AZITHROMYCIN, AZITHROMYCIN
  - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
  - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - CARVEDILOL, CARVEDILOL
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CETIRIZINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
  - CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - CLARITHROMYCIN, CLARITHROMYCIN
  - CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
  - CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
  - DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - DOXYCYCLINE, DOXYCYCLINE
  - FAMOTIDINE, FAMOTIDINE (OTC)
  - FELODIPINE, FELODIPINE
  - FENOFLIBRATE, FENOFLIBRATE
  - FLECAINIDE ACETATE, FLECAINIDE ACETATE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - GABAPENTIN, GABAPENTIN
  - GLYCOPYRRROLATE, GLYCOPYRRROLATE
  - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
  - LISINOPRIL AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE
  - LISINOPRIL, LISINOPRIL
  - LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
  - LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SUN PHARMACEUTICAL INDUSTRIES LTD
  - LORATADINE REDIDOSE, LORATADINE (OTC)
  - LORATADINE, LORATADINE (OTC)
  - LORAZEPAM, LORAZEPAM
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
  - NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
  - NARATRIPTAN, NARATRIPTAN HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON, ONDANSETRON
  - OPCICON ONE-STEP, LEVONORGESTREL (OTC)
  - OXCARBAZEPINE, OXCARBAZEPINE
  - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
  - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
  - RILUZOLE, RILUZOLE
  - RIOMET, METFORMIN HYDROCHLORIDE
  - RISPERIDONE, RISPERIDONE
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
  - TOPIRAMATE, TOPIRAMATE
  - ULTRAVATE, HALOBETASOL PROPIONATE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VALPROIC ACID, VALPROIC ACID
  - XIMINO, MINOCYCLINE HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**SUN PHARM INDUSTRIES**

- \* SUN PHARMACEUTICAL INDUSTRIES INC
  - ACETAZOLAMIDE, ACETAZOLAMIDE
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - ALLOPURINOL, ALLOPURINOL
  - ATENOLOL AND CHLORTHALIDONE, ATENOLOL
  - ATENOLOL, ATENOLOL
  - BACTRIM DS, SULFAMETHOXAZOLE
  - BACTRIM, SULFAMETHOXAZOLE
  - CARISOPRODOL, CARISOPRODOL
  - CARVEDILOL PHOSPHATE, CARVEDILOL PHOSPHATE
  - CHLORTHALIDONE, CHLORTHALIDONE
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
  - DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
  - DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
  - ERGOLOID MESYLATES, ERGOLOID MESYLATES
  - FELODIPINE, FELODIPINE
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
  - LEVETIRACETAM, LEVETIRACETAM
  - LOVASTATIN, LOVASTATIN
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MINOXIDIL, MINOXIDIL
  - MORPHINE SULFATE, MORPHINE SULFATE
  - NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
  - NYSTATIN, NYSTATIN
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PINDOLOL, PINDOLOL
  - PIROXICAM, PIROXICAM
  - PREDNISONE, PREDNISONE
  - PROPafenone HYDROCHLORIDE, PROPafenone HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SUN PHARMACEUTICAL INDUSTRIES INC
  - QUALAQUIN, QUININE SULFATE
  - QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
  - QUINIDINE SULFATE, QUINIDINE SULFATE
  - SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - SPIRONOLACTONE, SPIRONOLACTONE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
  - SULINDAC, SULINDAC
  - SYNALGOS-DC, ASPIRIN
  - TEMAZEPAM, TEMAZEPAM
  - THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
  - TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
  - ULTRAVATE, HALOBETASOL PROPIONATE

**SUN PHARMA GLOBAL**

- \* SUN PHARMA GLOBAL FZE
  - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - ALENDRONATE SODIUM, ALENDRONATE SODIUM
  - AMIFOSTINE, AMIFOSTINE
  - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
  - BROMSITE, BROMFENAC SODIUM
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CAFFEINE CITRATE, CAFFEINE CITRATE
  - CARBIDOPA AND LEVODOPA, CARBIDOPA
  - CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHLOROTHIAZIDE SODIUM, CHLOROTHIAZIDE SODIUM
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - DECITABINE, DECITABINE
  - DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
  - DESVENLAFAKINE, DESVENLAFAKINE FUMARATE
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DOXORUBICIN HYDROCHLORIDE (LIPOSOMAL), DOXORUBICIN HYDROCHLORIDE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ENTACAPONE, ENTACAPONE
  - ESOMEPRAZOLE SODIUM, ESOMEPRAZOLE SODIUM
  - ESZOPICLONE, ESZOPICLONE
  - FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE
  - FINASTERIDE, FINASTERIDE
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - IMATINIB MESYLATE, IMATINIB MESYLATE
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
  - LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - METHOTREXATE SODIUM, METHOTREXATE SODIUM
  - NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
  - NIACIN, NIACIN
  - NICARDIPIINE HYDROCHLORIDE, NICARDIPIINE HYDROCHLORIDE
  - ODOMZO, SONIDEGB PHOSPHATE
  - OXALIPLATIN, OXALIPLATIN
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RISEDRONATE SODIUM, RISEDRONATE SODIUM
  - ROSVUASTATIN CALCIUM, ROUVASTATIN CALCIUM
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - TETRABENAZINE, TETRABENAZINE
  - TOPIRAMATE, TOPIRAMATE
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - VECURONIUM BROMIDE, VECURONIUM BROMIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* S \*\*

- \* SUN PHARMA GLOBAL FZE  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE  
ZOLEDRONIC ACID, ZOLEDRONIC ACID  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- \* SUN PHARMA GLOBAL INC  
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE  
ALPRAZOLAM, ALPRAZOLAM  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
BICALUTAMIDE, BICALUTAMIDE  
CARBOPLATIN, CARBOPLATIN  
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM  
GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

**SUNOVION**

- \* SUNOVION PHARMACEUTICALS INC  
BROVANA, ARFORMOTEROL TARTRATE  
XOPENEX HFA, LEVALBUTEROL TARTRATE

**SUNOVION PHARMS INC**

- \* SUNOVION PHARMACEUTICALS INC  
APTIOM, ESLICARBAZEPINE ACETATE  
ARCAPTA NEOHALER, INDACATEROL MALEATE  
LATUDA, LURASIDONE HYDROCHLORIDE  
LUNESTA, ESZOPICLONE  
SEEBRI, GLYCOPYRROLATE  
UTIBRON, GLYCOPYRROLATE  
ZONEGRAN, ZONISAMIDE

**SUNOVION RESP**

- \* SUNOVION RESPIRATORY DEVELOPMENT INC  
LONHALA MAGNAIR KIT, GLYCOPYRROLATE

**SUNRISE PHARM INC**

- \* SUNRISE PHARMACEUTICAL INC  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE

**SUNSTAR AMERICAS**

- \* SUNSTAR AMERICAS INC  
PAROEX, CHLORHEXIDINE GLUCONATE

**SUPERNUS PHARMS**

- \* SUPERNUS PHARMACEUTICALS INC  
OXTELLAR XR, OXCARBAZEPINE  
TROKENDI XR, TOPIRAMATE

**SUVEN LIFE**

- \* SUVEN LIFE SCIENCES LTD  
MALATHION, MALATHION

**SVC PHARMA**

- \* SVC PHARMA LP  
DRONABINOL, DRONABINOL

**SWEDISH ORPHAN**

- \* SWEDISH ORPHAN BIOVITRUM AB PUBL  
ORFADIN, NITISINONE

**SYNERGY PHARMS**

- \* SYNERGY PHARMACEUTICALS INC  
TRULANCE, PLECANATIDE

**SYNTTHON PHARMS**

- \* SYNTTHON PHARMACEUTICALS INC  
LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE  
TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE

\*\* T \*\*

**ACME LABS**

- \* THE ACME LABORATORIES LTD  
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

**GEN HOSP**

- \* THE GENERAL HOSPITAL CORP  
AMMONIA N 13, AMMONIA N-13

**METHODIST HOSP RES**

- \* THE METHODIST HOSP RESEARCH INSTITUTE  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**RITEDOSE CORP**

- \* THE RITEDOSE CORP  
ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE  
ALBUTEROL SULFATE, ALBUTEROL SULFATE  
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE  
LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

**TAIHO ONCOLOGY**

- \* TAIHO ONCOLOGY INC  
LONSURF, TIPIRACIL HYDROCHLORIDE

**TAKEDA PHARMS USA**

- \* TAKEDA PHARMACEUTICALS USA INC  
ACTOPLUS MET XR, METFORMIN HYDROCHLORIDE  
ACTOPLUS MET, METFORMIN HYDROCHLORIDE  
ACTOS, PIOGLITAZONE HYDROCHLORIDE  
COLCRYS, COLCHICINE  
DEXILANT, DEXLANSOPRAZOLE  
DUETACT, GLIMEPIRIDE  
KAZANO, ALOGLIPTIN BENZOATE  
NESINA, ALOGLIPTIN BENZOATE  
OSENI, ALOGLIPTIN BENZOATE  
PREVACID, LANSOPRAZOLE  
PREVPAC, AMOXICILLIN  
ROZEREM, RAMELTEON  
TRINTELLIX, VORTioxETINE HYDROBROMIDE  
ULORIC, FEBUXOSTAT

**TALON THERAP**

- \* TALON THERAPEUTICS INC  
MARQIBO KIT, VINCRISTINE SULFATE

**TAMARANG**

- \* TAMARANG SA  
ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

**TARO**

- \* TARO PHARMACEUTICAL INDUSTRIES LTD  
ACETAZOLAMIDE, ACETAZOLAMIDE  
CARBAMAZEPINE, CARBAMAZEPINE  
CARVEDILOL, CARVEDILOL  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE  
CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE  
ENALAPRIL MALEATE, ENALAPRIL MALEATE  
ETODOLAC, ETODOLAC  
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM  
FLUCONAZOLE, FLUCONAZOLE  
FLUOROURACIL, FLUOROURACIL  
GABAPENTIN, GABAPENTIN  
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE  
IMIQUIMOD, IMIQUIMOD  
KETOCONAZOLE, KETOCONAZOLE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
LORATADINE, LORATADINE (OTC)  
MELOXICAM, MELOXICAM  
METRONIDAZOLE, METRONIDAZOLE  
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
OXCARBAZEPINE, OXCARBAZEPINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TARO PHARMACEUTICAL INDUSTRIES LTD  
PHENYTOIN, PHENYTOIN
- \* TARO PHARMACEUTICALS USA INC  
ACETIC ACID, ACETIC ACID, GLACIAL  
ACYCLOVIR, ACYCLOVIR  
ADAPALENE, ADAPALENE  
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE  
AMMONIUM LACTATE, AMMONIUM LACTATE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
BUTENAFINE HYDROCHLORIDE, BUTENAFINE HYDROCHLORIDE (OTC)  
CICLOPIROX, CICLOPIROX  
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CLOTRIMAZOLE, CLOTRIMAZOLE  
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)  
DAPSONE, DAPSONE  
DERMABET, BETAMETHASONE VALERATE  
DESONIDE, DESONIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE  
ECONAZOLE NITRATE, ECONAZOLE NITRATE  
ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE  
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE  
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE  
FLUOCINONIDE, FLUOCINONIDE  
GENTAMICIN SULFATE, GENTAMICIN SULFATE  
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE  
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL  
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE  
HYDROCORTISONE, HYDROCORTISONE  
IBUPROFEN, IBUPROFEN  
IBUPROFEN, IBUPROFEN (OTC)  
KETOZOLE, KETOCONAZOLE  
LIDOCAINE, LIDOCAINE  
LORATADINE, LORATADINE (OTC)  
MICONAZOLE 3, MICONAZOLE NITRATE (OTC)  
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)  
MOMETASONE FUROATE, MOMETASONE FUROATE  
MUPIROCIN, MUPIROCIN  
NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE  
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN  
NYSTATIN, NYSTATIN  
OXICONAZOLE NITRATE, OXICONAZOLE NITRATE  
PHENYTOIN, PHENYTOIN  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
RISPERIDONE, RISPERIDONE  
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE  
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, STERILE WATER FOR INJECTION  
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM  
TAZAROTENE, TAZAROTENE  
TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE (OTC)  
TERCONAZOLE, TERCONAZOLE  
TOPICORT, DESOXIMETASONE  
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE  
TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)  
U-CORT, HYDROCORTISONE ACETATE

**TARO PHARM**

- \* TARO PHARMACEUTICAL INDUSTRIES LTD  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE  
BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN  
CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TARO PHARMACEUTICAL INDUSTRIES LTD
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
  - DESLORATADINE, DESLORATADINE
  - DESONIDE, DESONIDE
  - FELBAMATE, FELBAMATE
  - FLUOROURACIL, FLUOROURACIL
  - GABAPENTIN, GABAPENTIN
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
  - LORATADINE, LORATADINE (OTC)
  - METRONIDAZOLE, METRONIDAZOLE
  - NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
  - NYSTATIN, NYSTATIN
  - OVIDE, MALATHION
  - TERIL, CARBAMAZEPINE
  - WARFARIN SODIUM, WARFARIN SODIUM

**TARO PHARM INDS**

- \* TARO PHARMACEUTICAL INDUSTRIES LTD
  - AMCINONIDE, AMCINONIDE
  - CARBAMAZEPINE, CARBAMAZEPINE
  - CICLOPIROX, CICLOPIROX
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - DESLORATADINE, DESLORATADINE
  - ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
  - ETODOLAC, ETODOLAC
  - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
  - LAMOTRIGINE, LAMOTRIGINE
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE

**TARO PHARMS**

- \* TARO PHARMACEUTICALS INC
  - PLIAGLIS, LIDOCAINE

**TARO PHARMS NORTH**

- \* TARO PHARMACEUTICALS NORTH AMERICA INC
  - ACETAMINOPHEN, ACETAMINOPHEN (OTC)
  - INFANTS' FEVERALL, ACETAMINOPHEN (OTC)

**TASMAN PHARMA**

- \* TASMAN PHARMA INC
  - VERSACLOZ, CLOZAPINE

**TCG FLUENT PHARMA**

- \* TCG FLUENT PHARMA INVESTORS LP
  - FLOLIPID, SIMVASTATIN

**TECH ORGANIZED**

- \* TECHNOLOGY ORGANIZED LLC
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - NEVIRAPINE, NEVIRAPINE

**TEDOR PHARM**

- \* TEDOR PHARMA INC
  - BENZPHETAMINE HYDROCHLORIDE, BENZPHETAMINE HYDROCHLORIDE

**TEDOR PHARMA INC**

- \* TEDOR PHARMA INC
  - BUTALBITAL AND ACETAMINOPHEN, ACETAMINOPHEN
  - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE

**TEIKOKU PHARMA USA**

- \* TEIKOKU PHARMA USA INC
  - LIDODERM, LIDOCAINE

**TELIGENT**

- \* TELIGENT OU
  - CEFOTAN, CEFOTETAN DISODIUM
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TELIGENT OU
  - FORTAZ IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
  - FORTAZ, CEFTAZIDIME
  - ZANTAC, RANITIDINE HYDROCHLORIDE
  - ZINACEF IN PLASTIC CONTAINER, CEFUROXIME SODIUM
  - ZINACEF, CEFUROXIME SODIUM

**TELIGENT PHARMA INC**

- \* TELIGENT PHARMA INC
  - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - DESONIDE, DESONIDE
  - DESOXIMETASONE, DESOXIMETASONE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - ECONAZOLE NITRATE, ECONAZOLE NITRATE
  - ERYTHROMYCIN, ERYTHROMYCIN
  - FLURANDRENOLIDE, FLURANDRENOLIDE
  - HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
  - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
  - LIDOCAINE, LIDOCAINE
  - NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

**TERSCERA THERAPS LLC**

- \* TERSCERA THERAPEUTICS LLC
  - ERGOMAR, ERGOTAMINE TARTRATE
  - ZOLADEX, GOSERELIN ACETATE

**TESARO INC**

- \* TESARO INC
  - VARUBI, ROLAPITANT HYDROCHLORIDE
  - ZEJULA, NIRAPARIB TOSYLATE

**TEVA**

- \* TEVA NEUROSCIENCE INC
  - AZILECT, RASAGILINE MESYLATE
- \* TEVA PHARMACEUTICALS USA INC
  - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - ACYCLOVIR, ACYCLOVIR
  - ADIPEX-P, PHENTERMINE HYDROCHLORIDE
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
  - AMOXICILLIN PEDIATRIC, AMOXICILLIN
  - AMOXICILLIN, AMOXICILLIN
  - ATENOLOL, ATENOLOL
  - AZITHROMYCIN, AZITHROMYCIN
  - BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
  - BICALUTAMIDE, BICALUTAMIDE
  - BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
  - CALCITRIOL, CALCITRIOL
  - CAPTOPRIL, CAPTOPRIL
  - CARVEDILOL, CARVEDILOL
  - CEFACLOR, CEFACLOR
  - CEFPROZIL, CEFPROZIL
  - CELECOXIB, CELECOXIB
  - CEPHALEXIN, CEPHALEXIN
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
  - CILOSTAZOL, CILOSTAZOL
  - CIMETIDINE, CIMETIDINE
  - CLARITHROMYCIN, CLARITHROMYCIN
  - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
  - CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
  - CLONAZEPAM, CLONAZEPAM
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TEVA PHARMACEUTICALS USA INC  
CLOTRIMAZOLE, CLOTRIMAZOLE  
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE  
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM  
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM  
DIFLUNISAL, DIFLUNISAL  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE  
ENALAPRIL MALEATE, ENALAPRIL MALEATE  
ENOXAPARIN SODIUM (PRESERVATIVE FREE), ENOXAPARIN SODIUM  
EPITOL, CARBAMAZEPINE  
ESZOPICLONE, ESZOPICLONE  
ETODOLAC, ETODOLAC  
FAMOTIDINE, FAMOTIDINE  
FAMOTIDINE, FAMOTIDINE (OTC)  
FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)  
FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)  
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE  
FINASTERIDE, FINASTERIDE  
FLUCONAZOLE, FLUCONAZOLE  
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE  
FLUOCINONIDE, FLUOCINONIDE  
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE  
FLURBIPROFEN, FLURBIPROFEN  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM  
GALZIN, ZINC ACETATE  
GEMFIBROZIL, GEMFIBROZIL  
GLIMEPIRIDE, GLIMEPIRIDE  
GLYBURIDE (MICRONIZED), GLYBURIDE  
GLYBURIDE, GLYBURIDE  
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE  
IRBESARTAN AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE  
KETOCONAZOLE, KETOCONAZOLE  
KETOPROFEN, KETOPROFEN  
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE  
LAMOTRIGINE, LAMOTRIGINE  
LEVOFLOXACIN, LEVOFLOXACIN  
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE  
LORATADINE, LORATADINE (OTC)  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
LOVASTATIN, LOVASTATIN  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
METOPROLOL TARTRATE, METOPROLOL TARTRATE  
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE  
MIRTAZAPINE, MIRTAZAPINE  
MOEXIPRIL HYDROCHLORIDE AND HYDROCHLORTHIAZIDE, HYDROCHLORTHIAZIDE  
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE  
MUPIROCIN, MUPIROCIN  
NAPROXEN SODIUM, NAPROXEN SODIUM  
NAPROXEN, NAPROXEN  
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE  
NEOMYCIN SULFATE, NEOMYCIN SULFATE  
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE  
NYSTATIN, NYSTATIN  
OFLOXACIN, OFLOXACIN  
ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE  
ONDANSETRON, ONDANSETRON  
ORAP, PIMOZIDE  
OXaprozin, OXaprozin

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TEVA PHARMACEUTICALS USA INC
  - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
  - PENICILLIN-VK, PENICILLIN V POTASSIUM
  - PIROXICAM, PIROXICAM
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PREDNISONE, PREDNISOLONE
  - QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RIBAVIRIN, RIBAVIRIN
  - RISPERIDONE, RISPERIDONE
  - ROSIGLITAZONE MALEATE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - ROSIGLITAZONE MALEATE, ROSIGLITAZONE MALEATE
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
  - SUCRALFATE, SUCRALFATE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
  - TERBINAFINE HYDROCHLORIDE, TERBINAFINE HYDROCHLORIDE
  - TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TOLMETIN SODIUM, TOLMETIN SODIUM
  - TOPIRAMATE, TOPIRAMATE
  - TORSEMIDE, TORSEMIDE
  - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**TEVA BRANDED PHARM**

- \* TEVA BRANDED PHARMACEUTICAL PRODUCTS R AND D INC
  - AUSTEDO, DEUTETRABENAZINE
  - DIAMOX, ACETAZOLAMIDE
  - LOSEASONIQUE, ETHINYLMESTRADIOL
  - PROAIR HFA, ALBUTEROL SULFATE
  - PROAIR RESPICLICK, ALBUTEROL SULFATE
  - PROGLYCEM, DIAZOXIDE
  - QNASL, BECLOMETHASONE DIPROPIONATE
  - QUARTETTE, ETHINYLMESTRADIOL
  - QVAR 40, BECLOMETHASONE DIPROPIONATE
  - QVAR 80, BECLOMETHASONE DIPROPIONATE
  - SEASONALE, ETHINYLMESTRADIOL
  - SEASONIQUE, ETHINYLMESTRADIOL
  - VANTRELA ER, HYDROCODONE BITARTRATE
  - ZIAC, BISOPROLOL FUMARATE

**TEVA PARENTERAL**

- \* TEVA PARENTERAL MEDICINES INC
  - DAPTOMYCIN, DAPTOMYCIN
  - LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE

**TEVA PHARM**

- \* TEVA PHARMACEUTICAL INDUSTRIES LTD
  - AIRDUO RESPICLICK, FLUTICASONE PROPIONATE
  - ARMONAIR RESPICLICK, FLUTICASONE PROPIONATE

**TEVA PHARMS**

- \* TEVA PHARMACEUTICALS USA
  - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
  - ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLORTIAZIDE, AMLODIPINE BESYLATE
  - ANASTROZOLE, ANASTROZOLE
  - AZITHROMYCIN, AZITHROMYCIN
  - BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
  - BUDESONIDE, BUDESONIDE
  - CARBAMAZEPINE, CARBAMAZEPINE
  - CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TEVA PHARMACEUTICALS USA
  - CEFDINIR, CEFdinir
  - CEFPROZIL, CEFprozil
  - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - CROMOLYN SODIUM, CROMOLYN SODIUM
  - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
  - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
  - DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
  - DOXEPIH HYDROCHLORIDE, DOXEPIH HYDROCHLORIDE
  - ETHOSUXIMIDE, ETHOSUXIMIDE
  - FAMCICLOVIR, FAMCICLOVIR
  - FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
  - GABAPENTIN, GABAPENTIN
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - GEMCITABINE HYDROCHLORIDE, GEMCITABINE HYDROCHLORIDE
  - GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
  - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - HYDROCORTISONE, HYDROCORTISONE
  - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
  - IRBESARTAN, IRBESARTAN
  - LAMIVUDINE AND ZIDOVUDINE, LAMIVUDINE
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LEFLUNOMIDE, LEFLUNOMIDE
  - LETROZOLE, LETROZOLE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOCETIRIZINE DIHYDROCHLORIDE, LEVOCETIRIZINE DIHYDROCHLORIDE
  - LINEZOLID, LINEZOLID
  - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - MEGESTROL ACETATE, MEGESTROL ACETATE
  - MELOXICAM, MELOXICAM
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
  - NARatriptan, NARatriptan HYDROCHLORIDE
  - OLANZAPINE AND FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - OLANZAPINE, OLANZAPINE
  - OXALIPLATIN, OXALIPLATIN
  - PACLITAXEL, PACLITAXEL
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
  - PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
  - PROGESTERONE, PROGESTERONE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - QUININE SULFATE, QUININE SULFATE
  - RAMIPRIL, RAMIPRIL
  - RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE
  - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TRANDOLAPRIL, TRANDOLAPRIL
  - URSODIOL, URSDIOL
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VANDAZOLE, METRONIDAZOLE
  - VORICONAZOLE, VORICONAZOLE
  - ZALEPLON, ZALEPLON

**TEVA PHARMS INTL**

- \* TEVA PHARMACEUTICALS INTERNATIONAL GMBH
  - AMRIX, CYCLOBENZAPRINE HYDROCHLORIDE
  - SYNRIBO, OMACETAXINE MEPESUCCINATE

**TEVA PHARMS USA**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TEVA PHARMACEUTICALS USA
  - ABACAVIR SULFATE AND LAMIVUDINE, ABACAVIR SULFATE
  - ACITRETIN, ACITRETIN
  - ADENOSINE, ADENOSINE
  - ALMOTRIPTAN MALATE, ALMOTRIPTAN MALATE
  - ALPROSTADIL, ALPROSTADIL
  - AMIKACIN SULFATE, AMIKACIN SULFATE
  - AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE
  - AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE
  - ARGATROBAN IN 0.9% SODIUM CHLORIDE, ARGATROBAN
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - ATAZANAVIR SULFATE, ATAZANAVIR SULFATE
  - ATOMOXETINE HYDROCHLORIDE, ATOMOXETINE HYDROCHLORIDE
  - ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
  - BUDESONIDE, BUDESONIDE
  - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - CARBOPLATIN, CARBOPLATIN
  - CLARAVIS, ISOTRETINOIN
  - CLOZAPINE, CLOZAPINE
  - COPAXONE, GLATIRAMER ACETATE
  - DACARBAZINE, DACARBAZINE
  - DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
  - DOCETAXEL, DOCETAXEL
  - DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ENALAPRILAT, ENALAPRILAT
  - ENTECAVIR, ENTECAVIR
  - EPOPROSTENOL SODIUM, EPOPROSTENOL SODIUM
  - EPTIFIBATIDE, EPTIFIBATIDE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ESTRADIOL, ESTRADIOL
  - ETOPOSIDE, ETOPOSIDE
  - EZETIMIBE, EZETIMIBE
  - FLUOROURACIL, FLUOROURACIL
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUVASTATIN SODIUM, FLUVASTATIN SODIUM
  - GABAPENTIN, GABAPENTIN
  - GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE
  - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
  - HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
  - HALOPERIDOL, HALOPERIDOL LACTATE
  - IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
  - IFOSFAMIDE, IFOSFAMIDE
  - IMATINIB MESYLATE, IMATINIB MESYLATE
  - IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE
  - LANSOPRAZOLE, AMOXICILLIN AND CLARITHROMYcin, AMOXICILLIN
  - LANSOPRAZOLE, LANSOPRAZOLE
  - LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
  - LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
  - LEVALBUTEROL HYDROCHLORIDE, LEVALBUTEROL HYDROCHLORIDE
  - LINEZOLID, LINEZOLID
  - LOGILIA, ULIPIRISTAL ACETATE
  - MESNA, MESNA
  - METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
  - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
  - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - MYCOPHENOLIC ACID, MYCOPHENOLIC ACID
  - NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
  - OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
  - OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TEVA PHARMACEUTICALS USA
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - OMEGA-3-ACID ETHYL ESTERS, OMEGA-3-ACID ETHYL ESTERS
  - OMEPRAZOLE, OMEPRAZOLE
  - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
  - PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
  - PARICALCITOL, PARICALCITOL
  - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
  - RISEDRONATE SODIUM, RISEDRONATE SODIUM
  - ROSIGLITAZONE MALEATE AND GLIMEPIRIDE, GLIMEPIRIDE
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - SOLIFENACIN SUCCINATE, SOLIFENACIN SUCCINATE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TENOFOVIR DISOPROXIL FUMARATE, TENOFOVIR DISOPROXIL FUMARATE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - TOBRAMYCIN, TOBRAMYCIN
  - TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
  - TOPOTECAN HYDROCHLORIDE, TOPOTECAN HYDROCHLORIDE
  - VECURONIUM BROMIDE, VECURONIUM BROMIDE
  - VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
  - VINORELBINE TARTRATE, VINORELBINE TARTRATE
  - ZANOSAR, STREPTOZOCIN
  - ZOLMITRIPTAN, ZOLMITRIPTAN
- \* TEVA PHARMACEUTICALS USA INC
  - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
  - CAPECITABINE, CAPECITABINE
  - DARUNAVIR ETHANOLATE, DARUNAVIR ETHANOLATE
  - ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
  - EMTRICITABINE AND TENOFOVIR DISOPROXIL FUMARATE, EMTRICITABINE
  - ESTRADIOL AND NORETHINDRONE ACETATE, ESTRADIOL
  - MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
  - METRONIDAZOLE, METRONIDAZOLE
  - OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - SELFEMRA, FLUOXETINE HYDROCHLORIDE
  - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE

**THE FEINSTEIN INST**

- \* THE FEINSTEIN INSTITUTE FOR MEDICAL RESEARCH
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**THE MEDICINES CO**

- \* THE MEDICINES CO
  - ANGIOMAX, BIVALIRUDIN
  - IONSYS, FENTANYL HYDROCHLORIDE
  - ORBACTIV, ORITAVANCIN DIPHOSPHATE

**THE PHARMA NETWORK**

- \* THE PHARMA NETWORK LLC
  - BENZONATATE, BENZONATATE

**THE PHARMANETWORK**

- \* THE PHARMANETWORK LLC
  - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE

**THEPHARMANETWORK LLC**

- \* THEPHARMANETWORK LLC
  - BENZONATATE, BENZONATATE
  - ISONIAZID, ISONIAZID
  - NIMODIPINE, NIMODIPINE
  - THERMAZENE, SILVER SULFADIAZINE

**THERATECHNOLOGIES**

- \* THERATECHNOLOGIES INC
  - EGRIFTA, TESAMORELIN ACETATE

**THERAVANCE BIOPHARMA**

- \* THERAVANCE BIOPHARMA R AND D INC

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* THERAVANCE BIOPHARMA R AND D INC  
VIBATIV, TELAVANCIN HYDROCHLORIDE

**TOLMAR**

- \* TOLMAR INC  
ACYCLOVIR, ACYCLOVIR  
ADAPALENE, ADAPALENE  
ATRIDOX, DOXYCYCLINE HYCLATE  
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CALCIPOTRIENE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE  
CALCIPOTRIENE, CALCIPOTRIENE  
CICLOPIROX, CICLOPIROX  
CLINDAMYCIN PHOSPHATE AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
IMIQUIMOD, IMIQUIMOD  
KETOCONAZOLE, KETOCONAZOLE  
LEVETIRACETAM, LEVETIRACETAM  
LIDOCAINE AND PRILOCAINE, LIDOCAINE  
METRONIDAZOLE, METRONIDAZOLE  
MOMETASONE FUROATE, MOMETASONE FUROATE  
NAFTIFINE HYDROCHLORIDE, NAFTIFINE HYDROCHLORIDE  
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

**TOLMAR THERAP**

- \* TOLMAR THERAPEUTICS INC  
ELIGARD, LEUPROLIDE ACETATE

**TORPHARM**

- \* TORPHARM INC  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

**TORRENT PHARMA INC**

- \* TORRENT PHARMA INC  
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

**TORRENT PHARMS**

- \* TORRENT PHARMACEUTICALS LIMITED  
LEVOFLOXACIN, LEVOFLOXACIN
- \* TORRENT PHARMACEUTICALS LTD  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
CARBAMAZEPINE, CARBAMAZEPINE  
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE  
LAMOTRIGINE, LAMOTRIGINE  
LEVETIRACETAM, LEVETIRACETAM  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM  
PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE  
RISPERIDONE, RISPERIDONE  
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE  
TOPIRAMATE, TOPIRAMATE  
ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
- \* TORRENT PHARMACEUTICALS LTD.  
ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE

**TORRENT PHARMS LLC**

- \* TORRENT PHARMACEUTICALS LLC  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)  
OLANZAPINE, OLANZAPINE  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

**TORRENT PHARMS LTD**

- \* TORRENT PHARMACEUTICALS LTD  
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE AND VALSARTAN, AMLODIPINE BESYLATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

- \* TORRENT PHARMACEUTICALS LTD
  - AMLODIPINE BESYLATE, VALSARTAN AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
  - ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
  - ARIPIPRAZOLE, ARIPIPRAZOLE
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CELECOXIB, CELECOXIB
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - DARIFENACIN HYDROBROMIDE, DARIFENACIN HYDROBROMIDE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ESOMEPRAZOLE MAGNESIUM, ESOMEPRAZOLE MAGNESIUM
  - FELODIPINE, FELODIPINE
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
  - LAMOTRIGINE, LAMOTRIGINE
  - LEVETIRACETAM, LEVETIRACETAM
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - OLANZAPINE, OLANZAPINE
  - OLMESARTAN MEDOXOMIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - OLMESARTAN MEDOXOMIL, AMLODIPINE AND HYDROCHLOROTHIAZIDE, AMLODIPINE BESYLATE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - PIOGLITAZONE HYDROCHLORIDE AND METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RABEPRAZOLE SODIUM, RABEPRAZOLE SODIUM
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - TELMISARTAN AND AMLODIPINE, AMLODIPINE BESYLATE
  - TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TELMISARTAN, TELMISARTAN
  - TOLTERODINE TARTRATE, TOLTERODINE TARTRATE
  - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
  - VALSARTAN, VALSARTAN

**TRIS PHARMA INC**

- \* TRIS PHARMA INC
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF, CETIRIZINE HYDROCHLORIDE (OTC)
  - DEXMETHYLPHENIDATE HYDROCHLORIDE, DEXMETHYLPHENIDATE HYDROCHLORIDE
  - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
  - DEXTROMETHORPHAN POLISTIREX, DEXTROMETHORPHAN POLISTIREX (OTC)
  - DYANAVEL XR, AMPHETAMINE
  - GABAPENTIN, GABAPENTIN
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - HYDROCODONE POLISTIREX AND CHLORPHENIRAMINE POLISTIREX, CHLORPHENIRAMINE POLISTIREX
  - IBUPROFEN, IBUPROFEN (OTC)
  - KARBINAL ER, CARBINOXAMINE MALEATE
  - LEVETIRACETAM, LEVETIRACETAM
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - MORPHINE SULFATE, MORPHINE SULFATE
  - POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
  - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - RISPERIDONE, RISPERIDONE
  - THEOPHYLLINE, THEOPHYLLINE

**TRUSTEES UNIV PA**

- \* TRUSTEES OF THE UNIV OF PENNSYLVANIA
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**TWI PHARMS INC**

- \* TWI PHARMACEUTICALS INC
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* T \*\*

\* TWI PHARMACEUTICALS INC  
 MEGESTROL ACETATE, MEGESTROL ACETATE  
 NIFEDIPINE, NIFEDIPINE

\*\* U \*\*

**UCB INC**

\* UCB INC  
 BRIVIACT, BRIVARACETAM  
 KEPPTRA XR, LEVETIRACETAM  
 KEPPTRA, LEVETIRACETAM  
 METADATE CD, METHYLPHENIDATE HYDROCHLORIDE  
 METADATE ER, METHYLPHENIDATE HYDROCHLORIDE  
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE  
 NEUPRO, ROTIGOTINE  
 TUSSIONEX PENNKinetic, CHLORPHENIRAMINE POLISTIREX  
 VIMPAT, LACOSAMIDE  
 ZAROXOLYN, METOLAZONE

**UCLA BIOMEDICAL**

\* UCLA BIOMEDICAL CYCLOTRON  
 AMMONIA N 13, AMMONIA N-13  
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**UCSF RODIOPHARM**

\* UCSF RADIOPHARMACEUTICAL FACILITY  
 AMMONIA N 13, AMMONIA N-13  
 CHOLINE C-11, CHOLINE C-11  
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
 SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**UIHC PET IMAGING**

\* UNIV IOWA HOSPS AND CLINICS PET IMAGING CENTER  
 FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**UNICHEM**

\* UNICHEM LABORATORIES LTD  
 BISOPROLOL FUMARATE AND HYDROCHLORTIAZIDE, BISOPROLOL FUMARATE  
 CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
 CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)  
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
 HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE  
 MELOXICAM, MELOXICAM  
 ZALEPLON, ZALEPLON

**UNICHEM LABS LTD**

\* UNICHEM LABORATORIES LIMITED  
 DIVALPROEX SODIUM, DIVALPROEX SODIUM  
 \* UNICHEM LABORATORIES LTD  
 ALFUZOSIN HYDROCHLORIDE, ALFUZOSIN HYDROCHLORIDE  
 AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
 DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
 IRBESARTAN AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE  
 IRBESARTAN, IRBESARTAN  
 LAMOTRIGINE, LAMOTRIGINE  
 LOSARTAN POTASSIUM AND HYDROCHLORTIAZIDE, HYDROCHLORTIAZIDE  
 LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
 MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
 METRONIDAZOLE, METRONIDAZOLE  
 MONTELUKAST SODIUM, MONTELUKAST SODIUM  
 PIROXICAM, PIROXICAM  
 QUETIAPINE FUMARATE, QUETIAPINE FUMARATE  
 RIZATRIPTAN BENZOATE, RIZATRIPTAN BENZOATE  
 TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE  
 TOPIRAMATE, TOPIRAMATE

**UNICHEM PHARMS (USA)**

\* UNICHEM PHARMACEUTICALS (USA) INC  
 BISOPROLOL FUMARATE, BISOPROLOL FUMARATE

**UNIMARK REMEDIES LTD**

\* UNIMARK REMEDIES LTD  
 MONTELUKAST SODIUM, MONTELUKAST SODIUM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* U \*\*

**UNIQUE PHARM LABS**

- \* UNIQUE PHARMACEUTICAL LABORATORIES A DIVISION OF J.B. CHEMICALS AND PHARMACEUTICALS LTD  
ATENOLOL, ATENOLOL  
CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)  
CETIRIZINE HYDROCHLORIDE HIVES, CETIRIZINE HYDROCHLORIDE (OTC)  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
DICLOFENAC SODIUM, DICLOFENAC SODIUM  
FLUCONAZOLE, FLUCONAZOLE  
GLIPIZIDE, GLIPIZIDE  
LITHIUM CARBONATE, LITHIUM CARBONATE  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE  
TINIDAZOLE, TINIDAZOLE

**UNITED BIOMEDCL**

- \* UNITED BIOMEDICAL INC  
TERBUTALINE SULFATE, TERBUTALINE SULFATE

**UNITED GUARDIAN**

- \* UNITED GUARDIAN INC  
RENACIDIN, CITRIC ACID

**UNITED THERAP**

- \* UNITED THERAPEUTICS CORP  
ORENITRAM, TREPROSTINIL DIOLAMINE  
REMODULIN, TREPROSTINIL  
TYVASO, TREPROSTINIL

**UNIV MICHIGAN**

- \* UNIV MICHIGAN PET RADIOPHARMACEUTICAL PRODUCTION PROGRAM  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**UNIV NORTH DAKOTA**

- \* UNIV NORTH DAKOTA  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**UNIV TX MD ANDERSON**

- \* UNIV TEXAS MD ANDERSON CANCER CENTER  
AMMONIA N 13, AMMONIA N-13  
CHOLINE C-11, CHOLINE C-11  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**UNIV UTAH CYCLOTRON**

- \* UNIV UTAH CYCLOTRON RADIOCHEMISTRY LAB  
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18  
SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**UPSHER-SMITH LABS**

- \* UPSHER-SMITH LABORATORIES LLC  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
EXEMESTANE, EXEMESTANE  
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE  
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM  
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
KLOR-CON M10, POTASSIUM CHLORIDE  
KLOR-CON M15, POTASSIUM CHLORIDE  
KLOR-CON M20, POTASSIUM CHLORIDE  
KLOR-CON, POTASSIUM CHLORIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE  
MIRTAZAPINE, MIRTAZAPINE  
MORPHINE SULFATE, MORPHINE SULFATE  
NYSTATIN, NYSTATIN  
ORVATEN, MIDODRINE HYDROCHLORIDE  
OXANDROLONE, OXANDROLONE  
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE  
PACERONE, AMIODARONE HYDROCHLORIDE  
PENTOXIL, PENTOXIFYLLINE  
PREVALITE, CHOLESTYRAMINE  
QUDEXY XR, TOPIRAMATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* U \*\*

- \* UPSHER-SMITH LABORATORIES LLC
  - SORINE, SOTALOL HYDROCHLORIDE
  - SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
  - TOPIRAMATE, TOPIRAMATE
  - VOGELXO, TESTOSTERONE

**US PHARM HOLDINGS**

- \* US PHARMACEUTICAL HOLDINGS II LLC
  - CLAFORAN, CEFOTAXIME SODIUM
  - DEMEROL, MEPERIDINE HYDROCHLORIDE
  - DRISDOL, ERGOCALCIFEROL
  - HIPREX, METHENAMINE HIPPURATE
  - LASIX, FUROSEMIDE
  - NORPRAMIN, DESIPRAMINE HYDROCHLORIDE

**US PHARMS HOLDINGS I**

- \* US PHARMACEUTICALS HOLDINGS I LLC
  - LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
  - LOPRESSOR, METOPROLOL TARTRATE
  - LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
  - LOTENSIN, BENAZEPRIL HYDROCHLORIDE
  - PARLODEL, BROMOCRIPTINE MESYLATE

**US WORLDMEDS**

- \* US WORLDMEDS LLC
  - APOKYN, APOMORPHINE HYDROCHLORIDE
  - REVONTO, DANTROLENE SODIUM

**US WORLDMEDS LLC**

- \* US WORLDMEDS LLC
  - CORGARD, NADOLOL
  - XADAGO, SAFINAMIDE MESYLATE

**USL PHARMA**

- \* USL PHARMA LLC
  - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
  - BACLOFEN, BACLOFEN
  - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
  - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
  - FLUOXYMESTERONE, FLUOXYMESTERONE
  - JANTOVEN, WARFARIN SODIUM

**USPHARMA**

- \* USPHARMA LTD
  - NITRO-DUR, NITROGLYCERIN

**USV NORTH AMERICA**

- \* USV NORTH AMERICA INC
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - ZOLEDRONIC ACID, ZOLEDRONIC ACID

\*\* V \*\*

**VALEANT**

- \* VALEANT PHARMACEUTICALS INTERNATIONAL
  - ANCOBON, FLUCYTOSINE
  - BONTRIL PDM, PHENDIMETRAZINE TARTRATE
  - D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
  - MIGRAL, DIHYDROERGOTAMINE MESYLATE
  - mysoline, PRIMIDONE

**VALEANT BERMUDA**

- \* VALEANT INTERNATIONAL BERMUDA
  - BENZACLIN, BENZOYL PEROXIDE
  - DERMATOP E EMOLIENT, PREDNICARBATE
  - ELIDEL, PIMECROLIMUS
  - PENLAC, CICLOPIROX
  - RETIN-A, TRETINOIN
  - XERESE, ACYCLOVIR
  - ZOVIRAX, ACYCLOVIR

**VALEANT INTL**

- \* VALEANT INTERNATIONAL BARBADOS SRL
  - ATIVAN, LORAZEPAM

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* V \*\*

- \* VALEANT INTERNATIONAL BARBADOS SRL  
CARDIZEM CD, DILTIAZEM HYDROCHLORIDE  
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE  
CARDIZEM, DILTIAZEM HYDROCHLORIDE  
RETIN-A MICRO, TRETINOIN  
RETIN-A, TRETINOIN  
RETIN-A-MICRO, TRETINOIN  
VASERETIC, ENALAPRIL MALEATE  
WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
- \* VALEANT INTERNATIONAL SRL  
BENZAMYCIN, BENZOYL PEROXIDE

**VALEANT LUXEMBOURG**

- \* VALEANT PHARMACEUTICALS LUXEMBOURG SARL  
ERTACZO, SERTACONAZOLE NITRATE  
TARGRETIN, BEXAROTENE  
VISUDYNE, VERTEPORFIN

**VALEANT PHARM INTL**

- \* VALEANT PHARMACEUTICALS INTERNATIONAL  
ANDROID 25, METHYLTESTOSTERONE  
EFUDEX, FLUOROURACIL  
LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE  
MESTINON, PYRIDOSTIGMINE BROMIDE  
TESTRED, METHYLTESTOSTERONE  
VIRAZOLE, RIBAVIRIN  
ZELAPAR, SELEGILINE HYDROCHLORIDE

**VALEANT PHARMS**

- \* VALEANT PHARMACEUTICALS NORTH AMERICA  
MEPHYTON, PHYTONADIONE
- \* VALEANT PHARMACEUTICALS NORTH AMERICA LLC  
LIBRAX, CHLORDIAZEPOXIDE HYDROCHLORIDE  
MESTINON, PYRIDOSTIGMINE BROMIDE  
MINITRAN, NITROGLYCERIN  
PENTOXIFYLLINE, PENTOXIFYLLINE

**VALEANT PHARMS INC**

- \* VALEANT PHARMACEUTICALS INTERNATIONAL INC  
GRIS-PEG, GRISEOFULVIN, ULTRAMICROSIZE

**VALEANT PHARMS INTL**

- \* VALEANT PHARMACEUTICALS INTERNATIONAL  
APRISO, MESALAMINE  
COLAZAL, BALSALAZIDE DISODIUM  
GIAZO, BALSALAZIDE DISODIUM  
UCERIS, BUDESONIDE

**VALEANT PHARMS LLC**

- \* VALEANT PHARMACEUTICALS NORTH AMERICA LLC  
MACUGEN, PEGAPTANIB SODIUM  
MESTINON, PYRIDOSTIGMINE BROMIDE  
TASMAR, TOLCAPONE  
TIMOPTIC-XE, TIMOLOL MALEATE

**VALEANT PHARMS NORTH**

- \* VALEANT PHARMACEUTICALS NORTH AMERICA LLC  
APLENZIN, BUPROPION HYDROBROMIDE  
CARAC, FLUOROURACIL  
DERMATOP, PREDNICARBATE  
DIASTAT ACUDIAL, DIAZEPAM  
DIASTAT, DIAZEPAM  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
FENOFLIBRATE, FENOFLIBRATE  
IPRIVASK, DESIRUDIN RECOMBINANT  
ISORDIL, ISOSORBIDE DINITRATE  
KLARON, SULFACETAMIDE SODIUM  
MINITRAN, NITROGLYCERIN  
NIFEDIPIINE, NIFEDIPINE  
NORITATE, METRONIDAZOLE  
PEPCID, FAMOTIDINE  
RENOVA, TRETINOIN

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* V \*\*

- \* VALEANT PHARMACEUTICALS NORTH AMERICA LLC
  - RETIN-A, TRETINOIN
  - SECONAL SODIUM, SECOBAREBITAL SODIUM
  - TIAZAC, DILTIAZEM HYDROCHLORIDE
  - VASOTEC, ENALAPRIL MALEATE
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - XENAZINE, TETRABENAZINE

**VALIDUS PHARMS**

- \* VALIDUS PHARMACEUTICALS LLC
  - BUMEX, BUMETANIDE
  - EQUETRO, CARBAMAZEPINE
  - NIFEDIPINE, NIFEDIPINE
  - ROCALTROL, CALCITRIOL

**VALIDUS PHARMS INC**

- \* VALIDUS PHARMACEUTICALS INC
  - MARPLAN, ISOCARBOXAZID

**VANDA PHARMS INC**

- \* VANDA PHARMACEUTICALS INC
  - FANAPT, ILOPERIDONE
  - HETLIOZ, TASIMELTEON

**VELOXIS PHARMS INC**

- \* VELOXIS PHARMACEUTICALS INC
  - ENVARSUS XR, TACROLIMUS

**VERNALIS R AND D LTD**

- \* VERNALIS R AND D LTD
  - MOXATAG, AMOXICILLIN
  - TUZISTRA XR, CHLORPHENIRAMINE POLISTIREX

**VEROSCIENCE**

- \* VEROSCIENCE LLC
  - CYCLOSET, BROMOCRIPTINE MESYLATE

**VERTEX PHARMS**

- \* VERTEX PHARMACEUTICALS INC
  - KALYDECO, IVACAFTOR

**VERTEX PHARMS INC**

- \* VERTEX PHARMACEUTICALS INC
  - KALYDECO, IVACAFTOR
  - ORKAMBI, IVACAFTOR

**VERTICAL PHARMS LLC**

- \* VERTICAL PHARMACEUTICALS LLC
  - DIVIGEL, ESTRADIOL

**VIB**

- \* VALEANT INTERNATIONAL BERMUDA
  - ZOVIRAX, ACYCLOVIR

**VICURON**

- \* VICURON PHARMACEUTICALS INC
  - ERAXIS, ANIDULAFUNGIN

**VIFOR FRESENIUS**

- \* VIFOR FRESENIUS MEDICAL CARE RENAL PHARMA FRANCE
  - VELPHORO, SUCROFERRIC OXYHYDROXIDE

**VIIV HLTHCARE**

- \* VIIV HEALTHCARE CO
  - COMBIVIR, LAMIVUDINE
  - EPIVIR, LAMIVUDINE
  - EPZICOM, ABACAVIR SULFATE
  - JULUCA, DOLUTEGRAVIR SODIUM
  - LEXIVA, FOSAMPRENAVIR CALCIUM
  - SCRIPTOR, DELAVIRDINE MESYLATE
  - RETROVIR, ZIDOVUDINE
  - SELZENTRY, MARAVIROC
  - TIVICAY, DOLUTEGRAVIR SODIUM
  - TRIUMEQ, ABACAVIR SULFATE
  - TRIZIVIR, ABACAVIR SULFATE
  - ZIAGEN, ABACAVIR SULFATE

**VINTAGE**

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* V \*\*

- \* VINTAGE PHARMACEUTICALS LLC
  - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - ACETIC ACID, ACETIC ACID, GLACIAL
  - ALBUTEROL SULFATE, ALBUTEROL SULFATE
  - ALPRAZOLAM, ALPRAZOLAM
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
  - CETIRIZINE HYDROCHLORIDE, CETIRIZINE HYDROCHLORIDE
  - FOLIC ACID, FOLIC ACID
  - HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
  - HYDROCORTISONE, HYDROCORTISONE
  - LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
  - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
  - NYSTATIN, NYSTATIN
  - PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE, PHENYLEPHRINE
  - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
  - PREDNISOLONE, PREDNISOLONE
  - PROMETH HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE, CODEINE
  - PROMETHAZINE DM, DEXTROMETHORPHAN HYDROBROMIDE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - PROMETHAZINE WITH CODEINE, CODEINE PHOSPHATE
  - RISPERIDONE, RISPERIDONE
  - SPIRONOLACTONE, SPIRONOLACTONE
  - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
  - TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
  - VALPROIC ACID, VALPROIC ACID
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

**VINTAGE PHARMS**

- \* VINTAGE PHARMACEUTICALS
  - ALPRAZOLAM, ALPRAZOLAM
  - BROMPHENIRAMINE MALEATE, PSEUDOEPHEDRINE HYDROCHLORIDE AND DEXTROMETHORPHAN
  - CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - CYCLAFEM 0.5/35, ETHINYL ESTRADIOL
  - DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
  - GILDAGIA, ETHINYL ESTRADIOL
  - GILDESS 24 FE, ETHINYL ESTRADIOL
  - GLYCOPYRROLATE, GLYCOPYRROLATE
  - GRISEOFULVIN, GRISEOFULVIN, MICROSIZE
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - KIMIDES, DESOGESTREL
  - LEVETIRACETAM, LEVETIRACETAM
  - METHSCOPOLAMINE BROMIDE, METHSCOPOLAMINE BROMIDE
  - METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
  - OXYCODONE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE

- \* VINTAGE PHARMACEUTICALS INC
  - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - ALLOPURINOL, ALLOPURINOL
  - AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
  - BACLOFEN, BACLOFEN
  - BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
  - BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - CARISOPRODOL, CARISOPRODOL
  - DEXAMETHASONE, DEXAMETHASONE
  - DIAZEPAM, DIAZEPAM
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
  - HYDROCORTISONE, HYDROCORTISONE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - IBUPROFEN, IBUPROFEN
  - IBUPROFEN, IBUPROFEN (OTC)
  - ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
  - LACTULOSE, LACTULOSE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* V \*\*

\* VINTAGE PHARMACEUTICALS INC  
 LEVETIRACETAM, LEVETIRACETAM  
 LORAZEPAM, LORAZEPAM  
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE  
 METHOCARBAMOL, METHOCARBAMOL  
 METHYLPREDNISOLONE, METHYLPREDNISOLONE  
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
 NYSTATIN, NYSTATIN  
 OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE  
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN  
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
 PERPHENAZINE, PERPHENAZINE  
 PREDNISONE, PREDNISONE  
 PRIMIDONE, PRIMIDONE  
 PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE  
 RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE  
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE  
 SULFASALAZINE, SULFASALAZINE  
 TORSEMIDE, TORSEMIDE

**VINTAGE PHARMS LLC**

\* VINTAGE PHARMACEUTICALS LLC  
 CYCLAFEM 1/35, ETHINYL ESTRADIOL  
 CYCLAFEM 7/7/7, ETHINYL ESTRADIOL  
 DUTASTERIDE, DUTASTERIDE  
 EMOQUETTE, DESOGESTREL  
 FELODIPINE, FELODIPINE  
 GILDESS 1.5/30, ETHINYL ESTRADIOL  
 GILDESS 1/20, ETHINYL ESTRADIOL  
 GILDESS FE 1.5/30, ETHINYL ESTRADIOL  
 GILDESS FE 1/20, ETHINYL ESTRADIOL  
 LETROZOLE, LETROZOLE  
 MEFENAMIC ACID, MEFENAMIC ACID  
 MONTELUKAST SODIUM, MONTELUKAST SODIUM  
 MORPHINE SULFATE, MORPHINE SULFATE  
 MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL  
 MYZILRA, ETHINYL ESTRADIOL  
 ORSYTHIA, ETHINYL ESTRADIOL  
 PERCOCET, ACETAMINOPHEN  
 PREVIFEM, ETHINYL ESTRADIOL  
 TRI-PREVIFEM, ETHINYL ESTRADIOL  
 TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

**VIRTUS PHARM**

\* VIRTUS PHARMACEUTICAL INC  
 ACARBOSE, ACARBOSE

**VIRTUS PHARMS**

\* VIRTUS PHARMACEUTICALS LLC  
 DAPSONE, DAPSONE  
 LEVETIRACETAM, LEVETIRACETAM  
 PROMETRIUM, PROGESTERONE  
 TRANEXAMIC ACID, TRANEXAMIC ACID

**VISTA PHARMS**

\* VISTA PHARMACEUTICALS INC  
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

**VISTAPHARM**

\* VISTAPHARM INC  
 ALBUTEROL SULFATE, ALBUTEROL SULFATE  
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN  
 LACTULOSE, LACTULOSE  
 METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE  
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE  
 MORPHINE SULFATE, MORPHINE SULFATE  
 NYSTATIN, NYSTATIN  
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE  
 PHENYTOIN, PHENYTOIN  
 PREDNISOLONE, PREDNISOLONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* V \*\*

- \* VISTAPHARM INC  
VALPROIC ACID, VALPROIC ACID

**VITRUVIAS THERAP**

- \* VITRUVIAS THERAPEUTICS LLC  
LIDOCAINE, LIDOCAINE

**VIVA HLTHCARE**

- \* VIVA HEALTHCARE FZ LLC  
GLIMEPIRIDE, GLIMEPIRIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
MEFENAMIC ACID, MEFENAMIC ACID  
SIMVASTATIN, SIMVASTATIN  
TRANEXAMIC ACID, TRANEXAMIC ACID

**VIVIMED GLOBAL**

- \* VIVIMED GLOBAL GENERICS PTE LTD  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
METRONIDAZOLE, METRONIDAZOLE

**VIVUS**

- \* VIVUS INC  
QSYMIA, PHENTERMINE HYDROCHLORIDE

**VPNA**

- \* VALEANT PHARMACEUTICALS NORTH AMERICA  
DICLOFENAC SODIUM, DICLOFENAC SODIUM

**VYERA PHARMS LLC**

- \* VYERA PHARMACEUTICALS LLC  
DARAPRIM, PYRIMETHAMINE

\*\* W \*\*

**WA UNIV SCH MED**

- \* WASHINGTON UNIV SCHOOL MEDICINE  
AMMONIA N 13, AMMONIA N-13  
CHOLINE C-11, CHOLINE C-11

**WARNER CHILCOTT LLC**

- \* WARNER CHILCOTT CO LLC  
CHOLEDYL SA, OXTRIPHYLLINE

**WATSON LABS**

- \* WATSON LABORATORIES  
FOLIC ACID, FOLIC ACID  
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
- \* WATSON LABORATORIES INC  
ACARBOSE, ACARBOSE  
AFEDITAB CR, NIFEDIPINE  
ALBUTEROL SULFATE, ALBUTEROL SULFATE  
ALENDRONATE SODIUM, ALENDRONATE SODIUM  
ALLOPURINOL, ALLOPURINOL  
AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
AMOXAPINE, AMOXAPINE  
ATENOLOL AND CHLORTHALIDONE, ATENOLOL  
CAPTOPRIL, CAPTOPRIL  
CARISOPRODOL, CARISOPRODOL  
CHLORZOXAZONE, CHLORZOXAZONE  
CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE  
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
CLONAZEPAM, CLONAZEPAM  
COL-PROBENECID, COLCHICINE  
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL  
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE  
DROSPIRENONE AND ETHINYL ESTRADIOL, DROSPIRENONE  
ESTAZOLAM, ESTAZOLAM  
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE  
GLIPIZIDE, GLIPIZIDE  
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE  
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WATSON LABORATORIES INC
  - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
  - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
  - LAMOTRIGINE, LAMOTRIGINE
  - LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - LISINOPRIL, LISINOPRIL
  - LORAZEPAM, LORAZEPAM
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
  - MEPROBAMATE, MEPROBAMATE
  - METHOCARBAMOL, METHOCARBAMOL
  - METHYLDOPA, METHYLDOPA
  - METHYLPREDNISOLONE, METHYLPREDNISOLONE
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - METRONIDAZOLE, METRONIDAZOLE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MINOXIDIL, MINOXIDIL
  - MIRTAZAPINE, MIRTAZAPINE
  - NABUMETONE, NABUMETONE
  - NALOXONE HYDROCHLORIDE AND PENTAZOCINE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
  - NATEGLINIDE, NATEGLINIDE
  - NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
  - NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
  - NIZATIDINE, NIZATIDINE
  - NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
  - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
  - OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
  - PRAVASTATIN SODIUM, PRAVASTATIN SODIUM
  - PREDNISOLONE, PREDNISOLONE
  - PREDNISONE, PREDNISONE
  - PRIMIDONE, PRIMIDONE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
  - QUASENSE, ETHINYL ESTRADIOL
  - QUINIDINE SULFATE, QUINIDINE SULFATE
  - RAMIPRIL, RAMIPRIL
  - RIVASTIGMINE TARTRATE, RIVASTIGMINE TARTRATE
  - SULFASALAZINE, SULFASALAZINE
  - SULINDAC, SULINDAC
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TELMISARTAN, TELMISARTAN
  - TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
  - TOPIRAMATE, TOPIRAMATE
  - TRANDOLAPRIL, TRANDOLAPRIL
  - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
  - TRIMETHOPRIM, TRIMETHOPRIM
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
  - ZOVIA 1/50E-28, ETHINYL ESTRADIOL
- \* WATSON LABS INC
  - LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

**WATSON LABS INC**

- \* WATSON LABORATORIES INC
  - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
  - AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE, AMLODIPINE BESYLATE
  - AMMONIUM LACTATE, AMMONIUM LACTATE
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CELECOXIB, CELECOXIB
  - CIPROFLOXACIN HYDROCHLORIDE, CIPROFLOXACIN HYDROCHLORIDE
  - DICLOFENAC SODIUM, DICLOFENAC SODIUM
  - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
  - DORZOLAMIDE HYDROCHLORIDE, DORZOLAMIDE HYDROCHLORIDE
  - DROSPIRENONONE, ETHINYL ESTRADIOL AND LEVOMEFOLATE CALCIUM, DROSPIRENONONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WATSON LABORATORIES INC
  - EZETIMIBE AND SIMVASTATIN, EZETIMIBE
  - EZETIMIBE, EZETIMIBE
  - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
  - METRONIDAZOLE, METRONIDAZOLE
  - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
  - MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
  - MODAFINIL, MODAFINIL
  - MOXIFLOXACIN HYDROCHLORIDE, MOXIFLOXACIN HYDROCHLORIDE
  - NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
  - PERPHENAZINE, PERPHENAZINE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
  - PROPOFOL, PROPOFOL
  - RALOXIFENE HYDROCHLORIDE, RALOXIFENE HYDROCHLORIDE
  - RASAGILINE MESYLATE, RASAGILINE MESYLATE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - ROSUVASTATIN CALCIUM, ROSUVASTATIN CALCIUM
  - SILDENAFIL CITRATE, SILDENAFIL CITRATE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VALSARTAN, VALSARTAN

**WATSON LABS TEVA**

- \* WATSON LABORATORIES INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - AFEDITAB CR, NIFEDIPINE
  - ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE, ALBUTEROL SULFATE
  - BICALUTAMIDE, BICALUTAMIDE
  - EZETIMIBE AND ATORVASTATIN CALCIUM, ATORVASTATIN CALCIUM
  - GLIPIZIDE, GLIPIZIDE
  - IBANDRONATE SODIUM, IBANDRONATE SODIUM
  - ISRADIPINE, ISRADIPINE
  - LEVOFLOXACIN, LEVOFLOXACIN
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
  - NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
  - PROBENECID, PROBENECID
  - SIMVASTATIN, SIMVASTATIN
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - VALSARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

**WATSON PHARMS INC**

- \* WATSON PHARMACEUTICALS INC
  - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
  - TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE

**WATSON PHARMS TEVA**

- \* WATSON PHARMACEUTICALS INC AN INDIRECT WHOLLY OWNED SUB OF TEVA PHARMACEUTICALS USA INC
  - RIFAMPIN, RIFAMPIN

**WELLSTAT THERAP**

- \* WELLSTAT THERAPEUTICS CORP
  - VISTOGARD, URIDINE TRIACETATE
  - XURIDEN, URIDINE TRIACETATE

**WES PHARMA INC**

- \* WES PHARMA INC
  - HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE

**WEST WARD PHARM CORP**

- \* WEST WARD PHARMACEUTICAL CORP
  - PHENYLEPHRINE HYDROCHLORIDE, PHENYLEPHRINE HYDROCHLORIDE
  - ROCURONIUM BROMIDE, ROCURONIUM BROMIDE

**WEST-WARD PHARM CORP**

- \* WEST-WARD PHARMACEUTICAL CORP
  - CEFOTETAN, CEFOTETAN DISODIUM

**WEST-WARD PHARMS INT**

- \* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
  - ACARBOSE, ACARBOSE
  - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
  - ADENOSINE, ADENOSINE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
  - ALENDRONATE SODIUM, ALENDRONATE SODIUM
  - ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
  - ALOSETRON HYDROCHLORIDE, ALOSETRON HYDROCHLORIDE
  - ALPRAZOLAM, ALPRAZOLAM
  - ALPROSTADIL, ALPROSTADIL
  - AMIKACIN SULFATE, AMIKACIN SULFATE
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
  - AMRINONE LACTATE, INAMRINONE LACTATE
  - ANASTROZOLE, ANASTROZOLE
  - ATIVAN, LORAZEPAM
  - ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
  - ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
  - AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
  - AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE
  - BALSALAZIDE DISODIUM, BALSALAZIDE DISODIUM
  - BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
  - BUMETANIDE, BUMETANIDE
  - BUPRENORPHINE HYDROCHLORIDE AND NALOXONE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
  - BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
  - BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
  - CAF'CIT, CAFFEINE CITRATE
  - CALCITRIOL, CALCITRIOL
  - CALCIUM ACETATE, CALCIUM ACETATE
  - CAPECITABINE, CAPECITABINE
  - CARBOPLATIN, CARBOPLATIN
  - CEFOXITIN, CEFOXITIN SODIUM
  - CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
  - CEVIMELINE HYDROCHLORIDE, CEVIMELINE HYDROCHLORIDE
  - CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
  - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
  - CILOSTAZOL, CILOSTAZOL
  - CISPLATIN, CISPLATIN
  - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
  - CLADRIBINE, CLADRIBINE
  - CLARITHROMYCYIN, CLARITHROMYCYIN
  - CLINDAMYCYIN PHOSPHATE, CLINDAMYCYIN PHOSPHATE
  - CLOTTRIMAZOLE, CLOTTRIMAZOLE
  - CODEINE SULFATE, CODEINE SULFATE
  - CYANOCOBALAMIN, CYANOCOBALAMIN
  - CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
  - CYCLOSPORINE, CYCLOSPORINE
  - CYTARABINE, CYTARABINE
  - DACARBAZINE, DACARBAZINE
  - DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
  - DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
  - DESVENLAFAKINE SUCCINATE, DESVENLAFAKINE SUCCINATE
  - DEXAMETHASONE INTENSOL, DEXAMETHASONE
  - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
  - DEXAMETHASONE, DEXAMETHASONE
  - DEXMEDETOMIDINE HYDROCHLORIDE, DEXMEDETOMIDINE HYDROCHLORIDE
  - DEXRAZOXANE HYDROCHLORIDE, DEXRAZOXANE HYDROCHLORIDE
  - DIAZEPAM INTENSOL, DIAZEPAM
  - DIAZEPAM, DIAZEPAM
  - DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE FREE), DICYCLOMINE HYDROCHLORIDE
  - DIGOXIN, DIGOXIN
  - DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
  - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
  - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
  - DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
  - DIPYRIDAMOLE, DIPYRIDAMOLE
  - DISULFIRAM, DISULFIRAM
  - DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD  
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE  
DOPRAM, DOXAPRAM HYDROCHLORIDE  
DOXERCALCIFEROL, DOXERCALCIFEROL  
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE  
DOXYCYCLINE, DOXYCYCLINE HYCLATE  
DURAMORPH PF, MORPHINE SULFATE  
DUTASTERIDE, DUTASTERIDE  
EPIRUBICIN HYDROCHLORIDE, EPIRUBICIN HYDROCHLORIDE  
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE  
ESZOPICLONE, ESZOPICLONE  
ETHACRYNIC ACID, ETHACRYNIC ACID  
ETOMIDATE, ETOMIDATE  
ETOPOSIDE, ETOPOSIDE  
EXEMESTANE, EXEMESTANE  
FAMCICLOVIR, FAMCICLOVIR  
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE  
FAMOTIDINE, FAMOTIDINE  
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE  
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE  
FLECAINIDE ACETATE, FLECAINIDE ACETATE  
FLOXURIDINE, FLOXURIDINE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE  
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE  
FLUCONAZOLE, FLUCONAZOLE  
FLUCYTOSINE, FLUCYTOSINE  
FLUMAZENIL, FLUMAZENIL  
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE  
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE (OTC)  
FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM  
FUROSEMIDE, FUROSEMIDE  
GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE  
GRANisetron HYDROCHLORIDE, GRANisetron HYDROCHLORIDE  
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE  
HALOPERIDOL, HALOPERIDOL LACTATE  
HEPARIN SODIUM, HEPARIN SODIUM  
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE  
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE  
IFOSFAMIDE, IFOSFAMIDE  
IMIPRAMINE PAMOATE, IMIPRAMINE PAMOATE  
INDOMETHACIN SODIUM, INDOMETHACIN SODIUM  
INFUMORPH, MORPHINE SULFATE  
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE  
IRBESARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
IRBESARTAN, IRBESARTAN  
IRINOTECAN HYDROCHLORIDE, IRINOTECAN HYDROCHLORIDE  
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE  
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE  
LACTULOSE, LACTULOSE  
LETROZOLE, LETROZOLE  
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM  
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM  
LEVOCARNITINE, LEVOCARNITINE  
LEVOLEUCOVORIN CALCIUM, LEVOLEUCOVORIN CALCIUM  
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE  
LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE  
LINEZOLID, LINEZOLID  
LITHIUM CARBONATE, LITHIUM CARBONATE  
LITHIUM CITRATE, LITHIUM CITRATE  
LORAZEPAM INTENSOL, LORAZEPAM  
LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE  
LOSARTAN POTASSIUM, LOSARTAN POTASSIUM  
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE  
MEGESTROL ACETATE, MEGESTROL ACETATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
  - MELPHALAN HYDROCHLORIDE, MELPHALAN HYDROCHLORIDE
  - MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
  - MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
  - MERCAPTOPURINE, MERCAPTOPURINE
  - MESNA, MESNA
  - METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
  - METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
  - METHAMPHETAMINE HYDROCHLORIDE, METHAMPHETAMINE HYDROCHLORIDE
  - METHOTREXATE SODIUM PRESERVATIVE FREE, METHOTREXATE SODIUM
  - METHOTREXATE SODIUM, METHOTREXATE SODIUM
  - METOPROLOL TARTRATE, METOPROLOL TARTRATE
  - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
  - MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
  - MILRINONE LACTATE, MILRINONE LACTATE
  - MITOMYCIN, MITOMYCIN
  - MITOXANTRONE HYDROCHLORIDE, MITOXANTRONE HYDROCHLORIDE
  - MONTELUKAST SODIUM, MONTELUKAST SODIUM
  - MORPHINE SULFATE, MORPHINE SULFATE
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL
  - NALOXONE, NALOXONE HYDROCHLORIDE
  - NAPROXEN, NAPROXEN
  - NARatriptan, NARatriptan HYDROCHLORIDE
  - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
  - NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
  - OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
  - OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
  - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
  - OXCARBAZEPINE, OXCARBAZEPINE
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - OXYMORPHONE HYDROCHLORIDE, OXYMORPHONE HYDROCHLORIDE
  - OXYTOCIN, OXYTOCIN
  - PACLITAXEL, PACLITAXEL
  - PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
  - PENTOSTATIN, PENTOSTATIN
  - PERINDOPRIL ERBUMINE, PERINDOPRIL ERBUMINE
  - PHENOXYBENZAMINE HYDROCHLORIDE, PHENOXYBENZAMINE HYDROCHLORIDE
  - PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
  - PHENYTOIN SODIUM, PHENYTOIN SODIUM
  - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - PREDNISONE INTENSOL, PREDNISONE
  - PREDNISONE, PREDNISONE
  - PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
  - PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
  - PROTRIPTYLINE HYDROCHLORIDE, PROTRIPTYLINE HYDROCHLORIDE
  - QUETIAPINE FUMARATE, QUETIAPINE FUMARATE
  - RAMIPRIL, RAMIPRIL
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RIFAMPIN, RIFAMPIN
  - RISPERIDONE, RISPERIDONE
  - RITONAVIR, RITONAVIR
  - ROBAXIN, METHOCARBAMOL
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - ROXICET, ACETAMINOPHEN
  - RUFINAMIDE, RUFINAMIDE
  - SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
  - SODIUM CHLORIDE 0.9%, SODIUM CHLORIDE
  - SODIUM FERRIC GLUCONATE COMPLEX IN SUCROSE, SODIUM FERRIC GLUCONATE COMPLEX
  - SODIUM OXYBATE, SODIUM OXYBATE
  - SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
  - STERILE WATER FOR INJECTION, STERILE WATER FOR INJECTION

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WEST-WARD PHARMACEUTICALS INTERNATIONAL LTD
  - SUFENTANIL CITRATE, SUFENTANIL CITRATE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
  - THIOTEP A, THIOTEP A
  - TINIDAZOLE, TINIDAZOLE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - TORSEMIDE, TORSEMIDE
  - TRIAZOLAM, TRIAZOLAM
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VECURONIUM BROMIDE, VECURONIUM BROMIDE
  - VINBLASTINE SULFATE, VINBLASTINE SULFATE
  - VINORELBINE TARTRATE, VINORELBINE TARTRATE
  - ZALEPLON, ZALEPLON
  - ZIDOVUDINE, ZIDOVUDINE

**WI MEDCL CYCLOTRON**

- \* WISCONSIN MEDICAL CYCLOTRON LLC
  - AMMONIA N 13, AMMONIA N-13
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**WILSHIRE PHARMS INC**

- \* WILSHIRE PHARMACEUTICALS INC
  - CARISOPRODOL, CARISOPRODOL
  - PERPHENAZINE, PERPHENAZINE
  - TIAGABINE HYDROCHLORIDE, TIAGABINE HYDROCHLORIDE

**WINDLAS HLTHCARE**

- \* WINDLAS HEALTHCARE PVT LTD
  - AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE

**WOCKHARDT**

- \* WOCKHARDT EU OPERATIONS (SWISS) AG
  - BROMFED-DM, BROMPHENIRAMINE MALEATE
  - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
  - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
  - LINDANE, LINDANE
  - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
- \* WOCKHARDT LTD
  - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
  - AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
  - AZITHROMYCIN, AZITHROMYCIN
  - BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
  - BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
  - CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
  - CEFOTAXIME, CEFOTAXIME SODIUM
  - CEFPROZIL, CEFPROZIL
  - CEFTAZIDIME, CEFTAZIDIME
  - CEFTRIAXONE, CEFTRIAXONE SODIUM
  - CEFUROXIME AXETIL, CEFUROXIME AXETIL
  - CETIRIZINE HYDROCHLORIDE ALLERGY, CETIRIZINE HYDROCHLORIDE (OTC)
  - CLARITHROMYCIN, CLARITHROMYCIN
  - DIVALPROEX SODIUM, DIVALPROEX SODIUM
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - FAMOTIDINE, FAMOTIDINE (OTC)
  - FOSPHENYTOIN SODIUM, FOSPHENYTOIN SODIUM
  - FUROSEMIDE, FUROSEMIDE
  - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LISINOPRIL, LISINOPRIL
  - METOPROLOL SUCCINATE, METOPROLOL SUCCINATE
  - NIACIN, NIACIN
  - NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
  - ONDANSETRON HYDROCHLORIDE, ONDANSETRON HYDROCHLORIDE
  - PANTOPRAZOLE SODIUM, PANTOPRAZOLE SODIUM
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE (OTC)
  - RISPERIDONE, RISPERIDONE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WOCKHARDT LTD
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE
  - SUMATRIPTAN SUCCINATE, SUMATRIPTAN SUCCINATE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TIMOLOL MALEATE, TIMOLOL MALEATE
  - VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE
  - ZONISAMIDE, ZONISAMIDE

**WOCKHARDT BIO AG**

- \* WOCKHARDT BIO AG
  - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
  - ACETIC ACID, ACETIC ACID, GLACIAL
  - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
  - AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
  - AMOXICILLIN, AMOXICILLIN
  - CARBAMAZEPINE, CARBAMAZEPINE
  - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
  - CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
  - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
  - CROMOLYN SODIUM, CROMOLYN SODIUM
  - CYCLOSPORINE, CYCLOSPORINE
  - DEXAMETHASONE, DEXAMETHASONE
  - DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
  - DOXE PIN HYDROCHLORIDE, DOXE PIN HYDROCHLORIDE
  - ERYTHROMYCIN, ERYTHROMYCIN
  - FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
  - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
  - FUROSEMIDE, FUROSEMIDE
  - GENERLAC, LACTULOSE
  - HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
  - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
  - LACTULOSE, LACTULOSE
  - LEVETIRACETAM, LEVETIRACETAM
  - LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
  - LINDANE, LINDANE
  - LITHIUM CITRATE, LITHIUM CITRATE
  - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
  - LORATADINE, LORATADINE (OTC)
  - MEGESTROL ACETATE, MEGESTROL ACETATE
  - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
  - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
  - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
  - NYSTATIN, NYSTATIN
  - OXACILLIN SODIUM, OXACILLIN SODIUM
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
  - PIPERACILLIN AND TAZOBACTAM, PIPERACILLIN SODIUM
  - PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
  - PREDNISOLONE, PREDNISOLONE
  - PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
  - PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
  - PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
  - SELENIUM SULFIDE, SELENIUM SULFIDE
  - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
  - VALPROIC ACID, VALPROIC ACID

**WOCKHARDT EU OPERATN**

- \* WOCKHARDT EU OPERATIONS SWISS AG
  - PHENYTOIN, PHENYTOIN

**WOCKHARDT LTD**

- \* WOCKHARDT LTD
  - BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
  - CAPTOPRIL, CAPTOPRIL
  - CARBIDOPA, LEVODOPA AND ENTACAPONE, CARBIDOPA

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* W \*\*

- \* WOCKHARDT LTD
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CHILDREN'S FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE
  - ENALAPRIL MALEATE, ENALAPRIL MALEATE
  - ENTACAPONE, ENTACAPONE
  - FAMOTIDINE, FAMOTIDINE
  - FEXOFENADINE HYDROCHLORIDE ALLERGY, FEXOFENADINE HYDROCHLORIDE (OTC)
  - FEXOFENADINE HYDROCHLORIDE HIVES, FEXOFENADINE HYDROCHLORIDE (OTC)
  - LAMOTRIGINE, LAMOTRIGINE
  - LANSOPRAZOLE, LANSOPRAZOLE (OTC)
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE

**WOCKHARDT USA**

- \* WOCKHARDT USA INC
  - GRANISETRON HYDROCHLORIDE, GRANISETRON HYDROCHLORIDE
- \* WOCKHARDT USA LLC
  - LANSOPRAZOLE, LANSOPRAZOLE

**WRASER PHARMS**

- \* WRASER PHARMACEUTICALS LLC
  - CETRAXAL, CIPROFLOXACIN HYDROCHLORIDE

**WRASER PHARMS LLC**

- \* WRASER PHARMACEUTICALS LLC
  - TREZIX, ACETAMINOPHEN

**WUSM CYCLOTRON**

- \* WASHINGTON UNIV SCH MEDICINE CYCLOTRON FACILITY
  - FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

**WYETH PHARMS INC**

- \* WYETH PHARMACEUTICALS INC
  - EFFEXOR XR, VENLAFAKINE HYDROCHLORIDE
  - PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE
  - PREMARIN, ESTROGENS, CONJUGATED
  - PREMPHASE 14/14, ESTROGENS, CONJUGATED
  - PREMPRO, ESTROGENS, CONJUGATED
  - PRISTIQ, DESVENLAFAKINE SUCCINATE
  - PROTONIX IV, PANTOPRAZOLE SODIUM
  - PROTONIX, PANTOPRAZOLE SODIUM
  - TRECATOR, ETHIONAMIDE
  - ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
  - ZOSYN, PIPERACILLIN SODIUM

**WYETH PHARMS PFIZER**

- \* WYETH PHARMACEUTICALS INC WHOLLY OWNED SUB PFIZER INC
  - DUAVEE, BAZEDOXIFENE ACETATE

\*\* X \*\*

**X GEN PHARMS**

- \* X GEN PHARMACEUTICALS INC
  - ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
  - AMPHOTERICIN B, AMPHOTERICIN B
  - BACIIM, BACITRACIN
  - COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
  - LEVETIRACETAM, LEVETIRACETAM
  - LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
  - NEOMYCIN AND POLYMYXIN B SULFATE, NEOMYCIN SULFATE
  - NEOMYCIN SULFATE, NEOMYCIN SULFATE
  - NYSTATIN, NYSTATIN
  - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

**X-GEN PHARMS**

- \* X-GEN PHARMACEUTICALS INC
  - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* X \*\*

**X-GEN PHARMS INC**

- \* X-GEN PHARMACEUTICALS INC
  - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
  - LINCOMYCIN, LINCOMYCIN HYDROCHLORIDE
  - TRANEXAMIC ACID, TRANEXAMIC ACID

**XELLIA PHARMS APS**

- \* XELLIA PHARMACEUTICALS APS
  - BACITRACIN, BACITRACIN
  - COLISTIMETHATE SODIUM, COLISTIMETHATE SODIUM
  - DAPTOXYCIN, DAPTOXYCIN
  - POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
  - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
  - VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
  - VORICONAZOLE, VORICONAZOLE

**XIAMEN LP PHARM CO**

- \* XIAMEN LP PHARMACEUTICAL CO LTD
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE

**XSPIRE PHARMA**

- \* XSPIRE PHARMA
  - NALFON, FENOPROFEN CALCIUM
- \* XSPIRE PHARMA LLC
  - DEXAMETHASONE, DEXAMETHASONE
  - FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

**XTTRIUM**

- \* XTTRIUM LABORATORIES INC
  - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
  - EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

\*\* Y \*\*

**YABAO PHARM**

- \* YABAO PHARMACEUTICAL CO LTD BEIJING
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE

**YAOPHARMA CO LTD**

- \* YAOPHARMA CO LTD
  - VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE

**YUNG SHIN PHARM**

- \* YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
  - CEFACLOR, CEFACLOR
  - CEPHALEXIN, CEPHALEXIN
  - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
  - MELOXICAM, MELOXICAM
  - ZOLPIDEM TARTRATE, ZOLPIDEM TARTRATE

\*\* Z \*\*

**ZAMBON SPA**

- \* ZAMBON SPA
  - DORZOLAMIDE HYDROCHLORIDE AND TIMOLOL MALEATE, DORZOLAMIDE HYDROCHLORIDE
  - OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
- \* ZAMBON SPA ITALY
  - MONUROL, FOSFOMYCIN TROMETHAMINE

**ZEVACOR PHARMA INC**

- \* ZEVACOR PHARMA INC
  - SODIUM FLUORIDE F-18, SODIUM FLUORIDE F-18

**ZHEJIANG HISUN PHARM**

- \* ZHEJIANG HISUN PHARMACEUTICAL CO LTD
  - DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE
  - MYCOPHENOLATE MOFETIL, MYCOPHENOLATE MOFETIL

**ZO SKIN HEALTH**

- \* ZO SKIN HEALTH
  - TRETINOIN, TRETINOIN

**ZYDUS HLTHCARE**

- \* ZYDUS HEALTHCARE USA LLC
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - LANSOPRAZOLE, LANSOPRAZOLE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* Z \*\*

\* ZYDUS HEALTHCARE USA LLC  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

**ZYDUS PHARMS USA**

\* ZYDUS PHARMACEUTICALS USA INC  
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE  
ATENOLOL, ATENOLOL  
AZATHIOPRINE, AZATHIOPRINE  
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE  
BENZONATATE, BENZONATATE  
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE  
HALOPERIDOL, HALOPERIDOL  
LAMOTRIGINE, LAMOTRIGINE  
MELOXICAM, MELOXICAM  
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE  
NAPROXEN, NAPROXEN  
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE  
PRAVASTATIN SODIUM, PRAVASTATIN SODIUM  
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE  
RAMIPRIL, RAMIPRIL  
RIBAVIRIN, RIBAVIRIN  
RISPERIDONE, RISPERIDONE  
SERTRALINE HYDROCHLORIDE, SERTRALINE HYDROCHLORIDE  
SIMVASTATIN, SIMVASTATIN  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE  
WARFARIN SODIUM, WARFARIN SODIUM  
ZONISAMIDE, ZONISAMIDE

**ZYDUS PHARMS USA INC**

\* ZYDUS PHARMACEUTICALS USA INC  
ACAMPROSATE CALCIUM, ACAMPROSATE CALCIUM  
ACETAZOLAMIDE, ACETAZOLAMIDE  
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM  
ACYCLOVIR, ACYCLOVIR  
ALLOPURINOL, ALLOPURINOL  
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE  
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE  
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE  
AMLODIPINE AND OLMESARTAN MEDOXOMIL, AMLODIPINE BESYLATE  
ANASTROZOLE, ANASTROZOLE  
ASPIRIN AND DIPYRIDAMOLE, ASPIRIN  
AZELASTINE HYDROCHLORIDE, AZELASTINE HYDROCHLORIDE  
BICALUTAMIDE, BICALUTAMIDE  
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE  
BUDESONIDE, BUDESONIDE  
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE  
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE  
CANDESARTAN CILEXETIL AND HYDROCHLOROTHIAZIDE, CANDESARTAN CILEXETIL  
CANDESARTAN CILEXETIL, CANDESARTAN CILEXETIL  
CARVEDILOL, CARVEDILOL  
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE  
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE  
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE  
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE  
CLOPIDOGREL BISULFATE, CLOPIDOGREL BISULFATE  
CLOZAPINE, CLOZAPINE  
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE  
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE  
DESOXIMETASONE, DESOXIMETASONE  
DESVENLAFAXINE SUCCINATE, DESVENLAFAXINE SUCCINATE  
DIFLUNISAL, DIFLUNISAL  
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE  
DIPYRIDAMOLE, DIPYRIDAMOLE  
DIVALPROEX SODIUM, DIVALPROEX SODIUM  
DONEPEZIL HYDROCHLORIDE, DONEPEZIL HYDROCHLORIDE  
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE  
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* Z \*\*

- \* ZYDUS PHARMACEUTICALS USA INC
  - DOXYCYCLINE, DOXYCYCLINE
  - DOXYCYCLINE, DOXYCYCLINE HYCLATE
  - DULOXETINE HYDROCHLORIDE, DULOXETINE HYDROCHLORIDE
  - DUTASTERIDE, DUTASTERIDE
  - ELETRIPTAN HYDROBROMIDE, ELETRIPTAN HYDROBROMIDE
  - ENTECAVIR, ENTECAVIR
  - ESCITALOPRAM OXALATE, ESCITALOPRAM OXALATE
  - ETHACRYNATE SODIUM, ETHACRYNATE SODIUM
  - ETODOLAC, ETODOLAC
  - ETOMIDATE, ETOMIDATE
  - EZETIMIBE, EZETIMIBE
  - FELBAMATE, FELBAMATE
  - FESOTERODINE FUMARATE, FESOTERODINE FUMARATE
  - FINASTERIDE, FINASTERIDE
  - FLUCONAZOLE, FLUCONAZOLE
  - GABAPENTIN, GABAPENTIN
  - GALANTAMINE HYDROBROMIDE, GALANTAMINE HYDROBROMIDE
  - GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
  - GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
  - GLYBURIDE, GLYBURIDE
  - HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
  - INDOMETHACIN, INDOMETHACIN
  - IRBESARTAN, IRBESARTAN
  - ITRACONAZOLE, ITRACONAZOLE
  - LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
  - LAMOTRIGINE, LAMOTRIGINE
  - LEVETIRACETAM, LEVETIRACETAM
  - LEVOFLOXACIN, LEVOFLOXACIN
  - LINEZOLID, LINEZOLID
  - LOSARTAN POTASSIUM AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - LOSARTAN POTASSIUM, LOSARTAN POTASSIUM
  - MEMANTINE HYDROCHLORIDE, MEMANTINE HYDROCHLORIDE
  - MESALAMINE, MESALAMINE
  - METHOTREXATE SODIUM, METHOTREXATE SODIUM
  - METRONIDAZOLE, METRONIDAZOLE
  - MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
  - MIRTAZAPINE, MIRTAZAPINE
  - MODAFINIL, MODAFINIL
  - MYCOPHENOLATE MOFETIL HYDROCHLORIDE, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
  - NADOLOL, NADOLOL
  - NATEGLINIDE, NATEGLINIDE
  - NIFEDIPINE, NIFEDIPINE
  - NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
  - OLMESARTAN MEDOXOMIL, OLMESARTAN MEDOXOMIL
  - OMEPRAZOLE, OMEPRAZOLE
  - OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
  - PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
  - PINDOLOL, PINDOLOL
  - PIOGLITAZONE HYDROCHLORIDE, PIOGLITAZONE HYDROCHLORIDE
  - POTASSIUM CITRATE, POTASSIUM CITRATE
  - PRAMIPEXOLE DIHYDROCHLORIDE, PRAMIPEXOLE DIHYDROCHLORIDE
  - PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
  - PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
  - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
  - RISPERIDONE, RISPERIDONE
  - ROPINIROLE HYDROCHLORIDE, ROPINIROLE HYDROCHLORIDE
  - SIROLIMUS, SIROLIMUS
  - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
  - TAMSULOSIN HYDROCHLORIDE, TAMSULOSIN HYDROCHLORIDE
  - TELMISARTAN AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
  - TELMISARTAN, TELMISARTAN
  - TEMOZOLOMIDE, TEMOZOLOMIDE
  - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
  - TOPIRAMATE, TOPIRAMATE

**APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**

\*\* Z \*\*

\* ZYDUS PHARMACEUTICALS USA INC  
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN  
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE  
TRANEXAMIC ACID, TRANEXAMIC ACID  
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE  
VALACYCLOVIR HYDROCHLORIDE, VALACYCLOVIR HYDROCHLORIDE  
VENLAFAXINE HYDROCHLORIDE, VENLAFAXINE HYDROCHLORIDE  
VORICONAZOLE, VORICONAZOLE  
ZIPRASIDONE HYDROCHLORIDE, ZIPRASIDONE HYDROCHLORIDE  
ZOLMITRIPTAN, ZOLMITRIPTAN  
ZYPITAMAG, PITAVASTATIN MAGNESIUM

**ZYDUS WORLDWIDE**

\* ZYDUS WORLDWIDE DMCC  
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE

**APPENDIX C****UNIFORM TERMS****DOSAGE FORMS**

AEROSOL, FOAM	OINTMENT, AUGMENTED
AEROSOL, METERED	PASTE
CAPSULE	PATCH
CAPSULE, DELAYED REL PELLETS	PELLET
CAPSULE, DELAYED RELEASE	POWDER
CAPSULE, EXTENDED RELEASE	POWDER, EXTENDED RELEASE
CAPSULE, PELLET	POWDER, METERED
CLOTH	RING
CONCENTRATE	SHAMPOO
CREAM	SOLUTION
CREAM, AUGMENTED	SOLUTION FOR SLUSH
ELIXIR	SOLUTION, EXTENDED RELEASE
EMULSION	SOLUTION, GEL FORMING/DROPS
ENEMA	SOLUTION, METERED
FILM	SOLUTION/DROPS
FILM, EXTENDED RELEASE	SPONGE
FOR SOLUTION	SPRAY
FOR SUSPENSION	SPRAY, METERED
FOR SUSPENSION, DELAYED RELEASE	SUPPOSITORY
FOR SUSPENSION, EXTENDED RELEASE	SUSPENSION
GAS	SUSPENSION, EXTENDED RELEASE
GEL	SUSPENSION/DROPS
GEL, AUGMENTED	SWAB
GEL, METERED	SYRUP
GRANULE	SYSTEM
GRANULE, DELAYED RELEASE	SYSTEM, EXTENDED RELEASE
GUM, CHEWING	TABLET
IMPLANT	TABLET, CHEWABLE
INHALANT	TABLET, COATED PARTICLES
INJECTABLE	TABLET, DELAYED RELEASE
INJECTABLE, LIPID COMPLEX	TABLET, DELAYED RELEASE, ORALLY DISINTEGRATING
INJECTABLE, LIPOSOMAL	TABLET, EFFERVESCENT
INJECTION, EXTENDED RELEASE	TABLET, EXTENDED RELEASE
INSERT	TABLET, EXTENDED RELEASE, CHEWABLE
INSERT, EXTENDED RELEASE	TABLET, FOR SUSPENSION
INTRAUTERINE DEVICE	TABLET, ORALLY DISINTEGRATING
JELLY	TABLET, ORALLY DISINTEGRATING, DELAYED RELEASE
LIQUID	TABLET, ORALLY DISINTEGRATING, EXTENDED RELEASE
LOTION	TAPE
LOTION, AUGMENTED	TROCHE/LOZENGE
LOTION/SHAMPOO	
OIL	
OIL/DROPS	
ointment	

Note: Terms comprise currently marketed products

**APPENDIX C****UNIFORM TERMS*****ROUTES OF ADMINISTRATION***

BUCCAL	IRRIGATION
DENTAL	IV (INFUSION)
ENDOCERVICAL	N/A
ENDOTRACHEL	NASAL
FOR RX COMPOUNDING	OPHTHALMIC
IMPLANTATION	ORAL
INHALATION	ORAL-21
INJECTION	ORAL-28
INTRA-ANAL	OTIC
INTRACRANIAL	PERFUSION, CARDIAC
INTRAMUSCULAR	PERIODONTAL
INTRAOCULAR	RECTAL
INTRAPERITONEAL	SPINAL
INTRAPLEURAL	SUBCUTANEOUS
INTRATHECAL	SUBLINGUAL
INTRATRACHEAL	TOPICAL
INTRAUTERINE	TRANSDERMAL
INTRAVENOUS	TRANSMUCOSAL
INTRAVESICAL	URETHRAL
INTRAVITREAL	VAGINAL
IONTOPHORESIS	

Note: Terms comprise currently marketed products

**APPENDIX C****UNIFORM TERMS*****ABBREVIATIONS***

AMP	AMPULE
AMPICIL	AMPICILLIN
APPROX	APPROXIMATELY
BOT	BOTTLE
CI	CURIE
CSR	CAROTID SINUS REFLEX
CU	CLINICAL UNITS
DIPROP	DIPROPIONATE
ELECT	ELECTROLYTE
EQ	EQUIVALENT TO
ER	EXTENDED RELEASE
GM	GRAM
HBR	HYDROBROMIDE
HCL	HYDROCHLORIDE
HR	HOUR
IM	INTRAMUSCULAR
INH	INHALATION
IU	INTERNATIONAL UNITS
IV	INTRAVENOUS
KIU	KALLIKREIN INHIBITOR UNITS
MCG	MICROGRAM
mCi	MILLCURIE
MEQ	MILLIEQUIVALENT
MG	MILLIGRAM
ML	MILLILITER
N/A	NOT APPLICABLE
PPM	PARTS PER MILLION
REL	RELEASE
SQ CM	SQUARE CENTIMETER
U	UNITS
uCi	MICROCURIE
UMOLAR	MICROMOLAR
USP	UNITED STATES PHARMACOPEIA

## PATENT AND EXCLUSIVITY INFORMATION ADDENDUM

This *Addendum* identifies drugs that qualify under the Federal Food, Drug, and Cosmetic Act (FD&C Act) for periods of exclusivity and provides patent information concerning the listed drug products.

### **Exclusivity**

During relevant exclusivity periods, certain abbreviated new drug applications (ANDAs) and applications described in Section 505(b)(2) of the FD&C Act (505(b)(2) applications) may not be submitted or approved as described below. This *Addendum* identifies drugs that qualify under the Drug Price Competition and Patent Term Restoration Act of 1984 (Hatch-Waxman Amendments) for periods of exclusivity. This *Addendum* also identifies those drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the FD&C Act, those drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A of the FD&C Act, those drugs that have qualified for Generating Antibiotics Incentives Now (GAIN) exclusivity pursuant to Section 505E of the FD&C Act, and those drugs that have qualified for Competitive Generic Therapy (CGT) exclusivity pursuant to Section 505(j)(5)(B)(v) of the FD&C Act. This section is arranged in alphabetical order by active ingredient name followed by the trade name. Active ingredient headings for multiple ingredient fixed-combination drug products are arranged alphabetically.

For an explanation of the codes used in the *Addendum*, see the *Patent and Exclusivity Terms Section*. The exclusivity codes are general shorthand descriptions and do not necessarily identify, with specificity, the actual scope of exclusivity. Please note that beginning with the publication of the 38<sup>th</sup> edition of the Orange Book, individual descriptions of the protected use have been added to each Orphan Drug Exclusivity entry listed in the Orange Book. In previous editions of the Orange Book, Orphan Drug Exclusivity was not described with any specificity.

The exclusivities identified in the *Addendum* do not prevent the submission or approval of an application submitted pursuant to Section 505(b)(1) of the FD&C Act that would otherwise be blocked if it had been submitted pursuant to Section 505(b)(2) or 505(j), except in the case of Orphan Drug Exclusivity. Drugs that may qualify for periods of exclusivity include:

- (1) A new chemical entity, submitted in a new drug application under Section 505(b) of the FD&C Act and approved after September 24, 1984. A new chemical entity is an active ingredient that contains "no active ingredient (including any ester or salt of the active ingredient)" that has been approved by FDA in any other application submitted under Section 505(b) of the FD&C Act. No subsequent ANDA or 505(b)(2) application for a drug that contains the same active moiety may be submitted for a period of five years from the date of approval of the original application, except that such an application may be submitted after four years if it contains a certification that a patent claiming the drug is invalid or will not be infringed by the product for which approval is sought. See Sections 505(j)(5)(F)(ii) and 505(c)(3)(E)(ii) of the FD&C Act.
- (2) A new drug application approved after September 24, 1984, for a drug product containing "an active ingredient (including any ester or salt of

the active ingredient)" that has been approved in an earlier new drug application and that includes reports of new clinical investigations (other than bioavailability studies). Such investigations must have been conducted or sponsored by the applicant and must have been essential to approval of the application. If these requirements are met, a subsequent ANDA or a 505(b)(2) application may not be approved for the exclusivity-protected "conditions of approval of such drug" before the expiration of three years from the date of approval of the original application. If an NDA has exclusivity only for a new indication or use, this exclusivity generally does not preclude the approval of an ANDA or 505(b)(2) application for indications and uses not covered by the exclusivity, assuming the proposed drug product will be safe and effective as labeled. See Sections 505(j)(5)(F)(iii) and 505(c)(3)(E)(iii) of the FD&C Act.

- (3) A supplement to a new drug application for a drug containing a previously approved "active ingredient (including any ester or salt of the active ingredient)" approved after September 24, 1984, that contains reports of new clinical investigations (other than bioavailability studies) essential to the approval of the supplement and conducted or sponsored by the applicant. A subsequent ANDA or 505(b)(2) application may not be approved for an exclusivity-protected change approved in the supplement for three years from the date of approval of the supplement. See Sections 505(j)(5)(F)(iv) and 505(c)(3)(E)(iv) of the FD&C Act.

## **Patent Information**

The FD&C Act requires that patent information be filed with all newly submitted Section 505(b) drug applications. No NDA may be approved after September 24, 1984, without the submission of patent information to the Agency. Effective August 18, 2003, this information must be filed using Form FDA 3542a "Patent Information Submitted with the Filing of an NDA, Amendment or Supplement".

Effective August 18, 2003, upon approval of an application, patent information for purposes of listing in the Orange Book must be submitted to the Agency within 30 days of the date of approval on Form FDA 3542 "Patent Information Submitted Upon and After Approval of an NDA or Supplement."<sup>1</sup> In November 2017, the Agency began including in the [Orange Book](#) the patent submission date (i.e., the date on which the FDA receives patent information from the NDA holder) for each newly listed patent. Patent information on unapproved applications or on patents beyond the scope of the FD&C Act (i.e., process or manufacturing patents) will not be published. Form FDA 3542 will be the only form used for the purposes of this publication.

The patents that FDA regards as covered by the statutory provisions for submission of patent information are: patents that claim the active ingredient(s); drug product patents, which include formulation/composition patents; method-of-use patents that claim one or more approved methods of using the approved drug product; and certain other patents as detailed on

---

<sup>1</sup> Please note that the date of approval for an NDA for a drug for which FDA intends to recommend controls under the Controlled Substances Act is the later of the date on the approval letter for the NDA or the date of issuance of the interim final rule controlling the drug (see section Section 505(x)(1) and (2) of the FD&C Act).

Form FDA 3542.<sup>2</sup> This information, as provided by the sponsor on Form FDA 3542, will be published as described above. As of December 5, 2016, an NDA holder submitting information on a patent that claims both the drug substance and the drug product (and is eligible for listing on either basis) is required only to specify that the patent claims either the drug substance or the drug product.

A requirement for submission of patent information to FDA for certain old antibiotics became effective October 7, 2008 under Section 4(b)(1) of the QI Program Supplemental Funding Act (Public Law 110-379) (QI Act).

Upon approval of an NDA, patent numbers and expiration dates for patent information submitted to FDA on Form FDA 3542 will be published daily in the Orange Book. The Addendum lists patent and exclusivity information up to January of the Edition year. The monthly Cumulative Supplements to the annual edition list patent and exclusivity information changes since the Annual Edition Addendum. Since all parts of this publication are subject to changes, additions, or deletions, the Orange Book, updated daily, should be consulted for the most recent patent and exclusivity information.

---

<sup>2</sup> See 21 CFR 314.53(c)(2)(ii)(M), (N)(2) and (N)(3).

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ABACAVIR SULFATE - ABACAVIR SULFATE</u></b>						
A 201107 001					PC	Mar 14, 2018
<b><u>ABACAVIR SULFATE - ZIAGEN</u></b>						
N 020977 001	6294540	May 14, 2018	DS DP U-65			
<b><u>ABACAVIR SULFATE - ZIAGEN</u></b>						
N 020978 001	6294540	May 14, 2018	DS DP U-65			
	6641843	Feb 04, 2019	DP			
<b><u>ABACAVIR SULFATE; DOLUTEGRAVIR SODIUM; LAMIVUDINE - TRIUMEQ</u></b>						
N 205551 001	6294540	May 14, 2018	DS DP U-1572			
	6294540*PED	Nov 14, 2018				
	8129385	Oct 05, 2027	DS DP			
	9242986	Dec 08, 2029	DS DP			
<b><u>ABACAVIR SULFATE; LAMIVUDINE - EPZICOM</u></b>						
N 021652 001	6294540	May 14, 2018	DS DP U-257			
<b><u>ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE - TRIZIVIR</u></b>						
N 021205 001	6294540	May 14, 2018	DS DP U-65			
<b><u>ABALOPARATIDE - TYMLOS</u></b>						
N 208743 001	7803770	Mar 26, 2028	U-2009			
	8148333	Nov 08, 2027	DP			
	8748382	Oct 03, 2027	U-2009			
<b><u>ABEMACICLIB - VERZENIO</u></b>						
N 208716 001	7855211	Dec 15, 2029	DS DP U-2132			
	7855211	Dec 15, 2029	DS DP U-2135			
<b><u>ABEMACICLIB - VERZENIO</u></b>						
N 208716 002	7855211	Dec 15, 2029	DS DP U-2132			
	7855211	Dec 15, 2029	DS DP U-2135			
<b><u>ABEMACICLIB - VERZENIO</u></b>						
N 208716 003	7855211	Dec 15, 2029	DS DP U-2132			
	7855211	Dec 15, 2029	DS DP U-2135			
<b><u>ABEMACICLIB - VERZENIO</u></b>						
N 208716 004	7855211	Dec 15, 2029	DS DP U-2132			
	7855211	Dec 15, 2029	DS DP U-2135			
<b><u>ABIRATERONE ACETATE - ZYTIGA</u></b>						
N 202379 001	8822438	Aug 24, 2027	U-1579			
	8822438	Aug 24, 2027	U-1580			
<b><u>ABIRATERONE ACETATE - ZYTIGA</u></b>						
N 202379 002	8822438	Aug 24, 2027	U-1579			
	8822438	Aug 24, 2027	U-1580			
<b><u>ACALABRUTINIB - CALQUENCE</u></b>						
N 210259 001	9290504	Jul 11, 2032	DS DP			
	9758524	Jul 11, 2032	U-2145			
	9796721	Jul 01, 2036	DS DP U-2145			
<b><u>ACETAMINOPHEN - OFIRMEV</u></b>						
N 022450 001	6028222	Aug 05, 2017	DP			
	6028222*PED	Feb 05, 2018			M-196	Jan 27, 2020
	6992218	Jun 06, 2021	DP		PED	Jul 27, 2020
	6992218*PED	Dec 06, 2021				
	9399012	Sep 11, 2031	U-1882			
	9399012*PED	Mar 11, 2032				
	9610265	Nov 13, 2028	U-2000			
<b><u>ACETAMINOPHEN; ASPIRIN; CAFFEINE - EXCEDRIN (MIGRAINE)</u></b>						
N 020802 001	5972916	Jul 14, 2017	U-296			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE - XARTEMIS XR</u></b>						
N 204031 001	6488962	Jun 20, 2020	DP			
	7976870	Jun 01, 2027		U-1498		
	8372432	Mar 11, 2029	DP	U-1499		
	8377453	Nov 19, 2029	DP	U-1499		
	8394408	Mar 11, 2029	DP			
	8597681	Dec 21, 2030	DP			
	8658631	May 16, 2032	DP			
	8668929	Mar 11, 2029		U-1499		
	8741885	May 16, 2032	DP	U-1499		
	8980319	Dec 21, 2030	DP			
	8992975	May 16, 2032	DP			
	9050335	May 16, 2032	DP			
	9468636	May 16, 2032		U-1499		
<b><u>ACETYLCYSTEINE - ACETADOTE</u></b>						
N 021539 001	8148356	May 21, 2026	DP			
	8399445	Aug 24, 2025		U-1373		
	8653061	Aug 24, 2025	U-1373			
	8722738	Apr 06, 2032	U-1373			
	9327028	Jul 21, 2031	U-1839			
<b><u>ACETYLCYSTEINE - CETYLEV</u></b>						
N 207916 001	8747894	May 08, 2032	DP	U-1373		
	9427421	May 08, 2032	DP			
	9561204	May 08, 2032		U-1373		
<b><u>ACETYLCYSTEINE - CETYLEV</u></b>						
N 207916 002	8747894	May 08, 2032	DP	U-1373		
	9427421	May 08, 2032	DP			
	9561204	May 08, 2032		U-1373		
<b><u>ACLIDINIUM BROMIDE - TUDORZA PRESSAIR</u></b>						
N 202450 001	6681768	Aug 07, 2022	DP		NCE	
	7078412	Jul 16, 2020	DS	DP U-1263		Jul 23, 2017
	8051851	Apr 22, 2027	DP			
	9056100	Jul 07, 2020	DP	U-1263		
	9333195	Jul 07, 2020	DP	U-1263		
	RE46417	Sep 05, 2020	DS	DP U-1263		
<b><u>ACYCLOVIR - SITAVIG</u></b>						
N 203791 001	8592434	Jun 16, 2030	DP	U-1460		
	8747896	Jun 03, 2027	DP	U-1460		
	8791127	Mar 23, 2027	DP	U-1460		
<b><u>ACYCLOVIR; HYDROCORTISONE - XERESE</u></b>						
N 022436 001	6514980	Jul 24, 2018	DP	U-1006		
	6514980	Jul 24, 2018	DP	U-1484		
	7223387	Nov 13, 2022	DP	U-1006		
	7223387	Nov 13, 2022	DP	U-1484		
<b><u>ADAPALENE - DIFFERIN</u></b>						
N 020380 002					RTO	Jul 08, 2019
<b><u>ADAPALENE - DIFFERIN</u></b>						
N 021753 001	7579377	Feb 23, 2025		U-818		
	7737181	Aug 29, 2024	DP			
	7834060	Mar 12, 2023		U-1078		
	7838558	Mar 12, 2023	DP			
	7868044	Mar 12, 2023		U-1078		
	8703820	Mar 12, 2023	U-1078			
<b><u>ADAPALENE - DIFFERIN</u></b>						
N 022502 001	7998467	May 31, 2028	DP	U-1078		
	8435502	Sep 15, 2026	DP	U-1078		
	8709392	Sep 15, 2026	DP	U-1078		
<b><u>ADAPALENE; BENZOYL PEROXIDE - ADAPALENE AND BENZOYL PEROXIDE</u></b>						
A 203790 001					PC	Jan 23, 2018

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO</u></b>						
N 022320 001	7820186	Nov 23, 2025	DP			
	7964202	Sep 01, 2024	DP U-1078			
	8071644	Jul 18, 2027	DP U-1078			
	8080537	Jul 18, 2027	U-1078			
	8105618	Dec 23, 2022	U-1078			
	8129362	Jul 18, 2027	U-1078			
	8241649	Dec 23, 2022	DP			
	8445543	Jul 12, 2027	U-1078			
	8809305	Dec 23, 2022	U-1078			
	8936800	Dec 23, 2022	DP U-1078			
<b><u>ADAPALENE; BENZOYL PEROXIDE - EPIDUO FORTE</u></b>						
N 207917 001	8445543	Dec 23, 2022	U-1078		NP	Jul 15, 2018
	8703820	Mar 12, 2023	U-1078			
	8729127	Mar 12, 2023	U-1078			
	8785420	Dec 23, 2022	U-1078			
	8809305	Dec 23, 2022	U-1078			
	8936800	Dec 23, 2022	DP U-1078			
	9381179	Mar 12, 2023	U-1078			
	9387187	Mar 12, 2023	U-1078			
	9814690	Dec 23, 2022	DP U-1078			
<b><u>ADEFOVIR DIPIVOXIL - HEPSSERA</u></b>						
N 021449 001	6451340	Jul 23, 2018	DS DP U-470			
<b><u>AFATINIB DIMALEATE - GILOTRIE</u></b>						
N 201292 001	6251912	Jul 29, 2018	DS DP U-1067		NCE	Jul 12, 2018
	8426586	Oct 10, 2029	DS		ODE-115	Apr 15, 2023
	8545884	Dec 19, 2029	DP		ODE-50	Jul 12, 2020
	9539258	Nov 09, 2026	U-1950			
	RE43431	Jan 22, 2022	DS			
<b><u>AFATINIB DIMALEATE - GILOTRIE</u></b>						
N 201292 002	6251912	Jul 29, 2018	DS DP U-1067		NCE	Jul 12, 2018
	8426586	Oct 10, 2029	DS		ODE-115	Apr 15, 2023
	8545884	Dec 19, 2029	DP		ODE-50	Jul 12, 2020
	9539258	Nov 09, 2026	U-1950			
	RE43431	Jan 22, 2022	DS			
<b><u>AFATINIB DIMALEATE - GILOTRIE</u></b>						
N 201292 003	6251912	Jul 29, 2018	DS DP U-1067		I-730	Apr 15, 2019
	8426586	Oct 10, 2029	DS		NCE	Jul 12, 2018
	8545884	Dec 19, 2029	DP		ODE-115	Apr 15, 2023
	9539258	Nov 09, 2026	U-1950		ODE-50	Jul 12, 2020
	RE43431	Jan 22, 2022	DS			
<b><u>ALBUMIN HUMAN - OPTISON</u></b>						
N 020899 001	6723303	Apr 20, 2021	DP			
<b><u>ALBUTEROL SULFATE - ACCUNEB</u></b>						
N 020949 001	6702997	Dec 28, 2021	U-558			
<b><u>ALBUTEROL SULFATE - ACCUNEB</u></b>						
N 020949 002	6702997	Dec 28, 2021	U-558			
<b><u>ALBUTEROL SULFATE - VENTOLIN HFA</u></b>						
N 020983 001	6161724	Jan 16, 2018	DP			
	6170717	Dec 23, 2017	DP			
	6315173	Dec 23, 2017	DP			
	6431168	Jun 08, 2018	DP			
	6435372	Jan 16, 2018	DP			
	6510969	Dec 23, 2017	DP			
	6938796	Jan 16, 2018	DP			
	6966467	Dec 23, 2017	DP			
	6997349	Jan 16, 2018	DP			
	7107986	Jun 08, 2018	DP			
	7143908	Jan 16, 2018	DP			
	7350676	Aug 24, 2018	DP			
	7500444	Feb 26, 2026	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ALBUTEROL SULFATE - VENTOLIN HFA</u></b>						
N 020983 001	7500444*PED 7832351	Aug 26, 2026 Jun 19, 2023	DP			
<b><u>ALBUTEROL SULFATE - PROAIR HFA</u></b>						
N 021457 001	6446627 7105152 8132712 9463289 9808587	Dec 18, 2017 Sep 12, 2023 Sep 07, 2028 May 18, 2031 May 18, 2031	DP			
<b><u>ALBUTEROL SULFATE - PROAIR RESPICLICK</u></b>						
N 205636 001	6446627 6701917 6718972 6748947 6871646 7540282 8006690 8651103 8978966 9216260 9463288 9731087	Dec 18, 2017 Jun 23, 2021 Jun 23, 2021 Jun 23, 2021 Jun 23, 2021 May 06, 2023 Jun 23, 2021 Mar 26, 2028 Jan 13, 2032 Jun 28, 2031 May 19, 2025 May 18, 2031	DP		NP NPP	Mar 12, 2018 Apr 28, 2019
<b><u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - DUONEB</u></b>						
N 020950 001	6632842	Dec 28, 2021	U-532			
<b><u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE - COMBIVENT RESPIMAT</u></b>						
N 021747 001	6846413 6977042 6988496 7284474 7396341 7802568 7837235 7896264 7988001 8733341 9027967	Aug 28, 2018 Aug 28, 2018 Feb 23, 2020 Aug 26, 2024 Oct 10, 2026 Feb 26, 2019 Mar 13, 2028 May 26, 2025 Aug 04, 2021 Oct 16, 2030 Mar 31, 2027	DP			
<b><u>ALCAFTADINE - LASTACAFT</u></b>						
N 022134 001	8664215	Dec 23, 2027	U-1493			
<b><u>ALECTINIB HYDROCHLORIDE - ALECENSA</u></b>						
N 208434 001	9126931 9365514 9440922	May 29, 2031 Mar 04, 2032 Jun 09, 2030	DS DP DP		I-756 NCE ODE ODE-105	Nov 06, 2020 Dec 11, 2020 Nov 06, 2024 Dec 11, 2022
<b><u>ALENDRONATE SODIUM - FOSAMAX</u></b>						
N 021575 001	5994329 6015801 6225294	Jul 17, 2018 Jul 17, 2018 Jul 17, 2018		Y Y Y		
<b><u>ALENDRONATE SODIUM - BINOSTO</u></b>						
N 202344 001	7488496 7964212	Aug 11, 2023 Mar 06, 2023	DS DP DS DP			
<b><u>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u></b>						
N 021762 001	5994329	Jul 17, 2018	U-647	Y		
<b><u>ALFUZOSIN HYDROCHLORIDE - UROXATRAL</u></b>						
N 021287 001	6149940	Aug 22, 2017				
<b><u>ALISKIREN HEMIFUMARATE - TEKTURNA</u></b>						
N 021985 001	5559111 5559111*PED 8617595 8617595*PED	Jul 21, 2018 Jan 21, 2019 Feb 19, 2026 Aug 19, 2026	DS DP U-3 DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ALISKIREN HEMIFUMARATE - TEKTURNA</u></b>						
N 021985 002	5559111	Jul 21, 2018	DS DP U-3			
	5559111*PED	Jan 21, 2019				
	8617595	Feb 19, 2026	DP			
	8617595*PED	Aug 19, 2026				
<b><u>ALISKIREN HEMIFUMARATE - TEKTURNA</u></b>						
N 210709 001					NP	Nov 14, 2020
					PED	May 14, 2021
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u></b>						
N 022545 001	5559111	Jul 21, 2018	DS DP U-3			
	8613949	Dec 21, 2029	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u></b>						
N 022545 002	5559111	Jul 21, 2018	DS DP U-3			
	8613949	Dec 21, 2029	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u></b>						
N 022545 003	5559111	Jul 21, 2018	DS DP U-3			
	8613949	Dec 21, 2029	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE - TEKAMLO</u></b>						
N 022545 004	5559111	Jul 21, 2018	DS DP U-3			
	8613949	Dec 21, 2029	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u></b>						
N 200045 001	5559111	Jul 21, 2018	DS DP U-3			
	8183295	May 16, 2023	DP			
	8618174	Nov 15, 2021	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u></b>						
N 200045 002	5559111	Jul 21, 2018	DS DP U-3			
	8183295	May 16, 2023	DP			
	8618174	Nov 15, 2021	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u></b>						
N 200045 003	5559111	Jul 21, 2018	DS DP U-3			
	8183295	May 16, 2023	DP			
	8618174	Nov 15, 2021	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u></b>						
N 200045 004	5559111	Jul 21, 2018	DS DP U-3			
	8183295	May 16, 2023	DP			
	8618174	Nov 15, 2021	DP			
<b><u>ALISKIREN HEMIFUMARATE; AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE - AMTURNIDE</u></b>						
N 200045 005	5559111	Jul 21, 2018	DS DP U-3			
	8183295	May 16, 2023	DP			
	8618174	Nov 15, 2021	DP			
<b><u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKURNA HCT</u></b>						
N 022107 001	5559111	Jul 21, 2018	DS DP U-3			
	5559111*PED	Jan 21, 2019				
	8618172	Jul 13, 2028	DP			
	9023893	Mar 03, 2022	DP			
<b><u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKURNA HCT</u></b>						
N 022107 002	5559111	Jul 21, 2018	DS DP U-3			
	5559111*PED	Jan 21, 2019				
	8618172	Jul 13, 2028	DP			
	9023893	Mar 03, 2022	DP			
<b><u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKURNA HCT</u></b>						
N 022107 003	5559111	Jul 21, 2018	DS DP U-3			
	5559111*PED	Jan 21, 2019				
	8618172	Jul 13, 2028	DP			
	9023893	Mar 03, 2022	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ALISKIREN HEMIFUMARATE; HYDROCHLOROTHIAZIDE - TEKturna HCT</u></b>						
N 022107 004	5559111	Jul 21, 2018	DS DP U-3			
	5559111*PED	Jan 21, 2019				
	8618172	Jul 13, 2028	DP			
	9023893	Mar 03, 2022	DP			
<b><u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALturna</u></b>						
N 022217 001	5559111	Jul 21, 2018	DS DP U-3			
	8168616	Jul 03, 2026	DP			
<b><u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALturna</u></b>						
N 022217 002	5559111	Jul 21, 2018	DS DP U-3			
	8168616	Jul 03, 2026	DP			
<b><u>ALLOPURINOL; LESINURAD - DUZALLO</u></b>						
N 209203 001	8003681	Aug 25, 2025	DS			
	8084483	Aug 17, 2029		U-2104		
	8283369	Nov 26, 2028		U-2104		
	8357713	Nov 26, 2028		DP U-2104		
	8546436	Feb 29, 2032	DS			
	8546437	Apr 29, 2029		U-2104		
	9216179	Aug 01, 2031		U-2104		
<b><u>ALLOPURINOL; LESINURAD - DUZALLO</u></b>						
N 209203 002	8003681	Aug 25, 2025	DS			
	8084483	Aug 17, 2029		U-2104		
	8283369	Nov 26, 2028		U-2104		
	8357713	Nov 26, 2028		DP U-2104		
	8546436	Feb 29, 2032	DS			
	8546437	Apr 29, 2029		U-2104		
	9216179	Aug 01, 2031		U-2104		
<b><u>ALOGLIPTIN BENZOATE - NESINA</u></b>						
N 022271 001	6890898	Feb 02, 2019		U-1335	M-177	Apr 05, 2019
	7078381	Feb 02, 2019		U-1335	NCE	Jan 25, 2018
	7459428	Feb 02, 2019		U-1336		
	7807689	Jun 27, 2028	DS DP	U-1337		
	8173663	Dec 02, 2025		U-1338		
	8288539	Mar 15, 2025	DS			
	8697125	Jun 16, 2029		DP		
<b><u>ALOGLIPTIN BENZOATE - NESINA</u></b>						
N 022271 002	6890898	Feb 02, 2019		U-1335	M-177	Apr 05, 2019
	7078381	Feb 02, 2019		U-1335	NCE	Jan 25, 2018
	7459428	Feb 02, 2019		U-1336		
	7807689	Jun 27, 2028	DS DP	U-1337		
	8173663	Dec 02, 2025		U-1338		
	8288539	Mar 15, 2025	DS			
	8697125	Jun 16, 2029		DP		
<b><u>ALOGLIPTIN BENZOATE - NESINA</u></b>						
N 022271 003	6890898	Feb 02, 2019		U-1335	M-177	Apr 05, 2019
	7078381	Feb 02, 2019		U-1335	NCE	Jan 25, 2018
	7459428	Feb 02, 2019		U-1336		
	7807689	Jun 27, 2028	DS DP	U-1337		
	8173663	Dec 02, 2025		U-1338		
	8288539	Mar 15, 2025	DS			
	8697125	Jun 16, 2029		DP		
<b><u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u></b>						
N 203414 001	6890898	Feb 02, 2019		U-1335	M-177	Apr 05, 2019
	7078381	Feb 02, 2019		U-1335	NCE	Jan 25, 2018
	7459428	Feb 02, 2019		U-1336		
	7807689	Jun 27, 2028	DS DP	U-1337		
	8173663	Mar 15, 2025		U-1338		
	8288539	Jun 24, 2025	DS			
	8900638	May 24, 2029		DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ALOGLIPTIN BENZOATE; METFORMIN HYDROCHLORIDE - KAZANO</u></b>						
N 203414 002	6890898	Feb 02, 2019	U-1335		M-177	Apr 05, 2019
	7078381	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Jun 24, 2025	DS			
	8900638	May 24, 2029	DP			
<b><u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u></b>						
N 022426 001	6329404	Jun 19, 2021	DP U-1334		M-177	Apr 05, 2019
	6890898	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7078381	Feb 02, 2019	U-1335			
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<b><u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u></b>						
N 022426 002	6329404	Jun 19, 2021	DP U-1334		M-177	Apr 05, 2019
	6890898	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7078381	Feb 02, 2019	U-1335			
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<b><u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u></b>						
N 022426 003	6329404	Jun 19, 2021	DP U-1334		M-177	Apr 05, 2019
	6890898	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7078381	Feb 02, 2019	U-1335			
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<b><u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u></b>						
N 022426 004	6329404	Jun 19, 2021	DP U-1334		M-177	Apr 05, 2019
	6890898	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7078381	Feb 02, 2019	U-1335			
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<b><u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u></b>						
N 022426 005	6329404	Jun 19, 2021	DP U-1334		M-177	Apr 05, 2019
	6890898	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7078381	Feb 02, 2019	U-1335			
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			
<b><u>ALOGLIPTIN BENZOATE; PIOGLITAZONE HYDROCHLORIDE - OSENI</u></b>						
N 022426 006	6890898	Feb 02, 2019	U-1335		M-177	Apr 05, 2019
	7078381	Feb 02, 2019	U-1335		NCE	Jan 25, 2018
	7459428	Feb 02, 2019	U-1336			
	7807689	Jun 27, 2028	DS DP U-1337			
	8173663	Mar 15, 2025	U-1338			
	8288539	Mar 15, 2025	DS			
	8637079	Jun 04, 2029	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ALPRAZOLAM - NIRAVAM</u></b>						
N 021726 001	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>ALPRAZOLAM - NIRAVAM</u></b>						
N 021726 002	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>ALPRAZOLAM - NIRAVAM</u></b>						
N 021726 003	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>ALPRAZOLAM - NIRAVAM</u></b>						
N 021726 004	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>ALVIMOPAN - ENTEREG</u></b>						
N 021775 001	6469030	Nov 29, 2020	U-879			
	8112290	Jul 31, 2030	U-1443	Y		
	8645160	Jun 18, 2029	U-1485	Y		
	8946262	Feb 12, 2030	U-1655			
<b><u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u></b>						
N 208944 001	8389578	Jan 22, 2028	U-2105		NP	Aug 24, 2020
	8741343	Dec 02, 2030	U-2106		ODE-153	Aug 24, 2024
	8796337	Nov 23, 2025	U-2106			
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025	U-2106			
	8895616	Nov 23, 2025	U-2106			
	8895617	Nov 23, 2025	U-2106			
	8895618	Nov 23, 2025	DP			
<b><u>AMANTADINE HYDROCHLORIDE - GOCOVRI</u></b>						
N 208944 002	8389578	Jan 22, 2028	U-2105		NP	Aug 24, 2020
	8741343	Dec 02, 2030	U-2106		ODE	Aug 24, 2024
	8796337	Nov 23, 2025	U-2106			
	8889740	Nov 23, 2025	DP			
	8895614	Nov 23, 2025	DP			
	8895615	Nov 23, 2025	U-2106			
	8895616	Nov 23, 2025	U-2106			
	8895617	Nov 23, 2025	U-2106			
	8895618	Nov 23, 2025	DP			
<b><u>AMBRISENTAN - LETAIRIS</u></b>						
N 022081 001	8377933	Dec 11, 2027	U-1754		I-716	Oct 02, 2018
	9474752	Dec 11, 2027	U-1754			
	9549926	Oct 14, 2031	U-1965			
	RE42462	Jul 29, 2018	DS			
<b><u>AMBRISENTAN - LETAIRIS</u></b>						
N 022081 002	8377933	Dec 11, 2027	U-1754		I-716	Oct 02, 2018
	9474752	Dec 11, 2027	U-1754			
	9549926	Oct 14, 2031	U-1965			
	RE42462	Jul 29, 2018	DS			
<b><u>AMIFOSTINE - ETHYOL</u></b>						
N 020221 001	5994409	Dec 08, 2017	U-305			
<b><u>AMIFOSTINE - ETHYOL</u></b>						
N 020221 002	5994409	Dec 08, 2017	U-305			
<b><u>AMINOLEVULINIC ACID HYDROCHLORIDE - LEVULAN</u></b>						
N 020965 001	5954703	Oct 31, 2017	U-289			
	6709446	May 01, 2018	U-289			
	7723910	Jun 17, 2019	U-289			
	8216289	May 01, 2018	U-289			
	8758418	May 01, 2018	U-289			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - AMBLUZ</u> N 208081 001 6559183		Nov 12, 2019	DP U-804		NP	May 10, 2019
<u>AMINOLEVULINIC ACID HYDROCHLORIDE - GLEOLAN</u> N 208630 001					ODE-146	Jun 06, 2024
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u> N 022325 001 6869939 7635773		May 04, 2022 Mar 13, 2029	DP DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u> N 022325 002 6869939 7635773		May 04, 2022 Mar 13, 2029	DP DP			
<u>AMIODARONE HYDROCHLORIDE - NEXTERONE</u> N 022325 003 6869939 7635773		May 04, 2022 Mar 13, 2029	DP DP			
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u> N 022026 001 6828339		Nov 20, 2022	DS			
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u> N 022026 002 6828339		Nov 20, 2022	DS			
<u>AMLODIPINE BESYLATE - AMLODIPINE BESYLATE</u> N 022026 003 6828339		Nov 20, 2022	DS			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 001 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 002 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 003 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 004 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 005 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 006 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 007 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 008 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 009 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 010 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM - CADUET</u> N 021540 011 6455574		Aug 11, 2018	U-552			
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u> N 020364 002 6162802		Dec 19, 2017	U-367			
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u> N 020364 003 6162802		Dec 19, 2017	U-367			
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u> N 020364 004 6162802		Dec 19, 2017	U-367			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u></b>						
N 020364 005	6162802	Dec 19, 2017		U-367		
<b><u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u></b>						
N 020364 006	6162802	Dec 19, 2017	DS DP	U-185		
<b><u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u></b>						
N 020364 007	6162802	Dec 19, 2017	DS DP	U-185		
<b><u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u></b>						
N 022314 001	8101599	May 16, 2023	DP			
	8475839	May 16, 2023	DP			
	8475839*PED	Nov 16, 2023				
<b><u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u></b>						
N 022314 002	8101599	May 16, 2023	DP			
	8475839	May 16, 2023	DP			
	8475839*PED	Nov 16, 2023				
<b><u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u></b>						
N 022314 003	8101599	May 16, 2023	DP			
	8475839	May 16, 2023	DP			
	8475839*PED	Nov 16, 2023				
<b><u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u></b>						
N 022314 004	8101599	May 16, 2023	DP			
	8475839	May 16, 2023	DP			
	8475839*PED	Nov 16, 2023				
<b><u>AMLODIPINE BESYLATE; HYDROCHLOROTHIAZIDE; VALSARTAN - EXFORGE HCT</u></b>						
N 022314 005	8101599	May 16, 2023	DP			
	8475839	May 16, 2023	DP			
	8475839*PED	Nov 16, 2023				
<b><u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u></b>						
N 205003 001	6696481	Apr 15, 2023	DS DP U-3		NP	Jan 21, 2018
	7846961	Oct 05, 2029	DS DP U-3			
<b><u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u></b>						
N 205003 002	6696481	Apr 15, 2023	DS DP U-3		NP	Jan 21, 2018
	7846961	Oct 05, 2029	DS DP U-3			
<b><u>AMLODIPINE BESYLATE; PERINDOPRIL ARGININE - PRESTALIA</u></b>						
N 205003 003	6696481	Apr 15, 2023	DS DP U-3		NP	Jan 21, 2018
	7846961	Oct 05, 2029	DS DP U-3			
<b><u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u></b>						
N 021990 002	6395728	Jul 08, 2019	DP			
<b><u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u></b>						
N 021990 003	6395728	Jul 08, 2019	DP			
<b><u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u></b>						
N 021990 004	6395728	Jul 08, 2019	DP			
<b><u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u></b>						
N 021990 005	6395728	Jul 08, 2019	DP			
<b><u>AMOXICILLIN - MOXATAG</u></b>						
N 050813 001	6544555	Oct 13, 2020	DS DP U-897			
	6669948	Oct 13, 2020	DS DP U-897			
	6723341	Oct 13, 2020	DS DP U-897			
	8299052	May 07, 2027		U-1304		
	8357394	Dec 08, 2026	DP			
	8778924	Dec 08, 2026	DS DP	U-897		
<b><u>AMOXICILLIN; CLAVULANATE POTASSIUM - AUGMENTIN XR</u></b>						
N 050785 001	6746692	Apr 04, 2020	DP			
	6783773	Apr 04, 2020	DP			
	6878386	Apr 04, 2020		U-926		
	7217430	Apr 04, 2020	DP	U-926		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>AMOXICILLIN; CLAVULANATE POTASSIUM - AUGMENTIN XR</u></b>						
N 050785 001	7250176	Apr 04, 2020		U-926		
<b><u>AMPHETAMINE - ADZENYS ER</u></b>						
N 204325 001	8709491	Jun 28, 2032	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - ADZENYS XR-ODT</u></b>						
N 204326 001	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - ADZENYS XR-ODT</u></b>						
N 204326 002	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - ADZENYS XR-ODT</u></b>						
N 204326 003	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - ADZENYS XR-ODT</u></b>						
N 204326 004	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - ADZENYS XR-ODT</u></b>						
N 204326 005	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - ADZENYS XR-ODT</u></b>						
N 204326 006	8709491	Jun 28, 2032	DP			
	8840924	Apr 09, 2026	DP			
	9017731	Jun 28, 2032	DP			
	9265737	Jun 28, 2032	DP			
<b><u>AMPHETAMINE - DYANAVEL XR</u></b>						
N 208147 001	8062667	Mar 29, 2029	DP		NP	Oct 19, 2018
	8597684	Mar 15, 2027	DP			
	8747902	Mar 15, 2027	DP			
	8883217	Mar 15, 2027	DP			
	9675703	Mar 15, 2027	DP			
<b><u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 10</u></b>						
N 011522 007	6384020	Jul 06, 2020				
<b><u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 20</u></b>						
N 011522 008	6384020	Jul 06, 2020				
<b><u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 5</u></b>						
N 011522 009	6384020	Jul 06, 2020				
<b><u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 30</u></b>						
N 011522 010	6384020	Jul 06, 2020				
<b><u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 7.5</u></b>						
N 011522 011	6384020	Jul 06, 2020				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 12.5</u>						
N 011522 012	6384020	Jul 06, 2020				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL 15</u>						
N 011522 013	6384020	Jul 06, 2020				
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 10</u>						
N 021303 001	6322819 6605300 RE41148 RE42096	Oct 21, 2018 Oct 21, 2018 Oct 21, 2018 Oct 21, 2018		DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 20</u>						
N 021303 002	6322819 6605300 RE41148 RE42096	Oct 21, 2018 Oct 21, 2018 Oct 21, 2018 Oct 21, 2018		DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 30</u>						
N 021303 003	6322819 6605300 RE41148 RE42096	Oct 21, 2018 Oct 21, 2018 Oct 21, 2018 Oct 21, 2018		DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 25</u>						
N 021303 004	6322819 6605300 RE41148 RE42096	Oct 21, 2018 Oct 21, 2018 Oct 21, 2018 Oct 21, 2018		DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 5</u>						
N 021303 005	6322819 6605300 RE41148 RE42096	Oct 21, 2018 Oct 21, 2018 Oct 21, 2018 Oct 21, 2018		DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - ADDERALL XR 15</u>						
N 021303 006	6322819 6605300 RE41148 RE42096	Oct 21, 2018 Oct 21, 2018 Oct 21, 2018 Oct 21, 2018		DP		
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 001	6913768 8846100 9173857 RE41148 RE42096	May 24, 2023 Aug 24, 2029 May 12, 2026 Oct 21, 2018 Oct 21, 2018	DP U-2025 DP U-2025 DP DP		NP	Jun 20, 2020
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 002	6913768 8846100 9173857 RE41148 RE42096	May 24, 2023 Aug 24, 2029 May 12, 2026 Oct 21, 2018 Oct 21, 2018	DP U-2025 DP U-2025 DP DP		NP	Jun 20, 2020
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 003	6913768 8846100 9173857 RE41148	May 24, 2023 Aug 24, 2029 May 12, 2026 Oct 21, 2018	DP U-2025 DP U-2025 DP		NP	Jun 20, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 003	RE42096	Oct 21, 2018	DP			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE - MYDAYIS</u>						
N 022063 004	6913768	May 24, 2023	DP U-2025		NP	Jun 20, 2020
	8846100	Aug 24, 2029	DP			
	9173857	May 12, 2026	U-2025			
	RE41148	Oct 21, 2018	DP			
	RE42096	Oct 21, 2018	DP			
<u>AMPHOTERICIN B - ABELCET</u>						
N 050724 001	6406713	Jun 18, 2019	DS			
<u>AMPRENAVIR - AGENERASE</u>						
N 021007 001	6730679	Nov 11, 2017	DP			
<u>AMPRENAVIR - AGENERASE</u>						
N 021007 002	6730679	Nov 11, 2017	DP			
<u>ANIDULAFUNGIN - ERAXIS</u>						
N 021632 001	5965525	Feb 17, 2020	DS DP U-540			
	6960564	Apr 12, 2021	DP U-540			
	7709444	Apr 12, 2021	DP U-540			
<u>ANIDULAFUNGIN - ERAXIS</u>						
N 021632 002	5965525	Feb 17, 2020	DS DP U-540			
	6960564	Apr 12, 2021	DP U-540			
	7709444	Apr 12, 2021	DP U-540			
<u>APIXABAN - ELIQUIS</u>						
N 202155 001	6413980	Dec 22, 2019	DS DP U-1200		I-661	Aug 21, 2017
	6413980	Dec 22, 2019	DS DP U-1301		I-690	Aug 21, 2017
	6413980	Dec 22, 2019	DS DP U-1302		I-691	Aug 21, 2017
	6413980	Dec 22, 2019	DS DP U-1501		NCE	Dec 28, 2017
	6967208	Nov 21, 2026	DS DP U-1167			
	6967208	Nov 21, 2026	DS DP U-1200			
	6967208	Nov 21, 2026	DS DP U-1301			
	6967208	Nov 21, 2026	DS DP U-1302			
	6967208	Nov 21, 2026	DS DP U-1323			
	6967208	Nov 21, 2026	DS DP U-1501			
	6967208	Nov 21, 2026	DS DP U-1502			
	6967208	Nov 21, 2026	DS DP U-1729			
	6967208	Nov 21, 2026	DS DP U-1730			
	9326945	Feb 24, 2031	DP			
<u>APIXABAN - ELIQUIS</u>						
N 202155 002	6413980	Dec 22, 2019	DS DP U-1200		I-661	Aug 21, 2017
	6413980	Dec 22, 2019	DS DP U-1301		I-690	Aug 21, 2017
	6413980	Dec 22, 2019	DS DP U-1302		I-691	Aug 21, 2017
	6967208	Nov 21, 2026	DS DP U-1200		NCE	Dec 28, 2017
	6967208	Nov 21, 2026	DS DP U-1301			
	6967208	Nov 21, 2026	DS DP U-1302			
	6967208	Nov 21, 2026	DS DP U-1323			
	9326945	Feb 24, 2031	DP			
<u>APREMILAST - OTEZIA</u>						
N 205437 001	6020358	Oct 30, 2018	DS DP U-1504		I-694	Sep 23, 2017
	6962940	Mar 19, 2023	U-1504		NCE	Mar 21, 2019
	7208516	Mar 19, 2023	U-1505			
	7427638	Nov 17, 2024	DS DP			
	7659302	Mar 19, 2023	U-1505			
	7659302	Mar 19, 2023	U-1595			
	7893101	Dec 09, 2023	DS DP			
	8455536	Mar 19, 2023	U-1505			
	8455536	Mar 19, 2023	U-1595			
	8802717	Mar 19, 2023	U-1561			
	9018243	Mar 19, 2023	U-1505			
	9018243	Mar 19, 2023	U-1595			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>APREMITLAST - OTEZLA</u></b>						
N 205437 002	6020358	Oct 30, 2018	DS DP U-1504		I-694	Sep 23, 2017
	6962940	Mar 19, 2023	U-1504		NCE	Mar 21, 2019
	7208516	Mar 19, 2023	U-1505			
	7427638	Nov 17, 2024	DS DP			
	7659302	Mar 19, 2023	U-1505			
	7659302	Mar 19, 2023	U-1595			
	7893101	Dec 09, 2023	DS DP			
	8455536	Mar 19, 2023	U-1505			
	8455536	Mar 19, 2023	U-1595			
	8802717	Mar 19, 2023	U-1561			
	9018243	Mar 19, 2023	U-1505			
	9018243	Mar 19, 2023	U-1595			
<b><u>APREMITLAST - OTEZLA</u></b>						
N 205437 003	6020358	Oct 30, 2018	DS DP U-1504		I-694	Sep 23, 2017
	6962940	Mar 19, 2023	U-1504		NCE	Mar 21, 2019
	7208516	Mar 19, 2023	U-1505			
	7427638	Nov 17, 2024	DS DP			
	7659302	Mar 19, 2023	U-1505			
	7659302	Mar 19, 2023	U-1595			
	7893101	Dec 09, 2023	DS DP			
	8455536	Mar 19, 2023	U-1505			
	8455536	Mar 19, 2023	U-1595			
	8802717	Mar 19, 2023	U-1561			
	9018243	Mar 19, 2023	U-1505			
	9018243	Mar 19, 2023	U-1595			
<b><u>APREPITANT - EMEND</u></b>						
N 021549 001	6096742	Jul 01, 2018	DS DP U-1743		NPP	Aug 28, 2018
	6096742	Jul 01, 2018	DS DP U-1744			
	6096742	Jul 01, 2018	DS DP U-745			
	8258132	Sep 26, 2027	DP U-1743			
	8258132	Sep 26, 2027	DP U-901			
<b><u>APREPITANT - EMEND</u></b>						
N 021549 002	6096742	Jul 01, 2018	DS DP U-1743		NPP	Aug 28, 2018
	6096742	Jul 01, 2018	DS DP U-1744			
	6096742	Jul 01, 2018	DS DP U-745			
	8258132	Sep 26, 2027	DP U-1743			
	8258132	Sep 26, 2027	DP U-901			
<b><u>APREPITANT - EMEND</u></b>						
N 021549 003	6096742	Jul 01, 2018	DS DP U-1743		NPP	Aug 28, 2018
	6096742	Jul 01, 2018	DS DP U-1744			
	6096742	Jul 01, 2018	DS DP U-745			
	8258132	Sep 26, 2027	DP U-1743			
	8258132	Sep 26, 2027	DP U-901			
<b><u>APREPITANT - EMEND</u></b>						
N 207865 001	6096742	Jul 01, 2018	DS DP U-1916		NPP	Aug 28, 2018
	8258132	Sep 26, 2027	DP U-1916			
<b><u>APREPITANT - CINVANTI</u></b>						
N 209296 001	9561229	Sep 18, 2035	DP U-2161			
	9808465	Sep 18, 2035	U-2161			
<b><u>ARFORMOTEROL TARTRATE - BROVANA</u></b>						
N 021912 001	6472563	Nov 09, 2021	DS			
	6667344	Jun 22, 2021	DP			
	6720453	Nov 09, 2021	DS			
	6814953	Jun 22, 2021	U-793			
	7145036	Nov 09, 2021	DS			
	7348362	Jun 22, 2021	DP U-793			
	7462645	Jun 22, 2021	U-793			
	7465756	Jun 22, 2021	DP			
	7473710	Jun 22, 2021	U-793			
	7541385	Jun 22, 2021	U-793			
	8110706	Nov 09, 2021	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>ARGATROBAN - ARGATROBAN IN SODIUM CHLORIDE</b>						
N 022434 001	7589106	Sep 26, 2027	DP	U-1163		
	7687516	Sep 26, 2027	DP	U-1164		
<b>ARIPIPRAZOLE - ABILIFY</b>						
N 021436 001	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			
<b>ARIPIPRAZOLE - ABILIFY</b>						
N 021436 002	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			
<b>ARIPIPRAZOLE - ABILIFY</b>						
N 021436 003	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			
<b>ARIPIPRAZOLE - ABILIFY</b>						
N 021436 004	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021436 005	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021436 006	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021713 001	6977257	Apr 24, 2022	DP		I-700	Dec 12, 2017
	6977257*PED	Oct 24, 2022			ODE-80	Dec 12, 2021
	7053092	Jan 28, 2022	U-839			
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8759350	Mar 02, 2027	U-1529			
	9387182	Dec 25, 2023	U-1529			
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021729 002	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8518421	Jan 24, 2021	DP			
	8518421*PED	Jul 24, 2021				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027	U-1529			
	9089567	Jan 28, 2022	U-543			
	9125939	Jul 28, 2026	U-1749			
	9358207	Apr 12, 2020	DP			
	9359302	Sep 25, 2022	DS DP U-1859			
	9387182	Dec 25, 2023	U-1529			
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021729 003	7053092	Jan 28, 2022	U-839		I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8518421	Jan 24, 2021	DP			
	8518421*PED	Jul 24, 2021				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022	U-1492			
	8642600*PED	Jul 28, 2022				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021729 003	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	8759350	Mar 02, 2027		U-1529		
	9089567	Jan 28, 2022		U-543		
	9125939	Jul 28, 2026		U-1749		
	9358207	Apr 12, 2020	DP			
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021729 004	7053092	Jan 28, 2022		U-839	I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8518421	Jan 24, 2021		DP		
	8518421*PED	Jul 24, 2021				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	9358207	Apr 12, 2020	DP			
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<b><u>ARIPIPRAZOLE - ABILIFY</u></b>						
N 021729 005	7053092	Jan 28, 2022		U-839	I-700	Dec 12, 2017
	8017615	Jun 16, 2024	DP		ODE-80	Dec 12, 2021
	8017615*PED	Dec 16, 2024				
	8518421	Jan 24, 2021		DP		
	8518421*PED	Jul 24, 2021				
	8580796	Sep 25, 2022	DS			
	8580796*PED	Mar 25, 2023				
	8642600	Jan 28, 2022		U-1492		
	8642600*PED	Jul 28, 2022				
	8642760	Sep 25, 2022	DS			
	8642760*PED	Mar 25, 2023				
	9358207	Apr 12, 2020	DP			
	9359302	Sep 25, 2022	DS DP	U-1859		
	9387182	Dec 25, 2023		U-1529		
<b><u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u></b>						
N 202971 001	7807680	Oct 19, 2024	DP		I-746	Jul 27, 2020
	8030313	Oct 19, 2024		U-1632	M-150	Dec 05, 2017
	8030313	Oct 19, 2024		U-543		
	8338427	Mar 15, 2025	DP	U-1633		
	8338427	Mar 15, 2025	DP	U-543		
	8338428	Aug 06, 2023	DP	U-1633		
	8338428	Aug 06, 2023	DP	U-543		
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024		DP		
	8759351	Aug 06, 2023	DP	U-1530		
	8759351	Aug 06, 2023	DP	U-1633		
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022		U-543		
<b><u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u></b>						
N 202971 002	7807680	Oct 19, 2024	DP		I-746	Jul 27, 2020
	8030313	Oct 19, 2024		U-1632	M-150	Dec 05, 2017
	8030313	Oct 19, 2024		U-543		
	8338427	Mar 15, 2025	DP	U-1633		
	8338427	Mar 15, 2025	DP	U-543		
	8338428	Aug 06, 2023	DP	U-1633		
	8338428	Aug 06, 2023	DP	U-543		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u></b>						
N 202971 002	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
	8759351	Aug 06, 2023	DP U-1530			
	8759351	Aug 06, 2023	DP U-1633			
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022	U-543			
<b><u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u></b>						
N 202971 003	7807680	Oct 19, 2024	DP	I-746	Jul 27, 2020	
	8030313	Oct 19, 2024	U-1632	M-150	Dec 05, 2017	
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8338428	Aug 06, 2023	DP U-1633			
	8338428	Aug 06, 2023	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
	8759351	Aug 06, 2023	DP U-1530			
	8759351	Aug 06, 2023	DP U-1633			
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022	U-543			
<b><u>ARIPIPRAZOLE - ABILIFY MAINTENA KIT</u></b>						
N 202971 004	7807680	Oct 19, 2024	DP	I-746	Jul 27, 2020	
	8030313	Oct 19, 2024	U-1632	M-150	Dec 05, 2017	
	8030313	Oct 19, 2024	U-543			
	8338427	Mar 15, 2025	DP U-1633			
	8338427	Mar 15, 2025	DP U-543			
	8338428	Aug 06, 2023	DP U-1633			
	8338428	Aug 06, 2023	DP U-543			
	8399469	Jun 29, 2025	DS			
	8722679	Oct 19, 2024	DP			
	8759351	Aug 06, 2023	DP U-1530			
	8759351	Aug 06, 2023	DP U-1633			
	8993761	Sep 25, 2022	DS			
	9089567	Jan 28, 2022	U-543			
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 001	7053092	Jan 28, 2022	U-1529	I-746	Jul 27, 2020	
	7978064	Sep 14, 2026	DP	M-150	Dec 05, 2017	
	8017615	Jun 16, 2024	DP			
	8114021	Nov 02, 2026	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9089567	Jan 28, 2022	U-543			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9270503	Sep 19, 2034	DP U-2169			
	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			
	9359302	Sep 25, 2022	DS DP U-543			
	9387182	Dec 25, 2023	U-1529			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 001	9577864	Oct 03, 2033	DP U-2169			
	9787511	Sep 19, 2034	DP U-2169			
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 002	7053092	Jan 28, 2022	U-1529		I-746	Jul 27, 2020
	7978064	Sep 14, 2026	DP		M-150	Dec 05, 2017
	8017615	Jun 16, 2024	DP			
	8114021	Nov 02, 2026	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9089567	Jan 28, 2022	U-543			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9270503	Sep 19, 2034	DP U-2169			
	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			
	9359302	Sep 25, 2022	DS DP U-543			
	9387182	Dec 25, 2023	U-1529			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9577864	Oct 03, 2033	DP U-2169			
	9787511	Sep 19, 2034	DP U-2169			
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 003	7053092	Jan 28, 2022	U-1529		I-746	Jul 27, 2020
	7978064	Sep 14, 2026	DP		M-150	Dec 05, 2017
	8017615	Jun 16, 2024	DP			
	8114021	Nov 02, 2026	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9089567	Jan 28, 2022	U-543			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9270503	Sep 19, 2034	DP U-2169			
	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			
	9359302	Sep 25, 2022	DS DP U-543			
	9387182	Dec 25, 2023	U-1529			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 003	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9577864	Oct 03, 2033	DP U-2169			
	9787511	Sep 19, 2034	DP U-2169			
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 004	7053092	Jan 28, 2022	U-1529		I-746	Jul 27, 2020
	7978064	Sep 14, 2026	DP		M-150	Dec 05, 2017
	8017615	Jun 16, 2024	DP			
	8114021	Nov 02, 2026	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9089567	Jan 28, 2022	U-543			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9270503	Sep 19, 2034	DP U-2169			
	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			
	9359302	Sep 25, 2022	DS DP U-543			
	9387182	Dec 25, 2023	U-1529			
	9433371	Sep 15, 2029	DP			
	9444503	Nov 19, 2027	DP U-2169			
	9577864	Oct 03, 2033	DP U-2169			
	9787511	Sep 19, 2034	DP U-2169			
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 005	7053092	Jan 28, 2022	U-1529		I-746	Jul 27, 2020
	7978064	Sep 14, 2026	DP		M-150	Dec 05, 2017
	8017615	Jun 16, 2024	DP			
	8114021	Nov 02, 2026	DP			
	8258962	Nov 25, 2030	DP			
	8545402	Apr 27, 2030	DP			
	8547248	Dec 18, 2030	DP U-2167			
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	8674825	Apr 09, 2029	DP U-2170			
	8718193	Dec 05, 2029	DP			
	8759350	Mar 02, 2027	U-1529			
	8847766	Mar 29, 2030	DP U-2167			
	8945005	Aug 19, 2029	DP U-2167			
	8956288	Jul 06, 2029	DP U-2167			
	8961412	Nov 17, 2030	DP			
	9060708	Mar 05, 2029	DP			
	9089567	Jan 28, 2022	U-543			
	9119554	Dec 16, 2028	DP			
	9125939	Jul 28, 2026	U-1749			
	9149577	Dec 15, 2029	DP			
	9258035	Mar 05, 2029	DP			
	9268909	Oct 15, 2033	DP U-2168			
	9270503	Sep 19, 2034	DP U-2169			
	9320455	Dec 15, 2031	DP			
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 005	9359302	Sep 25, 2022	DS DP U-543			
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029		DP		
	9444503	Nov 19, 2027		DP U-2169		
	9577864	Oct 03, 2033		DP U-2169		
	9787511	Sep 19, 2034		DP U-2169		
<b><u>ARIPIPRAZOLE - ABILIFY MYCITE KIT</u></b>						
N 207202 006	7053092	Jan 28, 2022		U-1529		
	7978064	Sep 14, 2026		DP	I-746	Jul 27, 2020
	8017615	Jun 16, 2024		DP	M-150	Dec 05, 2017
	8114021	Nov 02, 2026		DP		
	8258962	Nov 25, 2030		DP		
	8545402	Apr 27, 2030		DP		
	8547248	Dec 18, 2030		DP U-2167		
	8580796	Sep 25, 2022	DS			
	8642760	Sep 25, 2022	DS			
	8674825	Apr 09, 2029		DP U-2170		
	8718193	Dec 05, 2029		DP		
	8759350	Mar 02, 2027		U-1529		
	8847766	Mar 29, 2030		DP U-2167		
	8945005	Aug 19, 2029		DP U-2167		
	8956288	Jul 06, 2029		DP U-2167		
	8961412	Nov 17, 2030		DP		
	9060708	Mar 05, 2029		DP		
	9089567	Jan 28, 2022		U-543		
	9119554	Dec 16, 2028		DP		
	9125939	Jul 28, 2026		U-1749		
	9149577	Dec 15, 2029		DP		
	9258035	Mar 05, 2029		DP		
	9268909	Oct 15, 2033		DP U-2168		
	9270503	Sep 19, 2034		DP U-2169		
	9320455	Dec 15, 2031		DP		
	9359302	Sep 25, 2022	DS DP U-1529			
	9359302	Sep 25, 2022	DS DP U-1749			
	9359302	Sep 25, 2022	DS DP U-543			
	9387182	Dec 25, 2023		U-1529		
	9433371	Sep 15, 2029		DP		
	9444503	Nov 19, 2027		DP U-2169		
	9577864	Oct 03, 2033		DP U-2169		
	9787511	Sep 19, 2034		DP U-2169		
<b><u>ARIPIPRAZOLE LAUROXIL - ARISTADA</u></b>						
N 207533 001	8431576	Oct 26, 2030	DS		NCE	Oct 05, 2020
	8796276	Jun 24, 2030		U-543		
	9034867	Nov 07, 2032		DP U-543		
	9193685	Oct 24, 2033		DP U-543		
	9452131	Mar 19, 2035		U-543		
	9526726	Mar 19, 2035		DP		
<b><u>ARIPIPRAZOLE LAUROXIL - ARISTADA</u></b>						
N 207533 002	8431576	Oct 26, 2030	DS		NCE	Oct 05, 2020
	8796276	Jun 24, 2030		U-543		
	9034867	Nov 07, 2032		DP U-543		
	9193685	Oct 24, 2033		DP U-543		
	9452131	Mar 19, 2035		U-543		
	9526726	Mar 19, 2035		DP		
<b><u>ARIPIPRAZOLE LAUROXIL - ARISTADA</u></b>						
N 207533 003	8431576	Oct 26, 2030	DS		NCE	Oct 05, 2020
	8796276	Jun 24, 2030		U-543		
	9034867	Nov 07, 2032		DP U-543		
	9193685	Oct 24, 2033		DP U-543		
	9452131	Mar 19, 2035		U-543		
	9526726	Mar 19, 2035		DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ARIPIPRAZOLE LAUROXIL - ARISTADA</u></b>						
N 207533 004	8431576	Oct 26, 2030	DS		NCE	Oct 05, 2020
	8796276	Jun 24, 2030		U-543		
	9034867	Nov 07, 2032		DP U-543		
	9193685	Oct 24, 2033		DP U-543		
	9452131	Mar 19, 2035		U-543		
<b><u>ARMODAFINIL - NUvigil</u></b>						
N 021875 001	7132570	Dec 18, 2023	DS DP			
	7297346	Nov 29, 2023		DP		
<b><u>ARMODAFINIL - NUvigil</u></b>						
N 021875 002	7132570	Dec 18, 2023	DS DP			
	7297346	Nov 29, 2023		DP		
<b><u>ARMODAFINIL - NUvigil</u></b>						
N 021875 003	7132570	Dec 18, 2023	DS DP			
	7297346	Nov 29, 2023		DP		
<b><u>ARMODAFINIL - NUvigil</u></b>						
N 021875 004	7132570	Dec 18, 2023	DS DP			
	7297346	Nov 29, 2023		DP		
<b><u>ARMODAFINIL - NUvigil</u></b>						
N 021875 005	7132570	Dec 18, 2023	DS DP			
	7297346	Nov 29, 2023		DP		
<b><u>ARSENIC TRIOXIDE - TRISENOX</u></b>						
N 021248 001	6723351	Nov 10, 2018	U-573			
	6855339	Nov 10, 2018	U-617			
	6861076	Nov 10, 2018	U-617			
	6884439	Nov 10, 2018	U-651			
	6982096	Nov 10, 2018	U-651			
	8273379	Nov 10, 2018	U-1291			
<b><u>ARSENIC TRIOXIDE - TRISENOX</u></b>						
N 021248 002	6723351	Nov 10, 2018	U-2204			
	6855339	Nov 10, 2018	U-2204			
	6861076	Nov 10, 2018	U-2204			
	6884439	Nov 10, 2018	U-2204			
	6982096	Nov 10, 2018	U-2204			
	8273379	Nov 10, 2018	U-2204			
<b><u>ASCORBIC ACID - ASCOR</u></b>						
N 209112 001					ODE	Oct 02, 2024
<b><u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u></b>						
N 021881 001	7169381	Sep 01, 2024	DS DP			
	7658914	Sep 01, 2024	DS DP			
<b><u>ASENAPINE MALEATE - SAPHRIS</u></b>						
N 022117 001	5763476	Jun 09, 2020	DP U-1960		D-166	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1961		I-597	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1962		M-158	Mar 17, 2018
	5763476	Jun 09, 2020	DP U-1963		NPP	Mar 17, 2018
	5763476	Jun 09, 2020	DP U-326		PED	Sep 17, 2018
	5763476*PED	Dec 09, 2020			PED	Sep 17, 2018
	7741358	Apr 06, 2026	DS DP U-1064			
	7741358	Apr 06, 2026	DS DP U-1960			
	7741358	Apr 06, 2026	DS DP U-1961			
	7741358	Apr 06, 2026	DS DP U-1962			
	7741358	Apr 06, 2026	DS DP U-1963			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<b><u>ASENAPINE MALEATE - SAPHRIS</u></b>						
N 022117 002	5763476	Jun 09, 2020	DP U-1960		D-166	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1961		I-597	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1962		M-158	Mar 17, 2018

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ASENAPINE MALEATE - SAPHRIS</u></b>						
N 022117 002	5763476	Jun 09, 2020	DP U-1963		NPP	Mar 17, 2018
	5763476	Jun 09, 2020	DP U-326		PED	Sep 17, 2018
	5763476*PED	Dec 09, 2020			PED	Sep 17, 2018
	7741358	Apr 06, 2026	DS DP U-1064			
	7741358	Apr 06, 2026	DS DP U-1960			
	7741358	Apr 06, 2026	DS DP U-1961			
	7741358	Apr 06, 2026	DS DP U-1962			
	7741358	Apr 06, 2026	DS DP U-1963			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<b><u>ASENAPINE MALEATE - SAPHRIS</u></b>						
N 022117 003	5763476	Jun 09, 2020	DP U-1893		D-166	Jan 13, 2020
	5763476	Jun 09, 2020	DP U-1966		I-597	Jan 13, 2020
	5763476*PED	Dec 09, 2020				
	7741358	Apr 06, 2026	DS DP U-1893			
	7741358	Apr 06, 2026	DS DP U-1966			
	7741358*PED	Oct 06, 2026				
	8022228	Apr 06, 2026	DS DP			
	8022228*PED	Oct 06, 2026				
<b><u>ASPIRIN - ASPIRIN</u></b>						
N 203697 001	8865187	Mar 23, 2022	DP			
	9101637	Mar 23, 2022	U-1731			
	9101637	Mar 23, 2022	U-1732			
	9101637	Mar 23, 2022	U-1733			
	9216150	Sep 29, 2032	DP			
	9226892	Sep 29, 2032	U-1731			
	9226892	Sep 29, 2032	U-1732			
	9226892	Sep 29, 2032	U-1733			
	9351984	Dec 19, 2021	DP			
<b><u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u></b>						
N 205103 001	6926907	Feb 28, 2023	DP U-1902		NC	Sep 14, 2019
	8206741	Feb 28, 2023	DP U-1902			
	9364439	May 31, 2022	DP U-1902			
	9539214	Mar 13, 2033	U-1902			
<b><u>ASPIRIN; OMEPRAZOLE - YOSPRALA</u></b>						
N 205103 002	6926907	Feb 28, 2023	DP U-1902		NC	Sep 14, 2019
	8206741	Feb 28, 2023	DP U-1902			
	9364439	May 31, 2022	DP U-1902			
	9539214	Mar 13, 2033	U-1902			
<b><u>ATAZANAVIR SULFATE - ATAZANAVIR SULFATE</u></b>						
A 091673 001					PC	Jun 25, 2018
<b><u>ATAZANAVIR SULFATE - ATAZANAVIR SULFATE</u></b>						
A 091673 002					PC	Jun 25, 2018
<b><u>ATAZANAVIR SULFATE - ATAZANAVIR SULFATE</u></b>						
A 091673 003					PC	Jun 25, 2018
<b><u>ATAZANAVIR SULFATE - ATAZANAVIR SULFATE</u></b>						
A 091673 004					PC	Jun 25, 2018
<b><u>ATAZANAVIR SULFATE - REYATAZ</u></b>						
N 021567 001	5849911*PED	Dec 20, 2017				
	6087383	Dec 21, 2018	DS DP			
	6087383*PED	Jun 21, 2019				
<b><u>ATAZANAVIR SULFATE - REYATAZ</u></b>						
N 021567 002	5849911*PED	Dec 20, 2017				
	6087383	Dec 21, 2018	DS DP			
	6087383*PED	Jun 21, 2019				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ATAZANAVIR SULFATE - REYATAZ</u></b>						
N 021567 003	5849911*PED 6087383 6087383*PED	Dec 20, 2017 Dec 21, 2018 Jun 21, 2019		DS DP		
N 021567 004	5849911*PED 6087383 6087383*PED	Dec 20, 2017 Dec 21, 2018 Jun 21, 2019		DS DP		
<b><u>ATAZANAVIR SULFATE - REYATAZ</u></b>						
N 206352 001	5849911*PED 6087383 6087383*PED	Dec 20, 2017 Dec 21, 2018 Jun 21, 2019		DS DP	NPP PED	Sep 24, 2018 Mar 24, 2019
<b><u>ATAZANAVIR SULFATE; COBICISTAT - EVOTAZ</u></b>						
N 206353 001	5849911*PED 6087383 6087383*PED 8148374	Dec 20, 2017 Dec 21, 2018 Jun 21, 2019 Sep 03, 2029		DS DP	NCE	Aug 27, 2017
<b><u>ATORVASTATIN CALCIUM - LIPITOR</u></b>						
N 020702 001					M-204	Jun 23, 2020
<b><u>ATORVASTATIN CALCIUM - LIPITOR</u></b>						
N 020702 002					M-204	Jun 23, 2020
<b><u>AVANAFILE - STENDRA</u></b>						
N 202276 001	6656935 7501409	Apr 27, 2025 May 05, 2023		DS DP U-155 DP	D-140	Sep 17, 2017
<b><u>AVANAFILE - STENDRA</u></b>						
N 202276 002	6656935 7501409	Apr 27, 2025 May 05, 2023		DS DP U-155 DP	D-140	Sep 17, 2017
<b><u>AVANAFILE - STENDRA</u></b>						
N 202276 003	6656935 7501409	Apr 27, 2025 May 05, 2023		DS DP U-155 DP	D-140	Sep 17, 2017
<b><u>AVIBACTAM SODIUM; CEFTAZIDIME - AVYCAZ</u></b>						
N 206494 001	7112592 7612087 8178554 8471025 8835455 8969566 9284314 9695122	Feb 24, 2022 Nov 12, 2026 Jul 24, 2021 Aug 12, 2031 Oct 08, 2030 Jun 15, 2032 Jun 15, 2032 Jun 15, 2032		DS DP U-282 DP DS DP U-282 DS DP DS DS DS		
<b><u>AXITINIB - INLYTA</u></b>						
N 202324 001	6534524 7141581 8791140	Apr 29, 2025 Jun 30, 2020 Dec 14, 2030		DS DP U-1220 DS		
<b><u>AXITINIB - INLYTA</u></b>						
N 202324 002	6534524 7141581 8791140	Apr 29, 2025 Jun 30, 2020 Dec 14, 2030		DS DP U-1220 DS		
<b><u>AZELAIC ACID - FINACEA</u></b>						
N 021470 001	6534070	Nov 18, 2018				
<b><u>AZELAIC ACID - FINACEA</u></b>						
N 207071 001	6730288 7700076 8435498 8722021 8900554 9211259 9265725	Sep 08, 2019 Sep 18, 2027 Mar 01, 2024 Oct 24, 2023 Oct 24, 2023 Jan 26, 2029 Dec 08, 2027		DP DP U-1727 DP DP U-1796 DP	NP	Jul 29, 2018

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>AZELASTINE HYDROCHLORIDE - ASTEPRO</b>						
N 022203 001	8071073	Jun 04, 2028	DP		NPP	Feb 20, 2018
	8518919	Nov 22, 2025	U-1430		NPP	Feb 20, 2018
<b>AZELASTINE HYDROCHLORIDE - ASTEPRO</b>						
N 022203 002	8071073	Jun 04, 2028	DP			
	8518919	Nov 22, 2025	U-1430			
<b>AZELASTINE HYDROCHLORIDE; FLUTICASONE PROPIONATE - DYMISTA</b>						
N 202236 001	8163723	Aug 29, 2023	U-1667		NPP	Feb 20, 2018
	8163723	Aug 29, 2023	U-644		PED	Aug 20, 2018
	8163723	Aug 29, 2023	U-707			
	8163723	Aug 29, 2023	U-77			
	8163723	Aug 29, 2023	U-81			
	8163723*PED	Feb 29, 2024				
	8168620	Feb 24, 2026	DP			
	9259428	Jun 13, 2023	U-644			
	9259428*PED	Dec 13, 2023				
<b>AZILSARTAN KAMEDOXOMIL - EDARBI</b>						
N 200796 001	7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
<b>AZILSARTAN KAMEDOXOMIL - EDARBI</b>						
N 200796 002	7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
<b>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</b>						
N 202331 001	7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
	9169238	Feb 04, 2030	DP			
<b>AZILSARTAN KAMEDOXOMIL; CHLORTHALIDONE - EDARBYCLOR</b>						
N 202331 002	7157584	May 22, 2025	DS			
	7572920	Jan 07, 2025	DP U-3			
	9066936	Mar 26, 2028	DP			
	9169238	Feb 04, 2030	DP			
<b>AZITHROMYCIN - AZITHROMYCIN</b>						
A 065488 001					PC	Mar 26, 2018
<b>AZITHROMYCIN - AZITHROMYCIN</b>						
A 065488 002					PC	Mar 26, 2018
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050693 001	6268489	Jul 31, 2018	DS			
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050710 001	6268489	Jul 31, 2018	DS			
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050710 002	6268489	Jul 31, 2018	DS			
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050711 001	6268489	Jul 31, 2018	DS			
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050730 001	6268489	Jul 31, 2018	DS			
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050733 001	6268489	Jul 31, 2018	DS			
<b>AZITHROMYCIN - ZITHROMAX</b>						
N 050784 001	6268489	Jul 31, 2018	DS			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>AZITHROMYCIN - ZMAX</b>						
N 050797 001	6268489	Jul 31, 2018	DS			
	6984403	Feb 14, 2024	DP	U-282		
	7887844	Feb 14, 2024	DP			
<b>AZITHROMYCIN - AZASITE</b>						
N 050810 001	6159458	Nov 04, 2017	DP	U-709		
	6239113	Mar 31, 2019		U-709		
	6569443	Mar 31, 2019	DP	U-709		
	6861411	Nov 25, 2018		U-709		
	7056893	Mar 31, 2019	DP	U-709		
<b>AZTREONAM - CAYSTON</b>						
N 050814 001	7208141	Dec 20, 2021	DP	U-1031		
	7214364	Dec 20, 2021	DP			
	7427633	Dec 20, 2021	DP	U-1031		
	8399496	Dec 20, 2021	DP	U-1377		
<b>BACLOFEN - KEMSTRO</b>						
N 021589 001	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b>BACLOFEN - KEMSTRO</b>						
N 021589 002	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b>BALSALAZIDE DISODIUM - COLAZAL</b>						
N 020610 001	7452872	Aug 24, 2026		U-141		
	7625884	Aug 24, 2026		U-141		
<b>BALSALAZIDE DISODIUM - GIAZO</b>						
N 022205 001	6197341	Mar 13, 2018	DP	U-1229		
	7452872	Aug 24, 2026		U-1229		
	7625884	Aug 24, 2026		U-1229		
	8497256	Jun 23, 2031		U-1229		
	9192616	Aug 02, 2026		U-1229		
<b>BAZEDOXIFENE ACETATE; ESTROGENS, CONJUGATED - DUAVEE</b>						
N 022247 001	5998402	Apr 04, 2018	DS	DP U-594	NCE	Oct 13, 2018
	6479535	May 06, 2019	DP	U-594		
	6479535	May 06, 2019	DP	U-904		
	7683051	Mar 10, 2027	DS	DP U-594		
	7683051	Mar 10, 2027	DS	DP U-904		
	8815934	May 06, 2019	DP			
<b>BECLOMETHASONE DIPROPIONATE - QVAR 80</b>						
N 020911 001	6446627	Dec 18, 2017	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<b>BECLOMETHASONE DIPROPIONATE - QVAR 40</b>						
N 020911 002	6446627	Dec 18, 2017	DP			
	9463289	May 18, 2031	DP			
	9808587	May 18, 2031	DP			
<b>BECLOMETHASONE DIPROPIONATE - ONASL</b>						
N 202813 001	7780038	Jan 24, 2027	DP			
<b>BECLOMETHASONE DIPROPIONATE - QNASL</b>						
N 202813 002	7780038	Jan 24, 2027	DP		NS	Dec 17, 2017
<b>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</b>						
N 207921 001	6446627	Dec 18, 2017	DP			
	7637260	Aug 25, 2020	DP			
	8132712	Sep 07, 2028	DP			
	8931476	Jul 17, 2031	DP			
<b>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</b>						
N 207921 002	6446627	Dec 18, 2017	DP			
	7637260	Aug 25, 2020	DP			
	8132712	Sep 07, 2028	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BECLOMETHASONE DIPROPIONATE - QVAR REDIHALER</u></b>						
N 207921 002	8931476	Jul 17, 2031	DP			
<b><u>BEDAQUILINE FUMARATE - SIRTURO</u></b>						
N 204384 001	7498343	Dec 01, 2026	DS DP U-1321		NCE	Dec 28, 2017
	8546428	Mar 19, 2029	DS DP U-1321		ODE-38	Dec 28, 2019
<b><u>BELINOSTAT - BELEODAQ</u></b>						
N 206256 001	6888027	Sep 27, 2021	DS DP U-1544		NCE	Jul 03, 2019
	8835501	Oct 27, 2027	DP		ODE-68	Jul 03, 2021
<b><u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u></b>						
N 022249 001	8436190	Oct 26, 2030	DP			
	8436190*PED	Apr 26, 2031				
	8445524	Mar 26, 2029	DS DP U-1402			
	8445524*PED	Sep 26, 2029				
	8609863	Jan 12, 2026	DP			
	8609863*PED	Jul 12, 2026				
	8669279	Mar 26, 2029	DP U-1402			
	8669279*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
	8883836	Mar 26, 2029	DP U-1402			
	8883836*PED	Sep 26, 2029				
	8895756	Jan 12, 2026	DP			
	8895756*PED	Jul 12, 2026				
	9533955	Mar 26, 2029	DP U-1949			
	9533955	Mar 26, 2029	DP U-1952			
<b><u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u></b>						
N 022249 002	8436190	Oct 26, 2030	DP			
	8436190*PED	Apr 26, 2031				
	8445524	Mar 26, 2029	DS DP U-1402			
	8445524*PED	Sep 26, 2029				
	8609863	Jan 12, 2026	DP			
	8609863*PED	Jul 12, 2026				
	8669279	Mar 26, 2029	DP U-1402			
	8669279*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
	8883836	Mar 26, 2029	DP U-1402			
	8883836*PED	Sep 26, 2029				
	8895756	Jan 12, 2026	DP			
	8895756*PED	Jul 12, 2026				
	9533955	Mar 26, 2029	DP U-1949			
	9533955	Mar 26, 2029	DP U-1952			
<b><u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u></b>						
N 022249 003	8344006	Sep 23, 2029	DP U-1402			
	8344006*PED	Mar 23, 2030				
	8445524	Mar 26, 2029	DS			
	8445524*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
<b><u>BENDAMUSTINE HYDROCHLORIDE - TREANDA</u></b>						
N 022249 004	8344006	Sep 23, 2029	DP U-1402			
	8344006*PED	Mar 23, 2030				
	8445524	Mar 26, 2029	DS			
	8445524*PED	Sep 26, 2029				
	8791270	Jan 12, 2026	DP U-1542			
	8791270*PED	Jul 12, 2026				
<b><u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u></b>						
N 208194 001	8609707	Aug 11, 2031	DP U-1542			
	8791270	Jan 12, 2026	DP U-1790			
	8791270*PED	Jul 12, 2026				
	9000021	Mar 15, 2033	U-1542			
	9034908	Mar 15, 2033	U-1542			
	9144568	Mar 15, 2033	U-1542			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BENDAMUSTINE HYDROCHLORIDE - BENDEKA</u></b>						
N 208194 001	9265831	Jan 28, 2031	DP			
	9572796	Jan 28, 2031	DP U-1971			
	9572796	Jan 28, 2031	DP U-1972			
	9572797	Jan 28, 2031	U-1971			
	9572797	Jan 28, 2031	U-1972			
	9572887	Mar 15, 2033	U-1971			
	9572887	Mar 15, 2033	U-1972			
	9579384	Mar 15, 2033	U-1971			
	9579384	Mar 15, 2033	U-1972			
	9597397	Mar 15, 2033	U-1971			
	9597397	Mar 15, 2033	U-1972			
	9597398	Mar 15, 2033	U-1971			
	9597399	Mar 15, 2033	U-1971			
	9597399	Mar 15, 2033	U-1972			
<b><u>BENZNIDAZOLE - BENZNIDAZOLE</u></b>						
N 209570 001					NCE	Aug 29, 2022
					ODE-154	Aug 29, 2024
<b><u>BENZNIDAZOLE - BENZNIDAZOLE</u></b>						
N 209570 002					NCE	Aug 29, 2022
					ODE-154	Aug 29, 2024
<b><u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ACANYA</u></b>						
N 050819 001	8288434	Aug 05, 2029	DP U-124			
	8663699	Jun 03, 2029	U-124			
	8895070	Jun 03, 2029	U-124			
	9078870	Jun 03, 2029	DP			
<b><u>BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE - ONEXTON</u></b>						
N 050819 002	8288434	Aug 05, 2029	DP U-1033			
	8288434	Aug 05, 2029	DP U-124			
	8288434	Aug 05, 2029	DP U-134			
	8288434	Aug 05, 2029	DP U-818			
	8288434	Aug 05, 2029	DP U-916			
	8288434	Aug 05, 2029	DP U-921			
	9504704	Jun 03, 2029	DP U-124			
	9504704	Jun 03, 2029	DP U-134			
	9504704	Jun 03, 2029	DP U-818			
	9504704	Jun 03, 2029	DP U-916			
	9561208	Jun 03, 2029	DP U-916			
<b><u>BENZYL ALCOHOL - ULESFIA</u></b>						
N 022129 001	5858383	Aug 11, 2017	U-970			
	6139859	Aug 11, 2017	U-970			
	6793931	Jul 11, 2022	DP U-970			
	7294342	May 19, 2024	U-970			
<b><u>BEPOTASTINE BESILATE - BEPREVE</u></b>						
N 022288 001	6780877	Sep 19, 2019	DS DP			
	8784789	Sep 05, 2024	DP			
	8877168	Jul 30, 2023	DP			
<b><u>BESIFLOXACIN HYDROCHLORIDE - BESIVANCE</u></b>						
N 022308 001	6685958	Jun 29, 2021	DP U-80			
	6699492	Mar 31, 2019	DP U-80			
	8415342	Nov 07, 2030	U-80			
	8481526	Jan 09, 2031	DS			
	8604020	Mar 12, 2030	DP			
	8937062	Nov 13, 2029	U-80			
<b><u>BETAMETHASONE DIPROPIONATE - SERNIVO</u></b>						
N 208079 001	9364485	Aug 31, 2030	DP U-1858		NDF	Feb 05, 2019
	9433630	Aug 31, 2030	DP U-1858			
	9439911	Aug 31, 2030	DP U-1858			
	9655907	Aug 31, 2030	DP U-1858			
	9775851	Aug 31, 2030	DP U-1858			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE - ENSTILAR</u></b>						
N 207589 001	6753013	Jan 27, 2020	DP U-1761		NP	Oct 16, 2018
	9119781	Jun 10, 2031	DP U-1761			
<b><u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX</u></b>						
N 021852 001	6753013	Jan 27, 2020	DP U-193		NPP	Dec 23, 2017
	6753013	Jan 27, 2020	DP U-88			
<b><u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX</u></b>						
N 022185 001	6753013	Jan 27, 2020	DP U-1761		NPP	Aug 29, 2017
	6753013	Jan 27, 2020	DP U-193			
	6753013	Jan 27, 2020	DP U-88			
	6787529	Jan 27, 2020	DP U-1761			
	6787529	Jan 27, 2020	DP U-193			
	6787529	Jan 27, 2020	DP U-88			
<b><u>BETRIXABAN - BEVYXXA</u></b>						
N 208383 001	6376515	Sep 15, 2020	DS DP U-1167		NCE	Jun 23, 2022
	6376515	Sep 15, 2020	DS DP U-1502			
	6376515	Sep 15, 2020	DS DP U-2029			
	6376515	Sep 15, 2020	DS DP U-2030			
	6835739	Sep 15, 2020	DS DP			
	7598276	Nov 08, 2026	DS			
	8404724	Mar 29, 2031	DP U-2034			
	8518977	Sep 15, 2020	DS			
	8557852	Sep 08, 2028	U-1167			
	8557852	Sep 08, 2028	U-2030			
	8691847	Sep 15, 2020	DS DP U-2029			
	8691847	Sep 15, 2020	DS DP U-2035			
	8987463	Dec 28, 2030	DP			
	9555023	Nov 07, 2026	U-1502			
	9629831	Sep 15, 2020	U-1167			
	9629831	Sep 15, 2020	U-1502			
	9629831	Sep 15, 2020	U-2030			
	9629831	Sep 15, 2020	U-2035			
<b><u>BETRIXABAN - BEVYXXA</u></b>						
N 208383 002	6376515	Sep 15, 2020	DS DP U-1167		NCE	Jun 23, 2022
	6376515	Sep 15, 2020	DS DP U-1502			
	6376515	Sep 15, 2020	DS DP U-2029			
	6376515	Sep 15, 2020	DS DP U-2030			
	6835739	Sep 15, 2020	DS DP			
	7598276	Nov 08, 2026	DS			
	8404724	Mar 29, 2031	DP U-2034			
	8518977	Sep 15, 2020	DS			
	8557852	Sep 08, 2028	U-1167			
	8557852	Sep 08, 2028	U-2030			
	8691847	Sep 15, 2020	DS DP U-2029			
	8691847	Sep 15, 2020	DS DP U-2035			
	8987463	Dec 28, 2030	DP			
	9555023	Nov 07, 2026	U-1502			
	9629831	Sep 15, 2020	U-1167			
	9629831	Sep 15, 2020	U-1502			
	9629831	Sep 15, 2020	U-2030			
	9629831	Sep 15, 2020	U-2035			
<b><u>BEXAROTENE - TARGRETIN</u></b>						
N 021055 001					M-164	Jul 29, 2018
<b><u>BIMATOPROST - LUMIGAN</u></b>						
N 022184 001	7851504	Jun 13, 2027	DS DP			
	8278353	Mar 16, 2025	DP			
	8299118	Mar 16, 2025	U-1295			
	8309605	Mar 16, 2025	U-1293			
	8309605	Mar 16, 2025	U-1294			
	8338479	Mar 16, 2025	DP U-1295			
	8524777	Mar 16, 2025	U-1235			
	8586630	Mar 16, 2025	U-1458			
	8772338	Mar 16, 2025	DP U-1528			
	8933120	Mar 16, 2025	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BIMATOPROST - LUMIGAN</u></b>						
N 022184 001	8933127	Mar 16, 2025	DP			
	9155716	Mar 16, 2025	DP U-1528			
	9241918	Mar 16, 2025	DP U-1814			
<b><u>BIMATOPROST - LATISSE</u></b>						
N 022369 001	8038988	Aug 25, 2023	DS DP U-1208		M-140	Sep 04, 2017
	8101161	May 25, 2024	U-1217			
	8101161	May 25, 2024	U-1218			
	8263054	Jan 15, 2023	U-1277			
	8541466	Jan 31, 2021	U-1217			
	8632760	Jan 15, 2023	U-1487			
	8758733	Jan 15, 2023	U-1487			
	8906962	Jan 31, 2021	U-1217			
	8986715	Jan 15, 2023	U-1217			
	9216183	Jan 15, 2023	U-1487			
	9226931	Jan 15, 2023	U-1799			
	9579270	Jan 31, 2021	U-1975			
<b><u>BISACODYL; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - HALFLYTENE</u></b>						
N 021551 003	7291324	Oct 22, 2022	U-837			
<b><u>BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE - PYLERA</u></b>						
N 050786 001	6350468	Dec 14, 2018	U-932			
	6350468	Dec 14, 2018	U-956			
<b><u>BIVALIRUDIN - ANGIOMAX</u></b>						
N 020873 001	7582727	Jul 27, 2028	DP			
	7598343	Jul 27, 2028	DP			
<b><u>BOCEPREVIR - VICTRELIS</u></b>						
N 202258 001	7772178	Nov 11, 2027	DP U-1128			
	8119602	Mar 17, 2027	U-1233			
	RE43298	Dec 22, 2024	DS DP U-1128			
<b><u>BORTEZOMIB - VELCADE</u></b>						
N 021602 001	5780454*PED	Nov 03, 2017		D-141	Oct 08, 2017	
	6713446	Jan 25, 2022	DS DP	D-142	Oct 08, 2017	
	6713446*PED	Jul 25, 2022		I-695	Oct 08, 2017	
	6958319	Jan 25, 2022	DS DP	M-139	Aug 08, 2017	
	6958319*PED	Jul 25, 2022		M-165	Sep 14, 2018	
				ODE-76	Oct 08, 2021	
				PED	Feb 08, 2018	
				PED	Apr 08, 2018	
				PED	Apr 08, 2018	
				PED	Apr 08, 2018	
				PED	Mar 14, 2019	
				PED	Apr 08, 2022	
<b><u>BORTEZOMIB - BORTEZOMIB</u></b>						
N 205004 001	8962572	Nov 03, 2032	DP			
<b><u>BOSENTAN - TRACLEER</u></b>						
N 021290 001				NPP	Sep 05, 2020	
<b><u>BOSENTAN - TRACLEER</u></b>						
N 021290 002				NPP	Sep 05, 2020	
<b><u>BOSENTAN - TRACLEER</u></b>						
N 209279 001	7959945	Dec 28, 2027	DP	NP	Sep 05, 2020	
	8309126	May 15, 2026	DP	ODE	Sep 05, 2024	
<b><u>BOSUTINIB MONOHYDRATE - BOSULIF</u></b>						
N 203341 001	6002008	Mar 27, 2018	DS DP U-1284	I-759	Dec 19, 2020	
	7417148	Jan 23, 2026	U-1283	NCE	Sep 04, 2017	
	7767678	Nov 23, 2026	DS DP	ODE-30	Sep 04, 2019	
	7919625	Dec 11, 2025	DP			
	RE42376	Apr 13, 2024	DS			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BOSUTINIB MONOHYDRATE - BOSULIF</u></b>						
N 203341 002	6002008	Mar 27, 2018	DS DP U-1284		I-759	Dec 19, 2020
	7417148	Jan 23, 2026	U-1283		NCE	Sep 04, 2017
	7767678	Nov 23, 2026	DS DP		ODE-30	Sep 04, 2019
	7919625	Dec 11, 2025	DP			
	RE42376	Apr 13, 2024	DS			
<b><u>BOSUTINIB MONOHYDRATE - BOSULIF</u></b>						
N 203341 003	6002008	Mar 27, 2018	DS DP U-1284		I-759	Dec 19, 2020
	7417148	Jan 23, 2026	U-1283		ODE-30	Sep 04, 2019
	7767678	Nov 23, 2026	DS DP			
	7919625	Dec 11, 2025	DP			
	RE42376	Apr 13, 2024	DS			
<b><u>BREXIPRAZOLE - REXULTI</u></b>						
N 205422 001	7888362	Feb 23, 2027	DS		M-186	Sep 23, 2019
	8349840	Apr 12, 2026	DP U-1529		NCE	Jul 10, 2020
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<b><u>BREXIPRAZOLE - REXULTI</u></b>						
N 205422 002	7888362	Feb 23, 2027	DS		M-186	Sep 23, 2019
	8349840	Apr 12, 2026	DP U-1529		NCE	Jul 10, 2020
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<b><u>BREXIPRAZOLE - REXULTI</u></b>						
N 205422 003	7888362	Feb 23, 2027	DS		M-186	Sep 23, 2019
	8349840	Apr 12, 2026	DP U-1529		NCE	Jul 10, 2020
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<b><u>BREXIPRAZOLE - REXULTI</u></b>						
N 205422 004	7888362	Feb 23, 2027	DS		M-186	Sep 23, 2019
	8349840	Apr 12, 2026	DP U-1529		NCE	Jul 10, 2020
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<b><u>BREXIPRAZOLE - REXULTI</u></b>						
N 205422 005	7888362	Feb 23, 2027	DS		M-186	Sep 23, 2019
	8349840	Apr 12, 2026	DP U-1529		NCE	Jul 10, 2020
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<b><u>BREXIPRAZOLE - REXULTI</u></b>						
N 205422 006	7888362	Feb 23, 2027	DS		M-186	Sep 23, 2019
	8349840	Apr 12, 2026	DP U-1529		NCE	Jul 10, 2020
	8618109	Apr 12, 2026	U-543			
	9839637	Apr 12, 2026	DP U-1529			
	9839637	Apr 12, 2026	DP U-543			
<b><u>BRIGATINIB - ALUNBRIG</u></b>						
N 208772 001	9012462	Feb 06, 2031	DS		NCE	Apr 28, 2022
	9273077	May 21, 2029	U-1927		ODE-142	Apr 28, 2024
	9611283	Apr 10, 2034	U-1927			
<b><u>BRIGATINIB - ALUNBRIG</u></b>						
N 208772 002	9012462	Feb 06, 2031	DS		NCE	Apr 28, 2022
	9273077	May 21, 2029	U-1927		ODE-142	Apr 28, 2024
	9611283	Apr 10, 2034	U-1927			
<b><u>BRIGATINIB - ALUNBRIG</u></b>						
N 208772 003	9012462	Feb 06, 2031	DS		NCE	Apr 28, 2022
	9273077	May 21, 2029	U-1927		ODE-142	Apr 28, 2024
	9611283	Apr 10, 2034	U-1927			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BRIGATINIB - ALUNBRIG</u></b>						
N 208772 003	9012462	Feb 06, 2031	DS		NCE	Apr 28, 2022
	9273077	May 21, 2029		U-1927	ODE-142	Apr 28, 2024
	9611283	Apr 10, 2034		U-1927		
<b><u>BRIMONIDINE TARTRATE - ALPHAGAN P</u></b>						
N 021262 001	6562873	Jul 10, 2021				
	6627210	Jul 18, 2021		DP		
	6641834	Jul 28, 2021		DP		
	6673337	Jul 26, 2021		DP		
	9295641	Jul 10, 2021			U-1833	
	9295641*PED	Jan 10, 2022				
<b><u>BRIMONIDINE TARTRATE - OOLJANA</u></b>						
N 021764 001	7265117	Aug 19, 2025		DP		
<b><u>BRIMONIDINE TARTRATE - ALPHAGAN P</u></b>						
N 021770 001	6562873	Jul 10, 2021		DP		
	6627210	Jul 18, 2021		DP		
	6641834	Jul 28, 2021		DP		
	6673337	Jul 26, 2021		DP		
	8858961	Sep 02, 2023		DP		
	8858961*PED	Mar 02, 2024				
	9295641	Jul 10, 2021			U-1833	
	9295641*PED	Jan 10, 2022				
	9687443	Jul 10, 2021		DP		
<b><u>BRIMONIDINE TARTRATE - MIRVASO</u></b>						
N 204708 001	7439241	Aug 25, 2025		U-1428		
	8053427	Jun 13, 2031		DP U-1428		
	8163725	Jun 13, 2031		DP		
	8231885	May 24, 2025		DP		
	8410102	May 24, 2025			U-1428	
	8426410	May 24, 2025			U-1428	
	8513247	Mar 25, 2031		DP U-1428		
	8513249	Mar 25, 2031		DP U-1428		
	8859551	May 25, 2024			U-1428	
<b><u>BRIMONIDINE TARTRATE - LIJMIFY</u></b>						
N 208144 001					NP	Dec 22, 2020
<b><u>BRIMONIDINE TARTRATE; BRINZOLAMIDE - SIMBRINZA</u></b>						
N 204251 001	6316441	Dec 07, 2019		U-778		
	9044484	Oct 30, 2030		DP		
	9421265	Jun 17, 2030		DP		
<b><u>BRIMONIDINE TARTRATE; TIMOLOL MALEATE - COMBIGAN</u></b>						
N 021398 001	7030149	Apr 19, 2022		U-849		
	7320976	Apr 19, 2022			U-849	
	7323463	Jan 19, 2023		DP		Y
	7642258	Apr 19, 2022	DS	DP U-1024		
	8133890	Apr 19, 2022			U-1235	
	8354409	Apr 19, 2022		DP U-1371		
	8748425	Apr 19, 2022		DP U-1524		
	9474751	Apr 19, 2022		DP U-1524		
	9770453	Apr 19, 2022		DP U-2131		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205836 001	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		
	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205836 002	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205836 002	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205836 003	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		
	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205836 004	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		
	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205836 005	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		
	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205837 001	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		
	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BRIVARACETAM - BRIVIACT</u></b>						
N 205838 001	6784197	Feb 21, 2021	DS DP	U-1815	NCE	May 12, 2021
	6784197	Feb 21, 2021	DS DP	U-2130		
	6911461	Feb 21, 2021	DS DP	U-1815		
	6911461	Feb 21, 2021	DS DP	U-2130		
	8492416	Feb 21, 2021		U-1815		
	8492416	Feb 21, 2021		U-2130		
<b><u>BROMFENAC SODIUM - PROLENSA</u></b>						
N 203168 001	8129431	Sep 11, 2025	DS DP			
	8669290	Jan 16, 2024	DP			
	8754131	Jan 16, 2024	DP			
	8871813	Jan 16, 2024	DP			
	8927606	Jan 16, 2024		U-100		
	8927606	Jan 16, 2024		U-1095		
	8927606	Jan 16, 2024		U-810		
	9144609	Jan 16, 2024	DP			
	9517220	Nov 11, 2033		U-1933		
	9561277	Jan 16, 2024		U-1933		
<b><u>BROMFENAC SODIUM - BROMSITE</u></b>						
N 206911 001	8778999	Sep 03, 2029	DP	U-1834	NP	Apr 08, 2019
<b><u>BROMOCRIPTINE MESYLATE - CYCLOSET</u></b>						
N 020866 001	7888310	Jul 25, 2023		U-1433		
	8137992	Jul 25, 2023		U-1433		
	8137993	Jul 25, 2023		U-1433		
	8137994	Jul 25, 2023		U-1433		
	8431155	Apr 30, 2032	DP	U-976		
	8613947	Apr 30, 2032	DP	U-976		
	8877708	Jun 07, 2030	DP	U-1706		
	9192576	Apr 30, 2032	DP	U-976		
	9352025	Jun 07, 2030		U-2111		
	9352025	Jun 07, 2030		U-2112		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BROMOCRIPTINE MESYLATE - CYCLOSET</u></b>						
N 020866 001	9352025	Jun 07, 2030	U-2113			
	9352025	Jun 07, 2030	U-2114			
	9352025	Jun 07, 2030	U-2115			
	9352025	Jun 07, 2030	U-2116			
	9352025	Jun 07, 2030	U-2117			
	9352025	Jun 07, 2030	U-2118			
	9352025	Jun 07, 2030	U-2119			
	9522117	Apr 30, 2032	DP U-1939			
	9522117	Apr 30, 2032	DP U-976			
	9700555	Apr 30, 2032	DP U-2183			
	9700555	Apr 30, 2032	DP U-2184			
	9700555	Apr 30, 2032	DP U-2185			
	9700555	Apr 30, 2032	DP U-2186			
	9700555	Apr 30, 2032	DP U-2187			
	9700555	Apr 30, 2032	DP U-2188			
	9700555	Apr 30, 2032	DP U-2189			
	9700555	Apr 30, 2032	DP U-2190			
	9700555	Apr 30, 2032	DP U-2191			
	9700555	Apr 30, 2032	DP U-2192			
	9700555	Apr 30, 2032	DP U-2193			
	9700555	Apr 30, 2032	DP U-2194			
	9700555	Apr 30, 2032	DP U-2195			
	9700555	Apr 30, 2032	DP U-2196			
	9700555	Apr 30, 2032	DP U-2197			
	9700555	Apr 30, 2032	DP U-2198			
<b><u>BUDESONIDE - PULMICORT RESPULES</u></b>						
N 020929 001	6598603	Dec 23, 2018	U-529			
	6899099	Dec 23, 2018	U-529			
	7524834	Nov 11, 2018	DP U-966			
<b><u>BUDESONIDE - PULMICORT RESPULES</u></b>						
N 020929 002	6598603	Dec 23, 2018	U-529			
	6899099	Dec 23, 2018	U-529			
	7524834	Nov 11, 2018	DP U-966			
<b><u>BUDESONIDE - PULMICORT RESPULES</u></b>						
N 020929 003	6598603	Dec 23, 2018	U-529			
	6899099	Dec 23, 2018	U-529			
	7524834	Nov 11, 2018	DP U-966			
<b><u>BUDESONIDE - ENTOCORT EC</u></b>						
N 021324 001				M-178	Apr 29, 2019	
				NPP	Apr 29, 2019	
<b><u>BUDESONIDE - PULMICORT FLEXHALER</u></b>						
N 021949 001	6027714	Jan 09, 2018	DP U-787			
	6142145	May 08, 2018	DP			
	6287540	Jan 09, 2018	DP			
	7143764	Mar 13, 2018	DP			
<b><u>BUDESONIDE - PULMICORT FLEXHALER</u></b>						
N 021949 002	6027714	Jan 09, 2018	DP U-787			
	6142145	May 08, 2018	DP			
	6287540	Jan 09, 2018	DP			
	7143764	Mar 13, 2018	DP			
<b><u>BUDESONIDE - UCERIS</u></b>						
N 203634 001	7410651	Jun 09, 2020	DP U-1325			
	7431943	Jun 09, 2020	DP			
	8293273	Jun 09, 2020	DP			
	8784888	Jun 09, 2020	DP			
	8895064	Sep 07, 2031	DP			
	9132093	Sep 07, 2031	DP			
	9192581	Sep 07, 2031	DP U-1325			
	9320716	Jun 09, 2020	DP U-1325			
	9532954	Jun 09, 2020	DP U-1325			
	RE43799	Jun 09, 2020	DP U-1325			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BUDESONIDE - UCERIS</u></b>						
	N 205613 001				NP	Oct 07, 2017
<b><u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u></b>						
N 021929 001	6123924	Sep 26, 2017	DP		M-210	Sep 11, 2020
	7367333	Nov 11, 2018	DP		M-214	Dec 20, 2020
	7367333*PED	May 11, 2019			NPP	Jan 27, 2020
	7587988	Apr 10, 2026	DP		PED	Jul 27, 2020
	7587988*PED	Oct 10, 2026				
	7759328	Jan 29, 2023	DP U-2001			
	7759328	Jan 29, 2023	DP U-2002			
	7759328	Jan 29, 2023	DP U-2122			
	7759328*PED	Jul 29, 2023				
	7967011	Aug 11, 2021	DP			
	7967011*PED	Feb 11, 2022				
	8143239	Jan 29, 2023	DP U-2001			
	8143239	Jan 29, 2023	DP U-2002			
	8143239	Jan 29, 2023	DP U-2122			
	8143239*PED	Jul 29, 2023				
	8387615	Nov 10, 2024	DP			
	8387615*PED	May 10, 2025				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8575137	Jan 29, 2023	DP U-2001			
	8575137	Jan 29, 2023	DP U-2002			
	8575137	Jan 29, 2023	DP U-2122			
	8575137*PED	Jul 29, 2023				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699	Nov 10, 2024	DP			
	8875699*PED	May 10, 2025				
<b><u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u></b>						
N 021929 002	6123924	Sep 26, 2017	DP		M-210	Sep 11, 2020
	7367333	Nov 11, 2018	DP		M-214	Dec 20, 2020
	7367333*PED	May 11, 2019				
	7587988	Apr 10, 2026	DP			
	7587988*PED	Oct 10, 2026				
	7759328	Jan 29, 2023	DP U-2001			
	7759328	Jan 29, 2023	DP U-2002			
	7759328	Jan 29, 2023	DP U-2122			
	7759328*PED	Jul 29, 2023				
	7897646	Sep 09, 2018	U-2002			
	7897646	Sep 09, 2018	U-2122			
	7897646*PED	Mar 09, 2019				
	7967011	Aug 11, 2021	DP			
	7967011*PED	Feb 11, 2022				
	8143239	Jan 29, 2023	DP U-2001			
	8143239	Jan 29, 2023	DP U-2002			
	8143239	Jan 29, 2023	DP U-2122			
	8143239*PED	Jul 29, 2023				
	8387615	Nov 10, 2024	DP			
	8387615*PED	May 10, 2025				
	8461211	Sep 09, 2018	U-2002			
	8461211	Sep 09, 2018	U-2122			
	8461211*PED	Mar 09, 2019				
	8528545	Oct 16, 2028	DP			
	8528545*PED	Apr 16, 2029				
	8575137	Jan 29, 2023	DP U-2001			
	8575137	Jan 29, 2023	DP U-2002			
	8575137	Jan 29, 2023	DP U-2122			
	8575137*PED	Jul 29, 2023				
	8616196	Apr 07, 2029	DP			
	8616196*PED	Oct 07, 2029				
	8875699	Nov 10, 2024	DP			
	8875699*PED	May 10, 2025				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BUPIVACAINE - EXPAREL</u></b>						
N 022496 001	8182835	Sep 18, 2018	DP U-1246			
	8834921	Sep 18, 2018	DP U-1587			
	9585838	Dec 24, 2021	DP			
<b><u>BUPIVACAINE - EXPAREL</u></b>						
N 022496 002	8182835	Sep 18, 2018	DP U-1246			
	8834921	Sep 18, 2018	DP U-1587			
	9205052	Sep 18, 2018	U-1246			
	9585838	Dec 24, 2021	DP			
<b><u>BUPRENORPHINE - BUTRANS</u></b>						
N 021306 001	9642850	Sep 29, 2017	U-1556			
	RE41408	Sep 29, 2017	U-1072			
	RE41408	Sep 29, 2017	U-1556			
	RE41489	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1556			
	RE41571	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1556			
<b><u>BUPRENORPHINE - BUTRANS</u></b>						
N 021306 002	9642850	Sep 29, 2017	U-1556			
	RE41408	Sep 29, 2017	U-1072			
	RE41408	Sep 29, 2017	U-1556			
	RE41489	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1556			
	RE41571	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1556			
<b><u>BUPRENORPHINE - BUTRANS</u></b>						
N 021306 003	9642850	Sep 29, 2017	U-1556			
	RE41408	Sep 29, 2017	U-1072			
	RE41408	Sep 29, 2017	U-1556			
	RE41489	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1556			
	RE41571	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1556			
<b><u>BUPRENORPHINE - BUTRANS</u></b>						
N 021306 004	9642850	Sep 29, 2017	U-1556			
	RE41408	Sep 29, 2017	U-1072			
	RE41408	Sep 29, 2017	U-1556			
	RE41489	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1556			
	RE41571	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1556			
<b><u>BUPRENORPHINE - BUTRANS</u></b>						
N 021306 005	9642850	Sep 29, 2017	U-1556			
	RE41408	Sep 29, 2017	U-1072			
	RE41408	Sep 29, 2017	U-1556			
	RE41489	Sep 29, 2017	U-1072			
	RE41489	Sep 29, 2017	U-1556			
	RE41571	Sep 29, 2017	U-1072			
	RE41571	Sep 29, 2017	U-1556			
<b><u>BUPRENORPHINE - SUBLOCADÉ</u></b>						
N 209819 001	8921387	Jan 06, 2032	DP U-2173			
	8921387	Jan 06, 2032	DP U-2174			
	8975270	Sep 05, 2031	DP U-2175			
	8975270	Sep 05, 2031	DP U-2206			
	9272044	Jun 06, 2031	U-2176			
	9272044	Jun 06, 2031	U-2177			
	9272044	Jun 06, 2031	U-2178			
	9272044	Jun 06, 2031	U-2209			
	9498432	Jun 06, 2031	DP U-2179			
	9782402	Jun 06, 2031	DP U-2176			
	9782402	Jun 06, 2031	DP U-2180			
	9782402	Jun 06, 2031	DP U-2207			
	9782402	Jun 06, 2031	DP U-2208			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BUPRENORPHINE - SUBLOCADÉ</u></b>						
N 209819 001	9827241	Jun 06, 2031	DP U-2174			
	9827241	Jun 06, 2031	DP U-2181			
	9827241	Jun 06, 2031	DP U-2206			
	9827241	Jun 06, 2031	DP U-2210			
	9827241	Jun 06, 2031	DP U-2211			
<b><u>BUPRENORPHINE - SUBLOCADÉ</u></b>						
N 209819 002	8921387	Jan 06, 2032	DP U-2173			
	8921387	Jan 06, 2032	DP U-2174			
	8975270	Sep 05, 2031	DP U-2175			
	8975270	Sep 05, 2031	DP U-2206			
	9272044	Jun 06, 2031	U-2176			
	9272044	Jun 06, 2031	U-2177			
	9272044	Jun 06, 2031	U-2178			
	9272044	Jun 06, 2031	U-2209			
	9498432	Jun 06, 2031	DP U-2179			
	9782402	Jun 06, 2031	DP U-2176			
	9782402	Jun 06, 2031	DP U-2180			
	9782402	Jun 06, 2031	DP U-2207			
	9782402	Jun 06, 2031	DP U-2208			
	9827241	Jun 06, 2031	DP U-2174			
	9827241	Jun 06, 2031	DP U-2181			
	9827241	Jun 06, 2031	DP U-2206			
	9827241	Jun 06, 2031	DP U-2210			
	9827241	Jun 06, 2031	DP U-2211			
<b><u>BUPRENORPHINE HYDROCHLORIDE - PROBUPHINE</u></b>						
N 204442 001	7736665	Apr 25, 2024	U-1878		NP	May 26, 2019
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 001	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 002	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 003	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 004	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 005	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 006	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE - BELBUCA</u></b>						
N 207932 007	7579019	Jan 22, 2020	U-1769		NP	Oct 23, 2018
	8147866	Jul 23, 2027	DP U-1769			
	9655843	Jul 23, 2027	DP U-1556			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u></b>						
N 022410 001	8017150	Feb 13, 2023	DP			
	8475832	Mar 26, 2030	DP U-1411			
	8603514	Apr 03, 2024	DP U-1464			
	9687454	Aug 07, 2029	DP U-1464			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u></b>						
N 022410 001	9855221	Feb 14, 2022	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u></b>						
N 022410 002	8017150	Feb 13, 2023	DP			
	8475832	Mar 26, 2030	DP U-1411			
	8603514	Apr 03, 2024	DP U-1464			
	9687454	Aug 07, 2029	DP U-1464			
	9855221	Feb 14, 2022	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u></b>						
N 022410 003	8017150	Feb 13, 2023	DP			
	8475832	Mar 26, 2030	DP U-1411			
	8603514	Apr 03, 2024	DP U-1464			
	9687454	Aug 07, 2029	DP U-1464			
	9855221	Feb 14, 2022	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - SUBOXONE</u></b>						
N 022410 004	8017150	Feb 13, 2023	DP			
	8475832	Mar 26, 2030	DP U-1411			
	8603514	Apr 03, 2024	DP U-1464			
	9687454	Aug 07, 2029	DP U-1464			
	9855221	Feb 14, 2022	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u></b>						
N 204242 001	8454996	Sep 24, 2019	U-1421	I-713	Aug 10, 2018	
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u></b>						
N 204242 002	8454996	Sep 24, 2019	U-1421	I-713	Aug 10, 2018	
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u></b>						
N 204242 003	8454996	Sep 24, 2019	U-1421	I-713	Aug 10, 2018	
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u></b>						
N 204242 004	8454996	Sep 24, 2019	U-1421	I-713	Aug 10, 2018	
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u></b>						
N 204242 005	8454996	Sep 24, 2019	U-1421	I-713	Aug 10, 2018	
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			
	9439900	Sep 18, 2032	DP			
<b><u>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</u></b>						
N 204242 006	8454996	Sep 24, 2019	U-1421	I-713	Aug 10, 2018	
	8470361	May 22, 2030	DP U-1425			
	8658198	Dec 03, 2027	DP U-1494			
	8940330	Sep 18, 2032	DP			
	9259421	Sep 18, 2032	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - ZUBSOLV</b>						
N 204242 006	9439900	Sep 18, 2032	DP		Y	
<b>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</b>						
N 205637 001	7579019	Jan 22, 2020	U-1521			
	8147866	Jul 23, 2027	DP U-1521			
	8703177	Aug 20, 2032	DP			
	9522188	Apr 24, 2035	DP			
	9655843	Jul 23, 2027	DP U-2017			
<b>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</b>						
N 205637 002	7579019	Jan 22, 2020	U-1521			
	8147866	Jul 23, 2027	DP U-1521			
	8703177	Aug 20, 2032	DP			
	9522188	Apr 24, 2035	DP			
	9655843	Jul 23, 2027	DP U-2017			
<b>BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE - BUNAVAIL</b>						
N 205637 003	7579019	Jan 22, 2020	U-1521			
	8147866	Jul 23, 2027	DP U-1521			
	8703177	Aug 20, 2032	DP			
	9522188	Apr 24, 2035	DP			
	9655843	Jul 23, 2027	DP U-2017			
<b>BUPROPION HYDROBROMIDE - APLENZIN</b>						
N 022108 001	7241805	Jun 27, 2026	DP			
	7569610	Jun 27, 2026	U-997			
	7572935	Jun 27, 2026	DP			
	7585897	Jun 27, 2026	DP			
	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	7662407	Jun 27, 2026	DP			
	7671094	Jun 27, 2026	DP			
<b>BUPROPION HYDROBROMIDE - APLENZIN</b>						
N 022108 002	7241805	Jun 27, 2026	DP			
	7569610	Jun 27, 2026	U-997			
	7572935	Jun 27, 2026	DP			
	7585897	Jun 27, 2026	DP			
	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	7662407	Jun 27, 2026	DP			
	7671094	Jun 27, 2026	DP			
<b>BUPROPION HYDROBROMIDE - APLENZIN</b>						
N 022108 003	7241805	Jun 27, 2026	DP			
	7569610	Jun 27, 2026	U-997			
	7572935	Jun 27, 2026	DP			
	7585897	Jun 27, 2026	DP			
	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	7662407	Jun 27, 2026	DP			
	7671094	Jun 27, 2026	DP			
<b>BUPROPION HYDROCHLORIDE - WELLBUTRIN XL</b>						
N 021515 001	6096341	Oct 30, 2018				
<b>BUPROPION HYDROCHLORIDE - WELLBUTRIN XL</b>						
N 021515 002	6096341	Oct 30, 2018				
<b>BUPROPION HYDROCHLORIDE - FORFIVO XL</b>						
N 022497 001	7674479	Jun 25, 2027	DP			
<b>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</b>						
N 200063 001	7375111	Mar 26, 2025	DP		NC	Sep 10, 2017
	7462626	Jul 20, 2024	U-1583			
	8088786	Feb 03, 2029	DP			
	8318788	Nov 08, 2027	U-1584			
	8722085	Nov 08, 2027	U-1585			
	8815889	Jul 20, 2024	U-1586			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>BUPROPION HYDROCHLORIDE; NALTREXONE HYDROCHLORIDE - CONTRAVE</u></b>						
N 200063 001	8916195	Feb 02, 2030	U-1639			
	9107837	Jun 04, 2027	U-1639			
	9125868	Nov 08, 2027	U-1585			
	9248123	Jan 13, 2032	U-1808			
<b><u>BUTOCONAZOLE NITRATE - BUTOCONAZOLE NITRATE</u></b>						
N 019881 001	5993856	Nov 17, 2017	DP U-457			
<b><u>CABAZITAXEL - JEVTANA KIT</u></b>						
N 201023 001	5847170	Mar 26, 2021	DS DP		M-201	May 17, 2020
	5847170*PED	Sep 26, 2021			M-209	Sep 14, 2020
	7241907	Dec 10, 2025	DS		PED	Nov 17, 2020
	7241907*PED	Jun 10, 2026				
	8927592	Oct 27, 2030	U-1630			
	8927592*PED	Apr 27, 2031				
<b><u>CABOZANTINIB S-MALATE - COMETRIO</u></b>						
N 203756 001	7579473	Aug 14, 2026	DS DP		NCE	Nov 29, 2017
	8877776	Oct 08, 2030	DS DP U-1617		ODE-33	Nov 29, 2019
	9717720	Feb 10, 2032	DP			
<b><u>CABOZANTINIB S-MALATE - COMETRIO</u></b>						
N 203756 002	7579473	Aug 14, 2026	DS DP		NCE	Nov 29, 2017
	8877776	Oct 08, 2030	DS DP U-1617		ODE-33	Nov 29, 2019
	9717720	Feb 10, 2032	DP			
<b><u>CABOZANTINIB S-MALATE - CABOMETYX</u></b>						
N 208692 001	7579473	Aug 14, 2026	DS DP		I-760	Dec 19, 2020
	8497284	Sep 24, 2024	U-1220		NCE	Nov 29, 2017
	8877776	Oct 08, 2030	DS DP		NP	Apr 25, 2019
	9724342	Jul 09, 2033	DP			
<b><u>CABOZANTINIB S-MALATE - CABOMETYX</u></b>						
N 208692 002	7579473	Aug 14, 2026	DS DP		I-760	Dec 19, 2020
	8497284	Sep 24, 2024	U-1220		NCE	Nov 29, 2017
	8877776	Oct 08, 2030	DS DP		NP	Apr 25, 2019
	9724342	Jul 09, 2033	DP			
<b><u>CABOZANTINIB S-MALATE - CABOMETYX</u></b>						
N 208692 003	7579473	Aug 14, 2026	DS DP		I-760	Dec 19, 2020
	8497284	Sep 24, 2024	U-1220		NCE	Nov 29, 2017
	8877776	Oct 08, 2030	DS DP		NP	Apr 25, 2019
	9724342	Jul 09, 2033	DP			
<b><u>CALCIFEDIOL - RAYALDEE</u></b>						
N 208010 001	6582727	Aug 22, 2020	DP		NP	Jun 17, 2019
	8207149	Apr 25, 2028	U-1871			
	8361488	Jul 19, 2028	DP			
	8426391	Aug 27, 2028	DP U-1872			
	8778373	Apr 25, 2028	U-1873			
	8906410	Feb 02, 2027	DP			
	9408858	Apr 25, 2028	U-1888			
	9498486	Apr 25, 2028	U-1920			
<b><u>CALCIPOTRIENE - SORILUX</u></b>						
N 022563 001	8263580	Sep 27, 2028	DP U-1280			
	8629128	May 26, 2026	DP U-1280			
	8629128	May 26, 2026	DP U-1767			
<b><u>CALCITONIN SALMON RECOMBINANT - FORTICAL</u></b>						
N 021406 001	6440392	Feb 02, 2021	DP U-227			
	RE40812	Feb 02, 2021	DP			
	RE43580	Feb 02, 2021	DP U-227			
<b><u>CALCITRIOL - CALCIJEX</u></b>						
N 018874 001	6051567	Aug 02, 2019				
	6265392	Aug 02, 2019				
	6274169	Aug 02, 2019				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>CALCITRIOL - CALCIJEX</u></b>						
N 018874 002	6051567	Aug 02, 2019				
	6265392	Aug 02, 2019				
	6274169	Aug 02, 2019				
<b><u>CALCIUM ACETATE - PHOSLO</u></b>						
N 021160 002	6576665	Apr 03, 2021				
<b><u>CALCIUM ACETATE - PHOSLO GELCAPS</u></b>						
N 021160 003	6576665	Apr 03, 2021				
	6875445	Jul 30, 2021	DP			
<b><u>CALCIUM ACETATE - PHOSLYRA</u></b>						
N 022581 001	8591938	Feb 23, 2030	DP U-1469			
	8592480	Jul 20, 2027	U-1469			
	9089528	Jul 20, 2027	U-1469			
<b><u>CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE - PEPCID COMPLETE</u></b>						
N 020958 001	5989588	Sep 30, 2017	DP U-349			
	5989588*PED	Mar 30, 2018				
	6814978	Aug 26, 2021	DP			
<b><u>CALCIUM CARBONATE; RISEDRONATE SODIUM - ACTONEL WITH CALCIUM (COPACKAGED)</u></b>						
N 021823 001	5994329	Jul 17, 2018	U-353			
	6015801	Jul 17, 2018	U-353			
	6165513	Jun 10, 2018	DP			
	6432932	Jul 17, 2018	U-595			
	6465443	Aug 14, 2018	DP			
<b><u>CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; OXIGLUTATIONE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - NAVSTEL</u></b>						
N 022193 001	7084130	Nov 29, 2021	DP U-891			
<b><u>CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - PHOXILLUM BK 4/2.5 IN PLASTIC CONTAINER</u></b>						
N 207026 001				ODE-85	Jan 13, 2022	
<b><u>CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE - PHOXILLUM B22K 4/0 IN PLASTIC CONTAINER</u></b>						
N 207026 002				ODE-85	Jan 13, 2022	
<b><u>CANAGLIFLOZIN - INVOKANA</u></b>						
N 204042 001	7943582	Feb 26, 2029	DS DP U-493	I-733	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	M-197	Feb 01, 2020	
	8222219	Jul 30, 2024	U-493	NCE	Mar 29, 2018	
	8513202	Dec 03, 2027	DS DP U-493			
<b><u>CANAGLIFLOZIN - INVOKANA</u></b>						
N 204042 002	7943582	Feb 26, 2029	DS DP U-493	I-733	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	M-197	Feb 01, 2020	
	8222219	Jul 30, 2024	U-493	NCE	Mar 29, 2018	
	8513202	Dec 03, 2027	DS DP U-493			
<b><u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u></b>						
N 204353 001	7943582	Feb 26, 2029	DS DP U-493	I-735	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	M-197	Feb 01, 2020	
	8222219	Jul 30, 2024	U-493	NC	Aug 08, 2017	
	8513202	Dec 03, 2027	DS DP U-493	NCE	Mar 29, 2018	
	8785403	Jul 30, 2024	DP			
<b><u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u></b>						
N 204353 002	7943582	Feb 26, 2029	DS DP U-493	I-735	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	M-197	Feb 01, 2020	
	8222219	Jul 30, 2024	U-493	NC	Aug 08, 2017	
	8513202	Dec 03, 2027	DS DP U-493	NCE	Mar 29, 2018	
	8785403	Jul 30, 2024	DP			
<b><u>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</u></b>						
N 204353 003	7943582	Feb 26, 2029	DS DP U-493	I-735	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	M-197	Feb 01, 2020	
	8222219	Jul 30, 2024	U-493	NC	Aug 08, 2017	

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</b>						
N 204353 003	8513202	Dec 03, 2027	DS DP U-493		NCE	Mar 29, 2018
	8785403	Jul 30, 2024	DP			
<b>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET</b>						
N 204353 004	7943582	Feb 26, 2029	DS DP U-493	I-735	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	M-197	Feb 01, 2020	
	8222219	Jul 30, 2024	U-493	NC	Aug 08, 2017	
	8513202	Dec 03, 2027	DS DP U-493	NCE	Mar 29, 2018	
	8785403	Jul 30, 2024	DP			
<b>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</b>						
N 205879 001	6723340	Oct 25, 2021	DP	I-735	May 20, 2019	
	7943582	Feb 26, 2029	DS DP U-493	NC	Aug 08, 2017	
	7943788	Jul 14, 2027	DS DP	NCE	Mar 29, 2018	
	8222219	Jul 30, 2024	U-493			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<b>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</b>						
N 205879 002	7943582	Feb 26, 2029	DS DP U-493	I-735	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	NC	Aug 08, 2017	
	8222219	Jul 30, 2024	U-493	NCE	Mar 29, 2018	
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<b>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</b>						
N 205879 003	6723340	Oct 25, 2021	DP	I-735	May 20, 2019	
	7943582	Feb 26, 2029	DS DP U-493	NC	Aug 08, 2017	
	7943788	Jul 14, 2027	DS DP	NCE	Mar 29, 2018	
	8222219	Jul 30, 2024	U-493			
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<b>CANAGLIFLOZIN; METFORMIN HYDROCHLORIDE - INVOKAMET XR</b>						
N 205879 004	7943582	Feb 26, 2029	DS DP U-493	I-735	May 20, 2019	
	7943788	Jul 14, 2027	DS DP	NC	Aug 08, 2017	
	8222219	Jul 30, 2024	U-493	NCE	Mar 29, 2018	
	8513202	Dec 03, 2027	DS DP U-493			
	8785403	Jul 30, 2024	DP			
<b>CANGRELOR - KENGREAL</b>						
N 204958 001	6114313	Dec 11, 2017	DP U-1715	NCE	Jun 22, 2020	
	6130208	Jun 29, 2018	DP U-1715			
	8680052	Mar 09, 2033	U-1715			
	8759316	May 13, 2029	U-1715			
	9295687	Jul 10, 2035	DP			
	9427448	Nov 10, 2030	U-1926			
	9439921	Jul 10, 2035	DP			
	9700575	Jul 10, 2035	DP			
<b>CAPSAICIN - QUTENZA</b>						
N 022395 001	6239180	Jun 04, 2021	DP			
<b>CARBAMAZEPINE - EQUETRO</b>						
N 021710 001	6977253	May 19, 2024	U-693			
<b>CARBAMAZEPINE - EQUETRO</b>						
N 021710 002	6977253	May 19, 2024	U-693			
<b>CARBAMAZEPINE - EQUETRO</b>						
N 021710 003	6977253	May 19, 2024	U-693			
<b>CARBAMAZEPINE - CARNEXIV</b>						
N 206030 001	7635773	Mar 13, 2029	DP	ODE-124	Oct 07, 2023	
	8410077	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9629797	Nov 10, 2028	U-2004			
	9629797	Nov 10, 2028	U-2005			
	9629797	Nov 10, 2028	U-2006			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>CARBAMAZEPINE - CARNEXIV</u></b>						
N 206030 001	9750822	Mar 13, 2029	DP			
	9770407	Nov 10, 2028	DP			
<b><u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 50</u></b>						
N 021485 001	6500867	Jun 29, 2020	DP U-219			
	6797732	Jun 29, 2020	DP			
<b><u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100</u></b>						
N 021485 002	6500867	Jun 29, 2020	DP U-219			
	6797732	Jun 29, 2020	DP			
<b><u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150</u></b>						
N 021485 003	6500867	Jun 29, 2020	DP U-219			
	6797732	Jun 29, 2020	DP			
<b><u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 200</u></b>						
N 021485 004	6500867	Jun 29, 2020	DP U-219			
	6797732	Jun 29, 2020	DP			
<b><u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 75</u></b>						
N 021485 005	6500867	Jun 29, 2020	DP U-219			
	6797732	Jun 29, 2020	DP			
<b><u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 125</u></b>						
N 021485 006	6500867	Jun 29, 2020	DP U-219			
	6797732	Jun 29, 2020	DP			
<b><u>CARBIDOPA; LEVODOPA - RYTARY</u></b>						
N 203312 001	7094427	May 29, 2022	DP U-1645		NDF	Jan 07, 2018
	8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
<b><u>CARBIDOPA; LEVODOPA - RYTARY</u></b>						
N 203312 002	7094427	May 29, 2022	DP U-1645		NDF	Jan 07, 2018
	8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
<b><u>CARBIDOPA; LEVODOPA - RYTARY</u></b>						
N 203312 003	7094427	May 29, 2022	DP U-1645		NDF	Jan 07, 2018
	8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>CARBIDOPA; LEVODOPA - RYTARY</u></b>						
N 203312 003	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
<b><u>CARBIDOPA; LEVODOPA - RYTARY</u></b>						
N 203312 004	7094427	May 29, 2022	DP U-1645		NDF	Jan 07, 2018
	8377474	Dec 26, 2028	DP U-1645			
	8377474	Dec 26, 2028	DP U-219			
	8454998	Dec 26, 2028	DP U-1645			
	8454998	Dec 26, 2028	DP U-1646			
	8454998	Dec 26, 2028	DP U-1647			
	8454998	Dec 26, 2028	DP U-1649			
	8454998	Dec 26, 2028	DP U-219			
	8557283	Dec 26, 2028	DP U-1645			
	8557283	Dec 26, 2028	DP U-219			
	9089607	Dec 26, 2028	DP U-1645			
	9089607	Dec 26, 2028	DP U-1720			
	9089608	Dec 26, 2028	DP			
	9463246	Dec 26, 2028	DP U-219			
	9533046	Dec 26, 2028	DP U-219			
<b><u>CARBIDOPA; LEVODOPA - DUOPA</u></b>						
N 203952 001					NP	Jan 09, 2018
					ODE-84	Jan 09, 2022
<b><u>CARBINOXAMINE MALEATE - KARBINAL ER</u></b>						
N 022556 001	8062667	Mar 29, 2029	DP			
	9522191	Jun 15, 2027	DP			
<b><u>CARFILZOMIB - KYPROLIS</u></b>						
N 202714 001	7232818	Apr 14, 2025	DS DP		I-712	Jul 24, 2018
	7417042	Jul 20, 2026	DS DP		I-722	Jan 21, 2019
	7491704	Apr 14, 2025		U-1260	I-723	Jan 21, 2019
	7737112	Dec 07, 2027	DP		NCE	Jul 20, 2017
	8129346	Apr 14, 2025		U-1260	ODE-27	Jul 20, 2019
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-1260		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	9511109	Oct 21, 2029		U-1924		
<b><u>CARFILZOMIB - KYPROLIS</u></b>						
N 202714 002	7232818	Apr 14, 2025	DS DP		I-712	Jul 24, 2018
	7417042	Jul 20, 2026	DS DP		I-722	Jan 21, 2019
	7491704	Apr 14, 2025		U-1260	I-723	Jan 21, 2019
	7737112	Dec 07, 2027	DP		NCE	Jul 20, 2017
	8129346	Apr 14, 2025		U-1260	ODE-27	Jul 20, 2019
	8207125	Apr 14, 2025	DS DP			
	8207126	Apr 14, 2025	DP			
	8207127	Apr 14, 2025		U-1260		
	8207297	Apr 14, 2025	DS DP			
	9493582	Feb 27, 2033	DP			
	9511109	Oct 21, 2029		U-1924		
<b><u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u></b>						
N 204370 001	7737142	Mar 27, 2027	DS DP U-1750		M-213	Nov 09, 2020
	7943621	Dec 16, 2028	DS DP		NCE	Sep 17, 2020
<b><u>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</u></b>						
N 204370 002	7737142	Mar 27, 2027	DS DP U-1750		M-213	Nov 09, 2020
	7943621	Dec 16, 2028	DS DP		NCE	Sep 17, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</b>						
N 204370 003	7737142	Mar 27, 2027	DS DP U-1750		M-213	Nov 09, 2020
	7943621	Dec 16, 2028	DS DP		NCE	Sep 17, 2020
<b>CARIPRAZINE HYDROCHLORIDE - VRAYLAR</b>						
N 204370 004	7737142	Mar 27, 2027	DS DP U-1750		M-213	Nov 09, 2020
	7943621	Dec 16, 2028	DS DP		NCE	Sep 17, 2020
<b>CARVEDILOL PHOSPHATE - CARVEDILOL PHOSPHATE</b>						
A 090132 001					PC	May 07, 2018
<b>CARVEDILOL PHOSPHATE - CARVEDILOL PHOSPHATE</b>						
A 090132 002					PC	May 07, 2018
<b>CARVEDILOL PHOSPHATE - CARVEDILOL PHOSPHATE</b>						
A 090132 003					PC	May 07, 2018
<b>CARVEDILOL PHOSPHATE - CARVEDILOL PHOSPHATE</b>						
A 090132 004					PC	May 07, 2018
<b>CARVEDILOL PHOSPHATE - COREG CR</b>						
N 022012 001	7268156	Jun 27, 2023	DS DP U-3			
	7268156	Jun 27, 2023	DS DP U-313			
	8101209	Sep 11, 2025	DP			
<b>CARVEDILOL PHOSPHATE - COREG CR</b>						
N 022012 002	7268156	Jun 27, 2023	DS DP U-3			
	7268156	Jun 27, 2023	DS DP U-313			
	8101209	Sep 11, 2025	DP			
<b>CARVEDILOL PHOSPHATE - COREG CR</b>						
N 022012 003	7268156	Jun 27, 2023	DS DP U-3			
	7268156	Jun 27, 2023	DS DP U-313			
	8101209	Sep 11, 2025	DP			
<b>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</b>						
N 206110 001	9636407	Dec 21, 2032	DP			
<b>CASPOFUNGIN ACETATE - CASPOFUNGIN ACETATE</b>						
N 206110 002	9636407	Dec 21, 2032	DP			
<b>CEFIXIME - SUPRAX</b>						
N 202091 001	9233112	Dec 14, 2028	DP U-1676			
<b>CEFTAROLINE FOSAMIL - TEFLARO</b>						
N 200327 001	6417175	Apr 11, 2022	DS DP U-1676		NPP	May 27, 2019
	6906055	Dec 15, 2021	DS DP		NPP	May 27, 2019
	7419973	Dec 15, 2021	DP			
	8247400	Feb 10, 2031	DP U-282			
	9629861	Sep 21, 2030	DP			
<b>CEFTAROLINE FOSAMIL - TEFLARO</b>						
N 200327 002	6417175	Apr 11, 2022	DS DP U-1676		NPP	May 27, 2019
	6906055	Dec 15, 2021	DS DP		NPP	May 27, 2019
	7419973	Dec 15, 2021	DP			
	8247400	Feb 10, 2031	DP U-282			
	9629861	Sep 21, 2030	DP			
<b>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</b>						
N 206829 001	7129232	Oct 21, 2024	DS DP U-36		NCE	Dec 19, 2019
	8476425	Sep 27, 2032	DS		GAIN	Dec 19, 2024
	8685957	Sep 27, 2032	DS U-36			
	8906898	May 28, 2034	DS DP			
	8968753	Mar 14, 2034	U-1672			
	8968753	Mar 14, 2034	U-1673			
	9320740	Mar 14, 2034	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>CEFTOLOZANE SULFATE; TAZOBACTAM SODIUM - ZERBAXA</u></b>						
N 206829 001	7129232	Oct 21, 2024	DS DP U-36		NCE	Dec 19, 2019
	8476425	Sep 27, 2032	DS		GAIN	Dec 19, 2024
	8685957	Sep 27, 2032	DS U-36			
	8906898	May 28, 2034	DS DP			
	8968753	Mar 14, 2034		U-1672		
	8968753	Mar 14, 2034		U-1673		
	9320740	Mar 14, 2034	DP			
<b><u>CERITINIB - ZYKADIA</u></b>						
N 205755 001	7153964	Feb 26, 2021	DS DP		M-199	May 26, 2020
	7893074	Apr 25, 2026	DS DP		NCE	Apr 29, 2019
	7964592	Jan 13, 2027	DS DP		ODE-145	May 26, 2024
	8039479	Jun 29, 2030	DS DP		ODE-66	Apr 29, 2021
	8188276	Jan 31, 2023	DS DP			
	8377921	Nov 20, 2027		U-1179		
	8399450	Nov 20, 2027	DS DP			
	8703787	Feb 02, 2032		U-1179		
	8835430	Jan 31, 2023	DS DP			
	9018204	Jan 31, 2023	DS DP			
	9309229	Jan 18, 2032	DS DP			
	9416112	Jan 31, 2023	DS DP			
<b><u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC ALLERGY</u></b>						
N 021621 003	6455533	Jul 02, 2018	DP U-295			
<b><u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC ALLERGY</u></b>						
N 021621 004	6455533	Jul 02, 2018	DP U-295			
<b><u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC HIVES RELIEF</u></b>						
N 021621 005	6455533	Jul 02, 2018	DP U-295			
<b><u>CETIRIZINE HYDROCHLORIDE - CHILDREN'S ZYRTEC HIVES RELIEF</u></b>						
N 021621 006	6455533	Jul 02, 2018	DP U-295			
<b><u>CETIRIZINE HYDROCHLORIDE - ZERVIATE</u></b>						
N 208694 001	8829005	Mar 15, 2030		U-1680		
	8829005*PED	Sep 15, 2030			NDF	May 30, 2020
	9254286	Jul 09, 2032	DP		PED	Nov 30, 2020
	9254286*PED	Jan 09, 2033				
	9750684	Mar 15, 2030	DP			
<b><u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u></b>						
N 021150 002	6469009	Jul 13, 2019	DP U-295			
	7014867	Jun 10, 2022	DP			
	7226614	Jun 10, 2022	U-295			
<b><u>CETRORELIx - CETROTIDE</u></b>						
N 021197 001	6319192	Apr 23, 2019		U-426		
<b><u>CETRORELIx - CETROTIDE</u></b>						
N 021197 002	6319192	Apr 23, 2019		U-426		
<b><u>CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE</u></b>						
N 021669 001	7066916	Feb 17, 2024		U-737		
	7427574	Apr 25, 2026	DP			
	7595021	May 12, 2023	DP U-1022			
	7717889	Feb 27, 2025	DP U-1022			
	7935093	Oct 02, 2027	DP U-1022			
<b><u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u></b>						
N 020832 001	6536975	Nov 10, 2020	DP			
<b><u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u></b>						
N 020832 002	6729786	Mar 14, 2023	DP			
	6991394	Jan 31, 2024	DP			
	7182536	Dec 30, 2023	DP			
	7241065	Mar 14, 2023	DP			
	7422388	Apr 25, 2027	DP U-1397			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
N 020832 004	6536975	Nov 10, 2020		DP		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832 005	6536975	Nov 10, 2020		DP		
	6729786	Mar 14, 2023		DP		
	7241065	Mar 14, 2023		DP		
	7422388	Apr 25, 2027		DP U-1397		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>						
N 020832 006	6991394	Jan 31, 2024		DP		
	7182536	Dec 30, 2023		DP		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>						
N 020832 007	6536975	Nov 10, 2020		DP		
	6729786	Mar 14, 2023		DP		
	7241065	Mar 14, 2023		DP		
	7422388	Apr 25, 2027		DP U-1397		
<u>CHLOROPROCAINE HYDROCHLORIDE - CLOROTEKAL</u>						
N 208791 001					NP	Sep 26, 2020
<u>CHLORPHENIRAMINE MALEATE; CODEINE PHOSPHATE - CODEINE PHOSPHATE AND CHLORPHENIRAMINE MALEATE</u>						
N 206323 001	6248363	Nov 23, 2019		DP U-1716		
	6383471	Apr 06, 2019		DP U-1716		
	9066942	Jan 03, 2032		U-1716		
	9107921	Jan 03, 2032		DP		
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - ADVIL ALLERGY SINUS</u>						
N 021441 001	7863287	Feb 28, 2027		DP		
<u>CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX - TUZISTRA XR</u>						
N 207768 001	8062667	Mar 29, 2029		DP		
	8790700	Mar 15, 2027		DP		
<u>CHOLIC ACID - CHOLBAM</u>						
N 205750 001					NCE	Mar 17, 2020
					ODE-91	Mar 17, 2022
<u>CHOLIC ACID - CHOLBAM</u>						
N 205750 002					NCE	Mar 17, 2020
					ODE-91	Mar 17, 2022
<u>CHOLINE C-11 - CHOLINE C-11</u>						
N 203155 001					NCE	Sep 12, 2017
					W	Sep 12, 2017
<u>CHOLINE FENOFOBRATE - TRILIPIX</u>						
N 022224 001	7259186	Jan 07, 2025		DS		
<u>CHOLINE FENOFOBRATE - TRILIPIX</u>						
N 022224 002	7259186	Jan 07, 2025		DS		
<u>CHORIOGONADOTROPIN ALFA - OVIDREL</u>						
N 021149 002	6706681	Mar 16, 2021		DP		
<u>CICLESONIDE - ALVESCO</u>						
N 021658 002	5482934	Oct 24, 2017		DS DP U-1002		
	6120752	May 13, 2018		DP		
	6264923	May 13, 2018		DP		
	8371292	Feb 01, 2028		U-1355		
<u>CICLESONIDE - ALVESCO</u>						
N 021658 003	5482934	Oct 24, 2017		DS DP U-1002		
	6120752	May 13, 2018		DP		
	6264923	May 13, 2018		DP		
	8371292	Feb 01, 2028		U-1355		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>CICLESONIDE - OMNARIS</b>						
N 022004 001	5482934	Oct 24, 2017	DS DP U-557			
	6767901	Oct 21, 2020	DP			
	6939559	Apr 21, 2019	DP			
	7235247	Apr 21, 2019	DP			
	8371292	Feb 01, 2028	U-1356			
	8383611	Oct 20, 2020	DP			
<b>CICLESONIDE - ZETONNA</b>						
N 202129 001	5482934	Oct 24, 2017	DS DP U-1002			
	6120752	May 13, 2018	DP			
	6264923	May 13, 2018	DP			
	8371292	Feb 01, 2028	U-1357			
<b>CICLOPIROX - CICLOPIROX</b>						
N 020519 001	7018656	Sep 05, 2018	DP			
<b>CICLOPIROX - LOPROX</b>						
N 021159 001	7981909	Sep 16, 2017	U-1162			
	8227490	Sep 16, 2017	U-1256			
<b>CINACALCET HYDROCHLORIDE - SENSI PAR</b>						
N 021688 001	6011068	Mar 08, 2018	DS DP		M-200	May 23, 2020
	7829595	Sep 22, 2026	DP U-1098		ODE-78	Nov 21, 2021
	9375405	Sep 22, 2026	DP		ODE-8	Feb 25, 2018
<b>CINACALCET HYDROCHLORIDE - SENSI PAR</b>						
N 021688 002	6011068	Mar 08, 2018	DS DP		M-200	May 23, 2020
	7829595	Sep 22, 2026	DP U-1098		ODE-78	Nov 21, 2021
	9375405	Sep 22, 2026	DP		ODE-8	Feb 25, 2018
<b>CINACALCET HYDROCHLORIDE - SENSI PAR</b>						
N 021688 003	6011068	Mar 08, 2018	DS DP		M-200	May 23, 2020
	7829595	Sep 22, 2026	DP U-1098		ODE-78	Nov 21, 2021
	9375405	Sep 22, 2026	DP		ODE-8	Feb 25, 2018
<b>CIPROFLOXACIN - OTIPRIO</b>						
N 207986 001	8318817	Apr 27, 2030	U-1792		NP	Dec 10, 2018
	9205048	Apr 21, 2029	U-1793			
	9220796	Jul 01, 2035	DP			
	9233068	Dec 11, 2029	DP			
<b>CIPROFLOXACIN HYDROCHLORIDE - PROQUIN XR</b>						
N 021744 001	6488962	Jun 20, 2020	DP			
<b>CIPROFLOXACIN HYDROCHLORIDE; FLUOCINOLONE ACETONIDE - OTOVEL</b>						
N 208251 001	8932610	Mar 24, 2030	DP U-1578		NC	Apr 29, 2019
<b>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</b>						
N 021473 001	7709022	Jun 23, 2021	DP			
	8187632	Jun 23, 2021	DP			
	8187632*PED	Dec 23, 2021				
<b>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE - CIPRO XR</b>						
N 021473 002	7709022	Jun 23, 2021	DP			
	8187632	Jun 23, 2021	DP			
	8187632*PED	Dec 23, 2021				
<b>CIPROFLOXACIN; DEXAMETHASONE - CIPRODEX</b>						
N 021537 001	6284804	Aug 10, 2020				
	6359016	Aug 10, 2020				
	8846650	Jun 04, 2025	DP U-1578			
	9149486	Sep 13, 2022	DP U-1578			
	9345714	Sep 13, 2022	DP U-1578			
	9402805	Sep 13, 2022	DP U-1578			
	9402805	Sep 13, 2022	DP U-1679			
<b>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - PREPOPIK</b>						
N 202535 001	8450338	Oct 10, 2028	DP			
	8481083	Oct 10, 2028	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>CITRIC ACID; MAGNESIUM OXIDE; SODIUM PICOSULFATE - CLENPIQ</b>						
N 209589 001	9827231	Jun 23, 2034	DP U-2162			
<b>CLEVIDIPINE - CLEVIPREX</b>						
N 022156 001	5856346	Jan 05, 2021	DS DP U-893			
	8658676	Oct 10, 2031	DP			
<b>CLEVIDIPINE - CLEVIPREX</b>						
N 022156 002	5856346	Jan 05, 2021	DS DP U-893			
	8658676	Oct 10, 2031	DP			
<b>CLEVIDIPINE - CLEVIPREX</b>						
N 022156 003	5856346	Jan 05, 2021	DS DP U-893			
	8658676	Oct 10, 2031	DP			
<b>CLINDAMYCIN PHOSPHATE - CLEOCIN</b>						
N 050767 001	6495157	Jul 20, 2020	DP			
<b>CLINDAMYCIN PHOSPHATE - CLINDAGEL</b>						
N 050782 001	6387383	Aug 03, 2020	DP U-818			
<b>CLINDAMYCIN PHOSPHATE - CLINDESSE</b>						
N 050793 001	5993856	Nov 17, 2017	DP U-137			
	6899890	Apr 27, 2023	DP U-137			
	9789057	Dec 02, 2026	DP U-137			
<b>CLINDAMYCIN PHOSPHATE - EVOCLIN</b>						
N 050801 001	7141237	Jan 23, 2024	DS DP			
	7374747	Aug 09, 2026	DS DP U-921			
<b>CLINDAMYCIN PHOSPHATE; TRETINOIN - ZIANA</b>						
N 050802 001	6387383	Aug 03, 2020	DP U-916			
<b>CLOBAZAM - ONFI</b>						
N 202067 001				ODE-17	Oct 21, 2018	
<b>CLOBAZAM - ONFI</b>						
N 202067 002				ODE-17	Oct 21, 2018	
<b>CLOBAZAM - ONFI</b>						
N 202067 003				ODE-17	Oct 21, 2018	
<b>CLOBAZAM - ONFI</b>						
N 203993 001				ODE-18	Oct 21, 2018	
<b>CLOBETASOL PROPIONATE - CLOBEX</b>						
N 021535 001	6106848	Sep 22, 2017				
<b>CLOBETASOL PROPIONATE - CLOBEX</b>						
N 021644 001	7316810	Jun 17, 2019	DP			
	7700081	Jan 03, 2022	U-1044			
	8066975	Jun 17, 2019	DP			
	8066976	Jun 17, 2019	DP			
<b>CLOBETASOL PROPIONATE - CLOBEX</b>						
N 021835 001	5972920	Feb 12, 2018	DP			
	5990100	Mar 24, 2018	DP U-742			
<b>CLOBETASOL PROPIONATE - OLUX E</b>						
N 022013 001	6730288	Sep 08, 2019	DP			
	7029659	Sep 08, 2019	DP			
	8460641	Nov 05, 2028	DP U-1410			
	8962000	Aug 31, 2025	DP U-1410			
<b>CLOBETASOL PROPIONATE - IMPOYZ</b>						
N 209483 001	9855334	Mar 11, 2035	DP		NP	Nov 28, 2020
<b>CLOFARABINE - CLOFARABINE</b>						
A 204029 001					PC	Nov 05, 2017

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>CLOFARABINE - CLOLAR</b>						
N 021673	001 5661136	Jan 14, 2018		U-626		
<b>CLONIDINE HYDROCHLORIDE - KAPVAY</b>						
N 022331	003				M-149	Nov 20, 2017
<b>CLONIDINE HYDROCHLORIDE - KAPVAY</b>						
N 022331	004				M-149	Nov 20, 2017
<b>CLOPIDOGREL BISULFATE - PLAVIX</b>						
N 020839	001 6429210	Jun 10, 2019	DS DP			
	6504030	Jun 10, 2019	DS			
<b>CLOPIDOGREL BISULFATE - PLAVIX</b>						
N 020839	002 6429210	Jun 10, 2019	DS DP			
	6504030	Jun 10, 2019	DS			
<b>CLOZAPINE - FAZACLO ODT</b>						
N 021590	001 6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<b>CLOZAPINE - FAZACLO ODT</b>						
N 021590	002 6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<b>CLOZAPINE - FAZACLO ODT</b>						
N 021590	003 6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<b>CLOZAPINE - FAZACLO ODT</b>						
N 021590	004 6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<b>CLOZAPINE - FAZACLO ODT</b>						
N 021590	005 6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<b>CLOZAPINE - FAZACLO ODT</b>						
N 021590	006 6024981	Apr 09, 2018	DP			
	6106861	Dec 05, 2017	DP			
	6221392	Apr 09, 2018	DP			
<b>COBICISTAT - TYBOST</b>						
N 203094	001 8148374	Sep 03, 2029	DS DP U-1279		NCE	Aug 27, 2017
					NP	Sep 24, 2017
<b>COBICISTAT; DARUNAVIR ETHANOLATE - PREZCOBIX</b>						
N 205395	001 7470506	Jun 23, 2019		U-1660		
	7470506*PED	Dec 23, 2019				
	7700645	Dec 26, 2026	DS DP			
	7700645*PED	Jun 26, 2027				
	8148374	Sep 03, 2029	DS DP	U-1660		
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1660		
	8597876*PED	Dec 23, 2019				
	RE43596*PED	Nov 09, 2017				
<b>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</b>						
N 207561	001 5914331	Jul 02, 2017	DS		NCE	Aug 27, 2017
	5914331*PED	Jan 02, 2018			NCE	Nov 05, 2020
	6642245	Nov 04, 2020		U-257		
	6642245*PED	May 04, 2021			NPP	Sep 25, 2020
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
	7176220	Aug 27, 2026	DS DP	U-257		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - GENVOYA</u></b>						
N 207561 001	7390791	May 07, 2022	DS DP			
	7635704	Oct 26, 2026	DS DP	U-257		
	7803788	Feb 02, 2022		U-257		
	8148374	Sep 03, 2029	DS DP	U-1279		
	8633219	Apr 24, 2030		DP U-257		
	8754065	Aug 15, 2032	DS DP	U-257		
	8981103	Oct 26, 2026	DS DP			
	9296769	Aug 15, 2032	DS DP	U-257		
<b><u>COBICISTAT; ELVITEGRAVIR; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - STRIBILD</u></b>						
N 203100 001	5914331	Jul 02, 2017	DS		I-704	
	5922695	Jul 25, 2017	DS	U-257	NCE	Dec 17, 2017
	5935946	Jul 25, 2017	DS DP	U-257	NPP	Aug 27, 2017
	5977089	Jul 25, 2017	DS DP	U-257		Jan 27, 2020
	6043230	Jul 25, 2017		U-257		
	6642245	Nov 04, 2020		U-257		
	6703396	Mar 09, 2021	DS DP			
	7176220	Aug 27, 2026	DS DP	U-257		
	7635704	Oct 26, 2026	DS DP	U-257		
	8148374	Sep 03, 2029	DS DP	U-1279		
	8592397	Jan 13, 2024		DP U-257		
	8633219	Apr 24, 2030		DP U-257		
	8716264	Jan 13, 2024		DP U-257		
	8981103	Oct 26, 2026	DS DP			
	9457036	Jan 13, 2024		DP U-257		
	9744181	Jan 13, 2024		DP U-257		
<b><u>COBIMETINIB FUMARATE - COTELLIC</u></b>						
N 206192 001	7803839	Feb 01, 2027	DS DP		NCE	
	8362002	Oct 05, 2026		U-1776	ODE-101	Nov 10, 2020
<b><u>COLCHICINE - COLCRYS</u></b>						
N 022352 001	7601758	Feb 10, 2029		U-1007		
	7619004	Dec 03, 2028		U-1020		
	7820681	Feb 17, 2029		U-1020		
	7906519	Feb 17, 2029		U-1116		
	7915269	Feb 17, 2029		U-1007		
	7935731	Dec 03, 2028		U-1116		
	7964647	Oct 06, 2028		U-1007		
	7964648	Oct 06, 2028		U-1161		
	7981938	Oct 06, 2028		U-1166		
	8093296	Oct 06, 2028		U-1007		
	8093297	Oct 06, 2028		U-1161		
	8093298	Oct 06, 2028		U-1116		
	8097655	Oct 06, 2028		U-1020		
	8415395	Oct 06, 2028		U-1007		
	8415396	Oct 06, 2028		U-1007		
	8440721	Feb 17, 2029		U-1007		
	8440722	Feb 17, 2029		U-1020		
<b><u>COLCHICINE - MITIGARE</u></b>						
N 204820 001	8927607	Aug 22, 2033		U-1020		
	9399036	Aug 22, 2033		U-1020		
	9555029	Aug 22, 2033		U-1020		
	9675613	Aug 22, 2033		U-1020		
	9789108	Aug 22, 2033		U-1020		
<b><u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u></b>						
N 021176 001	7229613	Apr 17, 2022		U-851		
<b><u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u></b>						
N 022362 001	7229613	Apr 17, 2022		U-493		
<b><u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u></b>						
N 022362 002	7229613	Apr 17, 2022		U-493		
<b><u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u></b>						
N 021697 001	5723606	Dec 15, 2019	DS DP	U-698		
	5723606	Dec 15, 2019	DS DP	U-868		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u></b>						
N 021697 001	5723606	Dec 15, 2019	DS DP U-698			
	5723606	Dec 15, 2019	DS DP U-868			
<b><u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL IN 5% DEXTROSE IN PLASTIC CONTAINER</u></b>						
N 021697 002	5723606	Dec 15, 2019	DS DP U-698			
	5723606	Dec 15, 2019	DS DP U-868			
<b><u>COPANLISIB DIHYDROCHLORIDE - ALIOOPA</u></b>						
N 209936 001	7511041	May 13, 2024	DS DP		NCE	Sep 14, 2022
	8466283	Oct 22, 2029	DS DP U-2124		ODE-155	Sep 14, 2024
	9636344	Mar 29, 2032	U-2124			
<b><u>CORTICOTROPIN - H.P. ACTHAR GEL</u></b>						
N 008372 008					ODE-3	Oct 15, 2017
<b><u>CRISABOROLE - EUCRISA</u></b>						
N 207695 001	8039451	Jun 11, 2026	DS DP		NCE	Dec 14, 2021
	8168614	Jan 20, 2030	U-1932			
	8501712	Feb 16, 2027	U-1932			
	9682092	Feb 16, 2027	U-1932			
<b><u>CRIZOTINIB - XALKORI</u></b>						
N 202570 001	7230098	Aug 26, 2025	DS		M-163	Sep 14, 2018
	7825137	May 12, 2027	U-1179		ODE-111	Mar 11, 2023
	7858643	Oct 08, 2029	DS DP		ODE-15	Aug 26, 2018
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<b><u>CRIZOTINIB - XALKORI</u></b>						
N 202570 002	7230098	Aug 26, 2025	DS		M-163	Sep 14, 2018
	7825137	May 12, 2027	U-1179		ODE-111	Mar 11, 2023
	7858643	Oct 08, 2029	DS DP		ODE-15	Aug 26, 2018
	8217057	Nov 06, 2029	DS DP			
	8785632	Mar 01, 2025	DS			
<b><u>CROFELEMER - FULYZAQ</u></b>						
N 202292 001	7323195	Jun 07, 2018	DP		NCE	Dec 31, 2017
	7341744	Jun 16, 2018	U-1319			
	8574634	Jan 11, 2018	U-1319			
	8962680	Oct 31, 2031	U-1319			
	9585868	Oct 31, 2031	DS U-1319			
<b><u>CYANOCOBALAMIN - NASCOBAL</u></b>						
N 021642 001	7229636	Aug 01, 2024	DP U-817			
	7404489	Mar 12, 2024	DP			
	7879349	Aug 01, 2024	DP U-1152			
	8003353	Aug 01, 2024	U-817			
	8940714	Feb 26, 2024	U-1152			
	9415007	Jul 28, 2024	U-1896			
<b><u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u></b>						
N 021777 001	7387793	Feb 26, 2025	DP			
	7544372	Nov 14, 2023	U-979			
	7790199	Nov 14, 2023	DP			
	7820203	Nov 14, 2023	DP			
	7829121	Nov 14, 2023	U-1088			
	9375410	Nov 14, 2023	U-1088			
	9399025	Nov 14, 2023	DP U-979			
<b><u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u></b>						
N 021777 002	7387793	Feb 26, 2025	DP			
	7544372	Nov 14, 2023	U-979			
	7790199	Nov 14, 2023	DP			
	7820203	Nov 14, 2023	DP			
	7829121	Nov 14, 2023	U-1088			
	9375410	Nov 14, 2023	U-1088			
	9399025	Nov 14, 2023	DP U-979			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>CYCLOSPORINE - RESTASIS</u></b>						
N 050790 001	8629111	Aug 27, 2024	DP			
	8633162	Aug 27, 2024		U-1479		
	8642556	Aug 27, 2024	DP			
	8648048	Aug 27, 2024		U-1483		
	8685930	Aug 27, 2024	DP			
	9248191	Aug 27, 2024		U-1479		
<b><u>CYCLOSPORINE - RESTASIS MULTIDOSE</u></b>						
N 050790 002	8292129	Feb 25, 2031	DP			
	8561859	Apr 16, 2032	DP			
	8629111	Aug 27, 2024	DP			
	8633162	Aug 27, 2024		U-1479		
	8642556	Aug 27, 2024	DP			
	8648048	Aug 27, 2024		U-1483		
	8685930	Aug 27, 2024	DP			
	9248191	Aug 27, 2024		U-1479		
	9669974	May 11, 2034	DP			
	9676525	Feb 07, 2034	DP			
<b><u>CYSTEAMINE BITARTRATE - PROCYSBI</u></b>						
N 203389 001	8026284	Sep 22, 2027	U-1399		M-216	Dec 22, 2020
	8026284*PED	Mar 22, 2028			NPP	Aug 14, 2018
	9173851	Jun 17, 2034	DP		ODE-45	Apr 30, 2020
	9173851*PED	Dec 17, 2034			ODE-97	Aug 14, 2022
	9192590	Jan 26, 2027		U-1399	PED	Oct 30, 2020
	9192590*PED	Jul 26, 2027			PED	Jun 22, 2021
	9198882	Jan 26, 2027		U-1399	PED	Feb 14, 2023
	9198882*PED	Jul 26, 2027				
	9233077	Jun 17, 2034	DP			
	9233077*PED	Dec 17, 2034				
<b><u>CYSTEAMINE BITARTRATE - PROCYSBI</u></b>						
N 203389 002	8026284	Sep 22, 2027	U-1399		M-216	Dec 22, 2020
	8026284*PED	Mar 22, 2028			NPP	Aug 14, 2018
	9173851	Jun 17, 2034	DP		ODE-45	Apr 30, 2020
	9173851*PED	Dec 17, 2034			ODE-97	Aug 14, 2022
	9192590	Jan 26, 2027		U-1399	PED	Oct 30, 2020
	9192590*PED	Jul 26, 2027			PED	Jun 22, 2021
	9198882	Jan 26, 2027		U-1399	PED	Feb 14, 2023
	9198882*PED	Jul 26, 2027				
	9233077	Jun 17, 2034	DP			
	9233077*PED	Dec 17, 2034				
<b><u>CYSTEAMINE HYDROCHLORIDE - CYSTARAN</u></b>						
N 200740 001					ODE-31	Oct 02, 2019
<b><u>CYTARABINE; DAUNORUBICIN - VYXEOS</u></b>						
N 209401 001	7850990	Jan 23, 2027	DP U-2090		NP	Aug 03, 2020
	8022279	Sep 14, 2027	DP U-2090			
	8092828	Apr 01, 2029	U-2090			
	8431806	Apr 22, 2025	DP U-2090			
	8518437	Jun 07, 2026	DP			
	9271931	Jan 23, 2027	DP			
<b><u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u></b>						
N 022512 001	6087380	Dec 28, 2021	DS DP U-1931		M-168	Nov 20, 2018
	7866474	Aug 31, 2027	DP	Y		
	7932273	Sep 07, 2025	DS DP			
	9034822	Jan 20, 2031	U-1759			
<b><u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u></b>						
N 022512 002	6087380	Dec 28, 2021	DS DP U-1931		M-168	Nov 20, 2018
	7866474	Aug 31, 2027	DP	Y		
	7932273	Sep 07, 2025	DS DP			
	9034822	Jan 20, 2031	U-1759			
<b><u>DABIGATRAN ETEXILATE MESYLATE - PRADAXA</u></b>						
N 022512 003	6087380	Dec 28, 2021	DS DP U-1931		NS	Nov 20, 2018
	7866474	Aug 31, 2027	DP	Y		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DABIGATRAN ETEXilate MESYLATE - PRADAXA</b>						
N 022512 003	7932273	Sep 07, 2025	DS DP			
	9034822	Jan 20, 2031		U-1759		
<b>DABRAFENIB MESYLATE - TAFINLAR</b>						
N 202806 001	7994185	Jan 20, 2030	DS DP U-1406		I-745	Jun 22, 2020
	7994185	Jan 20, 2030	DS DP U-2031		M-170	Nov 20, 2018
	7994185	Jan 20, 2030	DS DP U-2032		NCE	May 29, 2018
	8415345	Jan 20, 2030	DS DP U-1406		ODE-147	Jun 22, 2024
	8415345	Jan 20, 2030	DS DP U-2031		ODE-47	May 29, 2020
	8415345	Jan 20, 2030	DS DP U-2032		ODE-58	Jan 09, 2021
	8703781	Oct 15, 2030	DS DP U-1713			
	8835443	Sep 13, 2025		U-2026		
	8835443	Sep 13, 2025		U-2027		
	8952018	Oct 15, 2030		U-2027		
	9233956	May 04, 2029		U-1811		
<b>DABRAFENIB MESYLATE - TAFINLAR</b>						
N 202806 002	7994185	Jan 20, 2030	DS DP U-1406		I-745	Jun 22, 2020
	7994185	Jan 20, 2030	DS DP U-2031		M-170	Nov 20, 2018
	7994185	Jan 20, 2030	DS DP U-2032		NCE	May 29, 2018
	8415345	Jan 20, 2030	DS DP U-1406		ODE-147	Jun 22, 2024
	8415345	Jan 20, 2030	DS DP U-2031		ODE-47	May 29, 2020
	8415345	Jan 20, 2030	DS DP U-2032		ODE-58	Jan 09, 2021
	8703781	Oct 15, 2030	DS DP U-1713			
	8835443	Sep 13, 2025		U-2026		
	8835443	Sep 13, 2025		U-2027		
	8952018	Oct 15, 2030		U-2027		
	9233956	May 04, 2029		U-1811		
<b>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</b>						
N 206843 001	8329159	Apr 13, 2028	DS		D-161	Feb 05, 2019
	8629171	Jun 13, 2031	DS DP U-1724		D-162	Feb 05, 2019
	8642025	Aug 11, 2027	DS DP U-1724		I-726	Feb 05, 2019
	8642025	Aug 11, 2027	DS DP U-1725		I-727	Feb 05, 2019
	8900566	Aug 08, 2027		U-1724		NCE
	8900566	Aug 08, 2027		U-1725		Jul 24, 2020
	9421192	Aug 08, 2027	DS	U-1724		
	9421192	Aug 08, 2027	DS	U-1725		
<b>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</b>						
N 206843 002	8329159	Apr 13, 2028	DS		D-161	Feb 05, 2019
	8629171	Jun 13, 2031	DS DP U-1724		D-162	Feb 05, 2019
	8642025	Aug 11, 2027	DS DP U-1724		I-726	Feb 05, 2019
	8642025	Aug 11, 2027	DS DP U-1725		I-727	Feb 05, 2019
	8900566	Aug 08, 2027		U-1724		NCE
	8900566	Aug 08, 2027		U-1725		Jul 24, 2020
	9421192	Aug 08, 2027	DS	U-1724		
	9421192	Aug 08, 2027	DS	U-1725		
<b>DACLATASVIR DIHYDROCHLORIDE - DAKLINZA</b>						
N 206843 003	9421192	Aug 08, 2027	DS	U-1724		
	9421192	Aug 08, 2027	DS	U-1725		
<b>DALBAVANCIN HYDROCHLORIDE - DALVANCE</b>						
N 021883 001	6900175	Dec 25, 2023		U-1517		D-154
	7115564	Nov 14, 2023	DP			NCE
	7119061	Nov 14, 2023	DP			GAIN
	8143212	Nov 14, 2023		U-1517		May 23, 2024
<b>DALFAMPRIDINE - AMPYRA</b>						
N 022250 001	5540938	Jul 30, 2018		U-1030		
	8007826	May 26, 2027		U-1030		
	8354437	Dec 22, 2026		U-1030		
	8440703	Apr 08, 2025		U-1030		
	8663685	Jan 18, 2025		U-1030		
<b>DANTROLENE SODIUM - RYANODEX</b>						
N 205579 001	7758890	Jul 01, 2025	DP			ODE-69
	8110225	Dec 24, 2022	DP			Jul 22, 2021

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DANTROLENE SODIUM - RYANODEX</b>						
N 205579	001	8604072	Dec 24, 2022	DP		
		8685460	Feb 15, 2023	U-1546		
<b>DAPAGLIFLOZIN PROPANEDIOL - FARXIGA</b>						
N 202293	001	6414126	Oct 04, 2020	DS DP U-2139	M-157	Mar 11, 2018
		6414126	Oct 04, 2020	DS DP U-493	M-212	Oct 20, 2020
		6479065	Aug 10, 2020	DP	Y	NCE
		6495164	May 25, 2020	DP	Y	
		6515117	Oct 04, 2020	DS DP U-2139		
		6515117	Oct 04, 2020	DS DP U-493		
		6667061	May 25, 2020	DP	Y	
		6824822	Oct 09, 2022	DP	Y	
		6872700	Jan 14, 2020	U-654	Y	
		6936590	Oct 04, 2020	U-493		
		6956026	Jan 07, 2018	U-687	Y	
		7223440	Aug 31, 2021	DP	Y	
		7456254	Jun 30, 2025	DP U-1223		
		7456254	Jun 30, 2025	DP U-2139		
		7563871	Apr 15, 2024	DP	Y	
		7612176	Apr 13, 2025	DP U-1223	Y	
		7741269	Jan 07, 2018	U-1224	Y	
		7851502	Aug 19, 2028	DP		
		7919598	Dec 16, 2029	DS		
		8216180	Jan 11, 2028	DP	Y	
		8221786	Mar 21, 2028	DP		
		8329648	Aug 18, 2026	U-1313		
		8329648	Aug 18, 2026	U-2139		
		8329648	Aug 18, 2026	U-2153		
		8329648	Aug 18, 2026	U-2155		
		8329648	Aug 18, 2026	U-2156		
		8329648	Aug 18, 2026	U-2212		
		8329648	Aug 18, 2026	U-2213		
		8361972	Mar 21, 2028	U-2139		
		8361972	Mar 21, 2028	U-493		
		8431685	Apr 13, 2025	DP U-2139		
		8431685	Apr 13, 2025	DP U-412		
		8439864	Mar 25, 2028	DP	Y	
		8461105	Apr 13, 2025	DP U-2139		
		8461105	Apr 13, 2025	DP U-412		
		8501698	Jun 20, 2027	DP U-493		
		8685934	May 26, 2030	U-1522		
		8690837	May 19, 2029	DP	Y	
		8716251	Mar 21, 2028	DP		
		8721615	Jan 18, 2030	DP		
		8758292	Nov 12, 2027	DP	Y	
		8827963	Feb 04, 2029	DP	Y	
		8906851	Aug 18, 2026	U-1313		
		8906851	Aug 18, 2026	U-2139		
		8998876	Jan 07, 2030	DP	Y	
		9198925	Oct 04, 2020	U-2139		
		9198925	Oct 04, 2020	U-493		
		9238076	Apr 15, 2024	DP U-2139		
		9238076	Apr 15, 2024	DP U-412		
		9320853	Mar 25, 2028	DP	Y	
<b>DAPAGLIFLOZIN PROPANEDIOL - FARXIGA</b>						
N 202293	002	6414126	Oct 04, 2020	DS DP U-2139	M-157	Mar 11, 2018
		6414126	Oct 04, 2020	DS DP U-493	M-212	Oct 20, 2020
		6479065	Aug 10, 2020	DP	Y	NCE
		6495164	May 25, 2020	DP	Y	
		6515117	Oct 04, 2020	DS DP U-2139		
		6515117	Oct 04, 2020	DS DP U-493		
		6667061	May 25, 2020	DP	Y	
		6824822	Oct 09, 2022	DP	Y	
		6872700	Jan 14, 2020	U-654	Y	
		6936590	Oct 04, 2020	U-493		
		6956026	Jan 07, 2018	U-687	Y	
		7223440	Aug 31, 2021	DP	Y	

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>DAPAGLIFLOZIN PROPANEDIOL - FARXIGA</b>						
N 202293 002	7456254	Jun 30, 2025	DP U-1223			
	7456254	Jun 30, 2025	DP U-2139			
	7563871	Apr 15, 2024	DP	Y		
	7612176	Apr 13, 2025	DP U-1223	Y		
	7741269	Jan 07, 2018	U-1224	Y		
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8216180	Jan 11, 2028	DP	Y		
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-1313			
	8329648	Aug 18, 2026	U-2139			
	8329648	Aug 18, 2026	U-2153			
	8329648	Aug 18, 2026	U-2155			
	8329648	Aug 18, 2026	U-2156			
	8329648	Aug 18, 2026	U-2212			
	8329648	Aug 18, 2026	U-2213			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-2139			
	8431685	Apr 13, 2025	DP U-412			
	8439864	Mar 25, 2028	DP	Y		
	8461105	Apr 13, 2025	DP U-2139			
	8461105	Apr 13, 2025	DP U-412			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8690837	May 19, 2029	DP	Y		
	8716251	Mar 21, 2028	DP			
	8721615	Jan 18, 2030	DP			
	8758292	Nov 12, 2027	DP	Y		
	8827963	Feb 04, 2029	DP	Y		
	8906851	Aug 18, 2026	U-1313			
	8906851	Aug 18, 2026	U-2139			
	8998876	Jan 07, 2030	DP	Y		
	9198925	Oct 04, 2020	U-2139			
	9198925	Oct 04, 2020	U-493			
	9238076	Apr 15, 2024	DP U-2139			
	9238076	Apr 15, 2024	DP U-412			
	9320853	Mar 25, 2028	DP	Y		
<b>DAPAGLIFLOZIN PROPANEDIOL; METFORMIN HYDROCHLORIDE - XIGDUO XR</b>						
N 205649 001	6414126	Oct 04, 2020	DS DP U-493		NCE	Jan 08, 2019
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
<b>DAPAGLIFLOZIN PROPANEDIOL; METFORMIN HYDROCHLORIDE - XIGDUO XR</b>						
N 205649 002	6414126	Oct 04, 2020	DS DP U-493		NCE	Jan 08, 2019
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			
<b>DAPAGLIFLOZIN PROPANEDIOL; METFORMIN HYDROCHLORIDE - XIGDUO XR</b>						
N 205649 003	6414126	Oct 04, 2020	DS DP U-493		NCE	Jan 08, 2019
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020	U-493			
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	9198925	Oct 04, 2020	U-493			
	9616028	Nov 12, 2030	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DAPAGLIFLOZIN PROPANEDIOL; METFORMIN HYDROCHLORIDE - XIGDUO XR</b>						
N 205649 004	6414126	Oct 04, 2020	DS DP U-493		NCE	Jan 08, 2019
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020		U-493		
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027		DP U-493		
	8685934	May 26, 2030		U-1522		
	9198925	Oct 04, 2020		U-493		
	9616028	Nov 12, 2030		DP		
<b>DAPAGLIFLOZIN PROPANEDIOL; METFORMIN HYDROCHLORIDE - XIGDUO XR</b>						
N 205649 005	6414126	Oct 04, 2020	DS DP U-493		NCE	Jan 08, 2019
	6515117	Oct 04, 2020	DS DP U-493			
	6936590	Oct 04, 2020		U-493		
	7919598	Dec 16, 2029	DS			
	8501698	Jun 20, 2027		DP U-493		
	8685934	May 26, 2030		U-1522		
	9198925	Oct 04, 2020		U-493		
	9616028	Nov 12, 2030		DP		
<b>DAPAGLIFLOZIN PROPANEDIOL; SAXAGLIPTIN HYDROCHLORIDE - QTERN</b>						
N 209091 001	6414126	Oct 04, 2020	DS DP U-1976		M-175	Apr 05, 2019
	6414126	Oct 04, 2020	DS DP U-1977		NC	Feb 27, 2020
	6515117	Oct 04, 2020	DS DP U-1976		NCE	Jan 08, 2019
	6515117	Oct 04, 2020	DS DP U-1977			
	6936590	Oct 04, 2020		U-1976		
	6936590	Oct 04, 2020		U-1977		
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028		DP		
	8361972	Mar 21, 2028		U-1976		
	8361972	Mar 21, 2028		U-1977		
	8501698	Jun 20, 2027		DP U-1976		
	8501698	Jun 20, 2027		DP U-1977		
	8628799	Jul 13, 2025		DP		
	8716251	Mar 21, 2028		DP		
	9198925	Oct 04, 2020		U-1976		
	9198925	Oct 04, 2020		U-1977		
	RE44186	Jul 31, 2023	DS DP U-1976		M-211	Sep 01, 2020
	RE44186	Jul 31, 2023	DS DP U-1977		NPP	Mar 29, 2020
<b>DAPSONE - ACZONE</b>						
N 207154 001	9161926	Nov 18, 2033	DP		NS	Feb 24, 2019
	9517219	Nov 18, 2033		U-1033		
<b>DAPTOMYCIN - CUBICIN</b>						
N 021572 002	8003673	Sep 04, 2028		U-1180	M-211	Sep 01, 2020
					NPP	Mar 29, 2020
<b>DAPTOMYCIN - CUBICIN RF</b>						
N 021572 003	9138456	Nov 23, 2030	DP		M-211	Sep 01, 2020
					NPP	Mar 29, 2020
<b>DARUNAVIR ETHANOATE - PREZISTA</b>						
N 021976 001	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7470506	Jun 23, 2019		U-935		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		
	8597876*PED	Dec 23, 2019				
<b>DARUNAVIR ETHANOATE - PREZISTA</b>						
N 021976 002	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7470506	Jun 23, 2019		U-935		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DARUNAVIR ETHANOLATE - PREZISTA</b>						
N 021976 002	8597876*PED	Dec 23, 2019				
<b>DARUNAVIR ETHANOLATE - PREZISTA</b>						
N 021976 003	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7470506	Jun 23, 2019		U-935		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		
	8597876*PED	Dec 23, 2019				
<b>DARUNAVIR ETHANOLATE - PREZISTA</b>						
N 021976 004	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7470506	Jun 23, 2019		U-935		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		
	8597876*PED	Dec 23, 2019				
<b>DARUNAVIR ETHANOLATE - PREZISTA</b>						
N 021976 005	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7470506	Jun 23, 2019		U-935		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		
	8597876*PED	Dec 23, 2019				
<b>DARUNAVIR ETHANOLATE - PREZISTA</b>						
N 021976 006	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7470506	Jun 23, 2019		U-935		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		
	8597876*PED	Dec 23, 2019				
<b>DARUNAVIR ETHANOLATE - PREZISTA</b>						
N 202895 001	7470506	Jun 23, 2019		U-1209		
	7470506	Jun 23, 2019		U-1305		
	7700645	Dec 26, 2026	DS DP			
	8518987	Feb 16, 2024	DS DP			
	8518987*PED	Aug 16, 2024				
	8597876	Jun 23, 2019		U-1305		
	8597876*PED	Dec 23, 2019				
<b>DASABUVIR SODIUM : OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA PAK (COPACKAGED)</b>						
N 206619 001	7148359	Jul 19, 2019	DP		D-163	Apr 22, 2019
	7364752	Nov 10, 2020	DP		NCE	Dec 19, 2019
	8188104	May 17, 2029	DS DP	U-1636		
	8268349	Aug 25, 2024	DP			
	8399015	Aug 25, 2024	DP			
	8420596	Apr 10, 2031	DS DP			
	8466159	Sep 04, 2032		U-1637		
	8492386	Sep 04, 2032		U-1840		
	8501238	Sep 17, 2028	DS DP	U-1636		
	8642538	Sep 10, 2029	DS DP	U-1638		
	8680106	Sep 04, 2032		U-1637		
	8685984	Sep 04, 2032		U-1840		
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030		U-1687		
	9044480	Apr 10, 2031		U-1638		
	9139536	Nov 09, 2028		U-1753		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DASABUVIR SODIUM : OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA PAK (COPACKAGED)</u>						
N 206619	001	9629841	Oct 18, 2033	DP U-1753		
<u>DASABUVIR SODIUM; OMBITASVIR; PARITAPREVIR; RITONAVIR - VIEKIRA XR</u>						
N 208624	001	7148359	Jul 19, 2019	DP		
		7364752	Nov 10, 2020	DP		
		8188104	May 17, 2029	DS DP U-1636		
		8268349	Aug 25, 2024	DP		
		8399015	Aug 25, 2024	DP		
		8420596	Apr 10, 2031	DS DP		
		8466159	Sep 04, 2032	U-1637		
		8492386	Sep 04, 2032	U-1840		
		8501238	Sep 17, 2028	DS DP U-1636		
		8642538	Sep 10, 2029	DS DP U-1638		
		8680106	Sep 04, 2032	U-1637		
		8685984	Sep 04, 2032	U-1840		
		8686026	Jun 09, 2031	DP		
		8691938	Apr 13, 2032	DS DP		
		9006387	Jun 10, 2030	U-1687		
		9044480	Apr 10, 2031	U-1638		
		9139536	Nov 09, 2028	U-1753		
		9333204	Jan 02, 2035	DP U-1889		
		9744170	Jan 02, 2035	DP U-1889		
<u>DASATINIB - SPRYCEL</u>						
N 021986	001	6596746	Jun 28, 2020	DS DP U-748		
		6596746	Jun 28, 2020	DS DP U-780		
		7125875	Apr 13, 2020	U-779		
		7125875	Apr 13, 2020	U-780		
		7153856	Apr 28, 2020	U-780		
		7491725	Mar 28, 2026	DS DP		
		8680103	Feb 04, 2025	DP		
<u>DASATINIB - SPRYCEL</u>						
N 021986	002	6596746	Jun 28, 2020	DS DP U-748		
		6596746	Jun 28, 2020	DS DP U-780		
		7125875	Apr 13, 2020	U-779		
		7125875	Apr 13, 2020	U-780		
		7153856	Apr 28, 2020	U-780		
		7491725	Mar 28, 2026	DS DP		
		8680103	Feb 04, 2025	DP		
<u>DASATINIB - SPRYCEL</u>						
N 021986	003	6596746	Jun 28, 2020	DS DP U-748		
		6596746	Jun 28, 2020	DS DP U-780		
		7125875	Apr 13, 2020	U-779		
		7125875	Apr 13, 2020	U-780		
		7153856	Apr 28, 2020	U-780		
		7491725	Mar 28, 2026	DS DP		
		8680103	Feb 04, 2025	DP		
<u>DASATINIB - SPRYCEL</u>						
N 021986	004	6596746	Jun 28, 2020	DS DP U-748		
		6596746	Jun 28, 2020	DS DP U-780		
		7125875	Apr 13, 2020	U-779		
		7125875	Apr 13, 2020	U-780		
		7153856	Apr 28, 2020	U-780		
		7491725	Mar 28, 2026	DS DP		
		8680103	Feb 04, 2025	DP		
<u>DASATINIB - SPRYCEL</u>						
N 021986	005	6596746	Jun 28, 2020	DS DP U-748		
		6596746	Jun 28, 2020	DS DP U-780		
		7125875	Apr 13, 2020	U-779		
		7125875	Apr 13, 2020	U-780		
		7153856	Apr 28, 2020	U-780		
		7491725	Mar 28, 2026	DS DP		
		8680103	Feb 04, 2025	DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DASATINIB - SPRYCEL</b>						
N 021986 006	6596746	Jun 28, 2020	DS DP U-748		NPP	Nov 09, 2020
	6596746	Jun 28, 2020	DS DP U-780			
	7125875	Apr 13, 2020		U-779		
	7125875	Apr 13, 2020		U-780		
	7153856	Apr 28, 2020		U-780		
	7491725	Mar 28, 2026	DS DP			
	8680103	Feb 04, 2025	DP			
<b>DEFERASIROX - EXJADE</b>						
N 021882 001	6465504	Apr 05, 2019	DS DP		ODE-39	Jan 23, 2020
<b>DEFERASIROX - EXJADE</b>						
N 021882 002	6465504	Apr 05, 2019	DS DP		ODE-39	Jan 23, 2020
<b>DEFERASIROX - EXJADE</b>						
N 021882 003	6465504	Apr 05, 2019	DS DP		ODE-39	Jan 23, 2020
<b>DEFERASIROX - JADENU</b>						
N 206910 001	6465504	Apr 05, 2019	DS DP		ODE-39	Jan 23, 2020
	9283209	Nov 21, 2034	DS DP			
<b>DEFERASIROX - JADENU</b>						
N 206910 002	6465504	Apr 05, 2019	DS DP		ODE-39	Jan 23, 2020
	9283209	Nov 21, 2034	DS DP			
<b>DEFERASIROX - JADENU</b>						
N 206910 003	6465504	Apr 05, 2019	DS DP		ODE-39	Jan 23, 2020
	9283209	Nov 21, 2034	DS DP			
<b>DEFERASIROX - JADENU SPRINKLE</b>						
N 207968 001	6465504	Apr 05, 2019	DS DP			
<b>DEFERASIROX - JADENU SPRINKLE</b>						
N 207968 002	6465504	Apr 05, 2019	DS DP			
<b>DEFERASIROX - JADENU SPRINKLE</b>						
N 207968 003	6465504	Apr 05, 2019	DS DP			
<b>DEFERIPRONE - FERRIPROX</b>						
N 021825 001	7049328	Jun 28, 2021		U-735	ODE-16	Oct 14, 2018
<b>DEFERIPRONE - FERRIPROX</b>						
N 208030 001	7049328	Jun 28, 2021		U-735	ODE-16	Oct 14, 2018
	8703156	Oct 29, 2029		DP U-735		
<b>DEFIBROTIDE SODIUM - DEFITELIO</b>						
N 208114 001					NCE	Mar 30, 2021
					ODE-112	Mar 30, 2023
<b>DEFLAZACORT - EMFLAZA</b>						
N 208684 001					NCE	Feb 09, 2022
					ODE-130	Feb 09, 2024
<b>DEFLAZACORT - EMFLAZA</b>						
N 208684 002					NCE	Feb 09, 2022
					ODE-130	Feb 09, 2024
<b>DEFLAZACORT - EMFLAZA</b>						
N 208684 003					NCE	Feb 09, 2022
					ODE-130	Feb 09, 2024
<b>DEFLAZACORT - EMFLAZA</b>						
N 208684 004					NCE	Feb 09, 2022
					ODE-130	Feb 09, 2024
<b>DEFLAZACORT - EMFLAZA</b>						
N 208685 001					NCE	Feb 09, 2022
					ODE-130	Feb 09, 2024

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DEGARELIX ACETATE - FIRMAGON</u></b>						
N 022201 001	5925730	May 18, 2021	DS DP U-943			
	9415085	Apr 27, 2032		U-1895		
	9579359	Feb 10, 2029		U-1978		
<b><u>DEGARELIX ACETATE - FIRMAGON</u></b>						
N 022201 002	5925730	May 18, 2021	DS DP U-943			
	9415085	Apr 27, 2032		U-1895		
	9579359	Feb 10, 2029		U-1978		
<b><u>DELAFLOXACIN MEGLUMINE - BAXDELA</u></b>						
N 208610 001	7728143	Nov 20, 2027	DS		NCE	Jun 19, 2022
	8252813	Oct 02, 2026		DP U-2028	GAIN	Jun 19, 2027
	8273892	Aug 06, 2026	DS			
	8497378	Dec 28, 2029	DS			
	8648093	Oct 07, 2025		DP U-2028		
	8871938	Sep 23, 2029	DS			
	8969569	Oct 07, 2025		DP U-2028		
	9539250	Oct 07, 2025	DS	DP U-2028		
	RE46617	Dec 28, 2029	DS			
<b><u>DELAFLOXACIN MEGLUMINE - BAXDELA</u></b>						
N 208611 001	7635773	Mar 13, 2029	DP		NCE	Jun 19, 2022
	7728143	Nov 20, 2027	DS		GAIN	Jun 19, 2027
	8252813	Oct 02, 2026		DP U-2028		
	8273892	Aug 06, 2026	DS			
	8410077	Mar 13, 2029	DP			
	8497378	Dec 28, 2029	DS			
	8648093	Oct 07, 2025		DP U-2028		
	8871938	Sep 23, 2029	DS			
	9200088	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
	9539250	Oct 07, 2025	DS	DP U-2028		
	9750822	Mar 13, 2029	DP			
	RE46617	Dec 28, 2029	DS			
<b><u>DELAVIRDINE MESYLATE - RESRIPTOR</u></b>						
N 020705 002	6177101	Jun 07, 2019				
<b><u>DEOXYCHOLIC ACID - KYBELLA</u></b>						
N 206333 001	7622130	Dec 10, 2027		U-1690		
	7754230	Dec 10, 2027		U-1690		
	8101593	Mar 02, 2030		DP		
	8242294	May 16, 2028	DS			
	8298556	Aug 03, 2025		U-1690		
	8367649	Mar 02, 2030	DP			
	8461140	Feb 21, 2028	DP			
	8546367	Feb 21, 2028		DP U-1690		
	8653058	Mar 02, 2030	DP			
	8846066	Feb 08, 2025		U-1690		
	8883770	Feb 21, 2028	DP			
	9522155	Feb 21, 2028		DP U-1940		
	9636349	Feb 21, 2028		U-1940		
<b><u>DESLORATADINE - CLARINEX</u></b>						
N 021165 001	6100274	Jul 07, 2019				
	7405223	Jul 07, 2019		U-886		
<b><u>DESLORATADINE - CLARINEX</u></b>						
N 021300 001	6514520	Jun 01, 2018	DP			
<b><u>DESLORATADINE - CLARINEX</u></b>						
N 021312 001	6100274	Jul 07, 2019	DP			
	7618649	Dec 19, 2020		DP U-1017		
<b><u>DESLORATADINE - CLARINEX</u></b>						
N 021312 002	6100274	Jul 07, 2019	DP			
	7618649	Dec 19, 2020		DP U-1017		

**PREScription AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR</u>						
N 021313 001	6100274	Jul 07, 2019	DP			
	6709676	Feb 18, 2021	DP U-707			
	7618649	Dec 19, 2020	DP U-1017			
	8187630	Dec 19, 2020	DP U-1017			
<u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX D 24 HOUR</u>						
N 021605 001	6100274	Jul 07, 2019	DP			
	6979463	Mar 28, 2022	DP			
	7618649	Dec 19, 2020	DP U-1017			
	7820199	Mar 28, 2022	DP			
<u>DESMOPRESSIN ACETATE - NOCTIVA</u>						
N 201656 001	7405203	May 06, 2023	U-1980		NP	Mar 03, 2020
	7579321	May 06, 2023	U-1980			
	7799761	Sep 26, 2024	DP			
	9539302	Jun 15, 2030	DP			
<u>DESMOPRESSIN ACETATE - NOCTIVA</u>						
N 201656 002	7405203	May 06, 2023	U-1980		NP	Mar 03, 2020
	7579321	May 06, 2023	U-1980			
	9539302	Jun 15, 2030	DP			
<u>DESONIDE - DESONATE</u>						
N 021844 001	6387383	Aug 03, 2020	DS DP U-783			
<u>DESONIDE - VERDESO</u>						
N 021978 001	6730288	Sep 08, 2019	DP			
	7029659	Sep 08, 2019	DP			
	8460641	Nov 05, 2028	DP U-1412			
	8962000	Aug 31, 2025	DP U-1412			
	9492384	Aug 31, 2025	DP U-1412			
<u>DESOXIMETASONE - TOPICORT</u>						
N 204141 001	5990100	Mar 24, 2018	DP U-1408			
	8277780	Sep 01, 2028	DP U-1408			
	8715624	May 26, 2026	DP U-1408			
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204003 001					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204003 002					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204028 001					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204028 002					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204065 002					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204065 003					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204082 001					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204083 001					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204095 001					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204095 002					PC	Aug 28, 2017
<u>DESVENLAFAKINE SUCCINATE - DESVENLAFAKINE SUCCINATE</u>						
A 204172 001					PC	Aug 28, 2017

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DESVENLAFAXINE SUCCINATE - DESVENLAFAXINE SUCCINATE</u></b>						
A 204172 002					PC	Aug 28, 2017
<b><u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u></b>						
N 021992 001	6673838	Mar 01, 2022	DS	U-1364		
	6673838	Mar 01, 2022	DS	U-860		
	8269040	Jul 05, 2027	DS			
<b><u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u></b>						
N 021992 002	6673838	Mar 01, 2022	DS	U-1364		
	6673838	Mar 01, 2022	DS	U-860		
	8269040	Jul 05, 2027	DS			
<b><u>DESVENLAFAXINE SUCCINATE - PRISTIQ</u></b>						
N 021992 003	6673838	Mar 01, 2022	DS	U-1364		
	6673838	Mar 01, 2022	DS	U-860		
	8269040	Jul 05, 2027	DS			
<b><u>DEUTETRABENAZINE - AUSTEDO</u></b>						
N 208082 001	8524733	Mar 27, 2031	DS DP		I-751	Aug 30, 2020
	9233959	Sep 18, 2033	DP		NCE	Apr 03, 2022
	9296739	Sep 18, 2033	DP		ODE-134	Apr 03, 2024
	9550780	Sep 18, 2033	DS DP	U-1846		
	9550780	Sep 18, 2033	DS DP	U-1995		
	9814708	Sep 18, 2033	DP			
<b><u>DEUTETRABENAZINE - AUSTEDO</u></b>						
N 208082 002	8524733	Mar 27, 2031	DS DP		I-751	Aug 30, 2020
	9233959	Sep 18, 2033	DP		NCE	Apr 03, 2022
	9296739	Sep 18, 2033	DP		ODE-134	Apr 03, 2024
	9550780	Sep 18, 2033	DS DP	U-1846		
	9550780	Sep 18, 2033	DS DP	U-1995		
	9814708	Sep 18, 2033	DP			
<b><u>DEUTETRABENAZINE - AUSTEDO</u></b>						
N 208082 003	8524733	Mar 27, 2031	DS DP		I-751	Aug 30, 2020
	9233959	Sep 18, 2033	DP		NCE	Apr 03, 2022
	9296739	Sep 18, 2033	DP		ODE-134	Apr 03, 2024
	9550780	Sep 18, 2033	DS DP	U-1846		
	9550780	Sep 18, 2033	DS DP	U-1995		
	9814708	Sep 18, 2033	DP			
<b><u>DEXAMETHASONE - OZURDEX</u></b>						
N 022315 001	6726918	Oct 20, 2020	DP	U-1204		
	6726918	Oct 20, 2020	DP	U-1205		
	6899717	Nov 01, 2023		U-1206		
	7033605	Oct 20, 2020	DP			
	7767223	Nov 28, 2021	DP			
	8034366	Jan 09, 2023	DP	U-1204		
	8034366	Jan 09, 2023	DP	U-1205		
	8034370	Jan 09, 2023	DP			
	8043628	Oct 20, 2020		U-1205		
	8063031	Oct 20, 2020	DP			
	8088407	Oct 20, 2020		U-1205		
	8506987	Jan 09, 2023		U-1204		
	8506987	Jan 09, 2023		U-1205		
	9012437	Oct 20, 2020		U-1205		
	9192511	Jan 09, 2023	DP			
	9283178	Oct 20, 2020		U-1205		
	9592242	Oct 20, 2020		U-1989		
	9592242	Oct 20, 2020		U-1990		
	9775849	Oct 20, 2020		U-1989		
	9775849	Oct 20, 2020		U-1990		
<b><u>DEXAMETHASONE; TOBRAMYCIN - TOBRADEX ST</u></b>						
N 050818 001	7795316	Aug 03, 2028	DP	U-1082		
	8101582	Dec 19, 2027	DP	U-1082		
	8450287	Dec 19, 2027	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DEXLANSOPRAZOLE - DEXILANT</u></b>						
N 022287 001	6462058	Jun 15, 2020	DS DP U-949		NPP	Jul 08, 2019
	6462058	Jun 15, 2020	DS DP U-950		NPP	Jul 08, 2019
	6462058	Jun 15, 2020	DS DP U-951		NPP	Jul 08, 2019
	6664276	Jan 30, 2023	DS DP U-1507			
	6664276	Jan 30, 2023	DS DP U-949			
	6664276	Jan 30, 2023	DS DP U-950			
	6664276	Jan 30, 2023	DS DP U-951			
	6664276*PED	Jul 30, 2023				
	6939971	Jun 15, 2020	U-949			
	6939971	Jun 15, 2020	U-950			
	6939971	Jun 15, 2020	U-951			
	7285668	Jun 15, 2020	DS			
	7790755	Aug 02, 2026	DP			
	8105626	Sep 27, 2026	DP			
	8173158	Mar 17, 2030	U-949			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-951			
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8722084	Oct 15, 2023	DP			
	8722084*PED	Apr 15, 2024				
	8784885	Oct 15, 2023	DP U-1552			
	8784885	Oct 15, 2023	DP U-1553			
	8784885	Oct 15, 2023	DP U-1554			
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	9011926	Feb 24, 2026	DP			
	9145389	Jun 15, 2020	DS DP			
	9233103	Mar 05, 2032	U-1805			
	9238029	Jan 17, 2026	DP			
<b><u>DEXLANSOPRAZOLE - DEXILANT</u></b>						
N 022287 002	6462058	Jun 15, 2020	DS DP U-949		NPP	Jul 08, 2019
	6462058	Jun 15, 2020	DS DP U-950		NPP	Jul 08, 2019
	6462058	Jun 15, 2020	DS DP U-951		NPP	Jul 08, 2019
	6664276	Jan 30, 2023	DS DP U-1507			
	6664276	Jan 30, 2023	DS DP U-949			
	6664276	Jan 30, 2023	DS DP U-950			
	6664276	Jan 30, 2023	DS DP U-951			
	6664276*PED	Jul 30, 2023				
	6939971	Jun 15, 2020	U-949			
	6939971	Jun 15, 2020	U-950			
	6939971	Jun 15, 2020	U-951			
	7285668	Jun 15, 2020	DS			
	7790755	Aug 02, 2026	DP			
	8105626	Sep 27, 2026	DP			
	8173158	Mar 17, 2030	U-949			
	8173158	Mar 17, 2030	U-950			
	8173158	Mar 17, 2030	U-951			
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8722084	Oct 15, 2023	DP			
	8722084*PED	Apr 15, 2024				
	8784885	Oct 15, 2023	DP U-1552			
	8784885	Oct 15, 2023	DP U-1553			
	8784885	Oct 15, 2023	DP U-1554			
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	9011926	Feb 24, 2026	DP			
	9145389	Jun 15, 2020	DS DP			
	9233103	Mar 05, 2032	U-1805			
	9238029	Jan 17, 2026	DP			
<b><u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u></b>						
N 208056 001	6328994	May 17, 2019	DP			
	6328994*PED	Nov 17, 2019				
	6462058	Jun 15, 2020	DS DP U-950			
	6462058	Jun 15, 2020	DS DP U-951			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DEXLANSOPRAZOLE - DEXILANT SOLUTAB</u></b>						
N 208056 001	6462058*PED	Dec 15, 2020				
	6664276	Jan 30, 2023	DS DP	U-950		
	6664276	Jan 30, 2023	DS DP	U-951		
	6664276*PED	Jul 30, 2023				
	6939971	Jun 15, 2020		U-950		
	6939971	Jun 15, 2020		U-951		
	6939971*PED	Dec 15, 2020				
	7285668	Jun 15, 2020	DS			
	7285668*PED	Dec 15, 2020				
	7399485	May 26, 2018	DP			
	7399485*PED	Nov 26, 2018				
	7431942	May 17, 2019	DP			
	7431942*PED	Nov 17, 2019				
	7875292	May 17, 2019	DP			
	7875292*PED	Nov 17, 2019				
	8461187	Jan 17, 2026	DP			
	8461187*PED	Jul 17, 2026				
	8784885	Oct 15, 2023	DP			
	8784885*PED	Apr 15, 2024				
	8871273	Jan 11, 2028	DP			
	8871273*PED	Jul 11, 2028				
	9011926	Feb 24, 2026	DP			
	9145389	Jun 15, 2020	DS DP			
	9238029	Jan 17, 2026	DP			
	9241910	Mar 10, 2029	DP			
<b><u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u></b>						
N 021038 001	6716867	Mar 31, 2019		U-1472		
	6716867*PED	Oct 01, 2019				
<b><u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u></b>						
N 021038 002	6716867	Mar 31, 2019		U-1472		
	6716867*PED	Oct 01, 2019				
	8242158	Jan 04, 2032	DP			
	8242158*PED	Jul 04, 2032				
	8338470	Jan 04, 2032	DP			
	8338470*PED	Jul 04, 2032				
	8455527	Jan 04, 2032	U-421			
	8455527*PED	Jul 04, 2032				
	8648106	Jan 04, 2032	DP			
	8648106*PED	Jul 04, 2032				
	9320712	Jan 04, 2032	DP			
	9320712*PED	Jul 04, 2032				
	9616049	Jan 04, 2032	DP			
	9616049*PED	Jul 04, 2032				
<b><u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u></b>						
N 021038 003	6716867	Mar 31, 2019		U-1472		
	6716867*PED	Oct 01, 2019				
	8242158	Jan 04, 2032	DP			
	8242158*PED	Jul 04, 2032				
	8338470	Jan 04, 2032	DP			
	8338470*PED	Jul 04, 2032				
	8455527	Jan 04, 2032	U-421			
	8455527*PED	Jul 04, 2032				
	8648106	Jan 04, 2032	DP			
	8648106*PED	Jul 04, 2032				
	9320712	Jan 04, 2032	DP			
	9320712*PED	Jul 04, 2032				
	9616049	Jan 04, 2032	DP			
	9616049*PED	Jul 04, 2032				
<b><u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u></b>						
N 021038 004	6716867	Mar 31, 2019		U-1472		
	6716867*PED	Oct 01, 2019				
	8242158	Jan 04, 2032	DP			
	8242158*PED	Jul 04, 2032				
	8338470	Jan 04, 2032	DP			
	8338470*PED	Jul 04, 2032				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DEXMEDETOMIDINE HYDROCHLORIDE - PRECEDEX</u></b>						
N 021038 004	8455527	Jan 04, 2032	U-421			
	8455527*PED	Jul 04, 2032				
	8648106	Jan 04, 2032	DP			
	8648106*PED	Jul 04, 2032				
	9320712	Jan 04, 2032	DP			
	9320712*PED	Jul 04, 2032				
	9616049	Jan 04, 2032	DP			
	9616049*PED	Jul 04, 2032				
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - DEXMETHYLPHENIDATE HYDROCHLORIDE</u></b>						
A 202842 005				PC		Jul 04, 2017
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - DEXMETHYLPHENIDATE HYDROCHLORIDE</u></b>						
A 202842 007				PC		Jul 04, 2017
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 001	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 002	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 003	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 004	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 005	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 006	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 007	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u></b>						
N 021802 008	6228398	Nov 01, 2019	DP U-676			
	6730325	Nov 01, 2019	DP U-676			
<b><u>DEXRAZOXANE HYDROCHLORIDE - TOTECT</u></b>						
N 022025 001	6727253	Mar 13, 2020	U-829			
<b><u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u></b>						
N 021620 001	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP U-685			
	7838032	Apr 28, 2020	DP			
<b><u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN - MUCINEX DM</u></b>						
N 021620 002	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP U-685			
	7838032	Apr 28, 2020	DP			
<b><u>DEXTROMETHORPHAN HYDROBROMIDE; QUINIDINE SULFATE - NUEDEXTA</u></b>						
N 021879 001	7659282	Aug 13, 2026	U-1093			
	8227484	Jul 17, 2023	U-1093			
<b><u>DICHLORPHENAMIDE - KEVEYIS</u></b>						
N 011366 002				ODE-96		Aug 07, 2022

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DICLOFENAC - ZORVOLEX</u></b>						
N 204592 001	8679544	Apr 23, 2030	DP		I-692	Aug 22, 2017
	8999387	Apr 23, 2030		U-55		
	9017721	Apr 23, 2030		DP		
	9173854	Apr 23, 2030		DP		
	9180095	Apr 23, 2030			U-55	
	9180096	Apr 23, 2030		DP		
	9186328	Apr 23, 2030			U-55	
<b><u>DICLOFENAC - ZORVOLEX</u></b>						
N 204592 002	8679544	Apr 23, 2030	DP		I-692	Aug 22, 2017
	8999387	Apr 23, 2030		U-55		
	9017721	Apr 23, 2030		DP		
	9173854	Apr 23, 2030		DP		
	9180095	Apr 23, 2030			U-55	
	9180096	Apr 23, 2030		DP		
	9186328	Apr 23, 2030			U-55	
<b><u>DICLOFENAC EPOLAMINE - FLECTOR</u></b>						
N 021234 001	5607690	Apr 13, 2019	DP			
<b><u>DICLOFENAC POTASSIUM - CAMBIA</u></b>						
N 022165 001	7759394	Jun 16, 2026	DS DP	U-436		
	8097651	Jun 16, 2026	DS DP	U-436		
	8927604	Jun 16, 2026		U-436		
	9827197	Jun 16, 2026	DP			
<b><u>DICLOFENAC POTASSIUM - ZIPSOR</u></b>						
N 022202 001	6287594	Jan 15, 2019	DP			
	6365180	Jul 15, 2019	DP	U-980		
	7662858	Feb 24, 2029		U-1035		
	7884095	Feb 24, 2029		U-1111		
	7939518	Feb 24, 2029		U-980		
	8110606	Feb 24, 2029		U-980		
	8623920	Feb 24, 2029		U-1482		
	9561200	Feb 24, 2029		U-1482		
<b><u>DICLOFENAC SODIUM - PENNSAID</u></b>						
N 020947 001	8217078	Jul 10, 2029		U-1248		
	8546450	Aug 09, 2030		U-1435		
	8546450	Aug 09, 2030		U-1436		
	8618164	Jul 10, 2029		U-1477		
	8741956	Jul 10, 2029		U-1435		
<b><u>DICLOFENAC SODIUM - DYLOJECT</u></b>						
N 022396 001	6407079	Jun 18, 2019	DP		NP	
	8946292	Mar 22, 2027		U-1659		Dec 23, 2017
<b><u>DICLOFENAC SODIUM - PENNSAID</u></b>						
N 204623 001	8217078	Jul 10, 2029		U-1477		
	8252838	Apr 21, 2028	DP	U-1489		
	8546450	Aug 09, 2030		U-1435		
	8546450	Aug 09, 2030		U-1436		
	8563613	Oct 17, 2027	DP	U-1488		
	8618164	Jul 10, 2029		U-1477		
	8741956	Jul 10, 2029		U-1435		
	8871809	Oct 17, 2027		U-1614		
	9066913	Oct 17, 2027	DP	U-1488		
	9101591	Oct 17, 2027	DP	U-1488		
	9132110	Oct 17, 2027		U-1488		
	9168304	Oct 17, 2027	DP			
	9168305	Oct 17, 2027		U-1488		
	9220784	Oct 17, 2027		U-1488		
	9339551	Oct 17, 2027		U-1488		
	9339552	Oct 17, 2027	DP	U-1488		
	9370501	Jul 10, 2029		U-1614		
	9375412	Jul 10, 2029		U-1614		
	9415029	Jul 10, 2029		U-1614		
	9539335	Oct 17, 2027		U-1614		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DIENOGEST; DIENOGEST; DIENOGEST; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE; ESTRADIOL VALERATE - NATAZIA</u></b>						
N 022252 001	8071577 8153616	May 13, 2026 Jan 30, 2028		DP U-1 U-1240		
<b><u>DIFLUPREDNATE - DUREZOL</u></b>						
N 022212 001	6114319 6114319*PED	May 18, 2019 Nov 18, 2019		DP	ODE-26 PED	Jun 13, 2019 Dec 13, 2019
<b><u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u></b>						
N 021392 001	6923984 7108866	Feb 25, 2021 Dec 17, 2019		DP DP U-107		
<b><u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u></b>						
N 021392 002	6923984 7108866	Feb 25, 2021 Dec 17, 2019		DP DP U-107		
<b><u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u></b>						
N 021392 003	6923984 7108866	Feb 25, 2021 Dec 17, 2019		DP DP U-107		
<b><u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u></b>						
N 021392 004	6923984 7108866	Feb 25, 2021 Dec 17, 2019		DP DP U-107		
<b><u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u></b>						
N 021392 005	6923984 7108866	Feb 25, 2021 Dec 17, 2019		DP DP U-107		
<b><u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u></b>						
N 021392 006	6923984 7108866	Feb 25, 2021 Dec 17, 2019		DP DP U-107		
<b><u>DIMETHYL FUMARATE - TECFIDERA</u></b>						
N 204063 001	6509376 7320999 7619001 7803840 8399514 8524773 8759393	Oct 29, 2019 May 18, 2020 Apr 01, 2018 Apr 01, 2018 Feb 07, 2028 Apr 01, 2018 Oct 29, 2019		DP U-1384 U-1384 U-1385 U-1384 U-1384 DP	NCE	Mar 27, 2018
<b><u>DIMETHYL FUMARATE - TECFIDERA</u></b>						
N 204063 002	6509376 7320999 7619001 7803840 8399514 8524773 8759393	Oct 29, 2019 May 18, 2020 Apr 01, 2018 Apr 01, 2018 Feb 07, 2028 Apr 01, 2018 Oct 29, 2019		DP U-1384 U-1384 U-1385 U-1384 U-1384 DP	NCE	Mar 27, 2018
<b><u>DIPHENHYDRAMINE CITRATE; IBUPROFEN - ADVIL PM</u></b>						
N 021394 001	8263647	May 30, 2022		DP		
<b><u>DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN - ADVIL PM</u></b>						
N 021393 001	8883849 9155718	Jan 17, 2022 Jan 17, 2022		U-1618 DP		
<b><u>DIVALPROEX SODIUM - DEPAKOTE ER</u></b>						
N 021168 001	6419953 6511678 6528090 6528091 6713086 6720004	Dec 18, 2018 Dec 18, 2018 Dec 18, 2018 Dec 18, 2018 Dec 18, 2018 Dec 18, 2018		DP U-106 DP U-579 DP		
<b><u>DIVALPROEX SODIUM - DEPAKOTE ER</u></b>						
N 021168 002	6511678 6528090 6713086 6720004	Dec 18, 2018 Dec 18, 2018 Dec 18, 2018 Dec 18, 2018		DP DP U-579 DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DIVALPROEX SODIUM - DEPAKOTE ER</b>						
N 021168 002	6511678	Dec 18, 2018				
	6528090	Dec 18, 2018	DP			
	6713086	Dec 18, 2018	DP U-579			
	6720004	Dec 18, 2018	DP			
<b>DOCETAXEL - DOCETAXEL</b>						
N 205934 001	8940786	Sep 30, 2033	DP U-1789			
	9308195	Sep 30, 2033	DP			
<b>DOCETAXEL - DOCETAXEL</b>						
N 205934 002	8940786	Sep 30, 2033	DP U-1789			
	9308195	Sep 30, 2033	DP			
<b>DOCETAXEL - DOCETAXEL</b>						
N 205934 003	8940786	Sep 30, 2033	DP U-1789			
	9308195	Sep 30, 2033	DP			
<b>DOFETILIDE - TIKOSYN</b>						
N 020931 001	6124363	Oct 09, 2018				
<b>DOFETILIDE - TIKOSYN</b>						
N 020931 002	6124363	Oct 09, 2018				
<b>DOFETILIDE - TIKOSYN</b>						
N 020931 003	6124363	Oct 09, 2018				
<b>DOLUTEGRAVIR SODIUM - TIVICAY</b>						
N 204790 001	8129385	Oct 05, 2027	DS DP		I-758	Nov 21, 2020
	9242986	Dec 08, 2029	DS DP		M-166	Jul 30, 2018
					NCE	Aug 12, 2018
<b>DOLUTEGRAVIR SODIUM - TIVICAY</b>						
N 204790 002	8129385	Oct 05, 2027	DS DP		I-758	Nov 21, 2020
	9242986	Dec 08, 2029	DS DP		NCE	Aug 12, 2018
<b>DOLUTEGRAVIR SODIUM - TIVICAY</b>						
N 204790 003	8129385	Oct 05, 2027	DS DP		I-758	Nov 21, 2020
	9242986	Dec 08, 2029	DS DP		NCE	Aug 12, 2018
<b>DOLUTEGRAVIR SODIUM; RILPIVIRINE HYDROCHLORIDE - JULUCA</b>						
N 210192 001	6838464	Feb 26, 2021	DS DP		NC	Nov 21, 2020
	7067522	Dec 20, 2019	DS DP		NCE	Aug 12, 2018
	7125879	Apr 21, 2025	DS DP U-257			
	8080551	Apr 11, 2023	DS DP			
	8101629	Aug 09, 2022	DP			
	8129385	Oct 05, 2027	DS DP			
	9242986	Dec 08, 2029	DS DP			
<b>DONEPEZIL HYDROCHLORIDE - ARICEPT</b>						
N 022568 001	8481565	Oct 04, 2026	DP			
<b>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</b>						
N 206439 001	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</b>						
N 206439 001	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<b>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</b>						
N 206439 002	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<b>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</b>						
N 206439 003	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<b>DONEPEZIL HYDROCHLORIDE; MEMANTINE HYDROCHLORIDE - NAMZARIC</b>						
N 206439 004	8039009	Mar 24, 2029	U-1641			
	8039009*PED	Sep 24, 2029				
	8058291	Dec 05, 2029	U-1641			
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-1641			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-1641			
	8283379*PED	May 22, 2026				
	8293794	Nov 22, 2025	DP			
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8338485	Nov 22, 2025	DP			
	8338486	Nov 22, 2025	U-1641			
	8362085	Nov 22, 2025	U-1641			
	8362085*PED	May 22, 2026				
	8580858	Nov 22, 2025	U-1641			
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DORIPENEM - DORIBAX</u></b>						
N 022106 001	8247402	Mar 30, 2021	DS DP			
<b><u>DORIPENEM - DORIBAX</u></b>						
N 022106 002	8247402	Mar 30, 2021	DS DP			
<b><u>DOXEPIN HYDROCHLORIDE - SILENOR</u></b>						
N 022036 001	6211229 7915307 8513299 9107898 9486437 9532971 9572814	Feb 17, 2020 Aug 24, 2027 Sep 07, 2030 May 01, 2028 May 18, 2027 Jun 01, 2029 Jul 20, 2027	U-620 U-620 U-620 U-620 U-620 DP U-620			
<b><u>DOXEPIN HYDROCHLORIDE - SILENOR</u></b>						
N 022036 002	6211229 7915307 8513299 9107898 9486437 9532971 9572814	Feb 17, 2020 Aug 24, 2027 Sep 07, 2030 May 01, 2028 May 18, 2027 Jun 01, 2029 Jul 20, 2027	U-620 U-620 U-620 U-620 U-620 DP U-620			
<b><u>DOXYCYCLINE - ORACEA</u></b>						
N 050805 001	7211267 7232572 7749532 8206740 8394405 8394406 8470364 8603506 8709478 9241946	Apr 05, 2022 Apr 05, 2022 Dec 19, 2027 Dec 24, 2025 Apr 07, 2024 Apr 07, 2024 Apr 07, 2024 Apr 05, 2022 Apr 07, 2024 Apr 05, 2022	U-925 U-925 DP U-1063 DP U-925 DP U-925 DP U-925 DP U-925 U-1063 U-1063 U-1063			
<b><u>DOXYCYCLINE HYCLATE - DORYX</u></b>						
N 050795 001	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX</u></b>						
N 050795 002	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX</u></b>						
N 050795 003	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX</u></b>						
N 050795 004	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX</u></b>						
N 050795 005	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX</u></b>						
N 050795 006	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX MPC</u></b>						
N 050795 007	6958161 8715724 9295652 9446057 9511031	Dec 15, 2022 Feb 03, 2028 Oct 23, 2034 Dec 23, 2034 Oct 23, 2034	DP U-918 DP DP U-918 DP U-918 DP			
<b><u>DOXYCYCLINE HYCLATE - DORYX MPC</u></b>						
N 050795 008	6958161 8715724	Dec 15, 2022 Feb 03, 2028	DP U-918 DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>DOXYCYCLINE HYCLATE - DORYX MPC</u></b>						
N 050795 008	9295652	Oct 23, 2034	DP U-918			
	9446057	Dec 23, 2034	DP U-918			
	9511031	Oct 23, 2034	DP			
<b><u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - DICLEGIS</u></b>						
N 021876 001	6340695	Jun 21, 2021	DP U-1382			
	7560122	Jan 25, 2019	DP			
<b><u>DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE - BONJESTA</u></b>						
N 209661 001	7560122	Jan 25, 2019	DP			
	9089489	Feb 18, 2033	DP U-1382			
	9375404	Feb 18, 2033	DP U-1382			
	9526703	Feb 18, 2033	DP U-1382			
<b><u>DRONABINOL - SYNDROS</u></b>						
N 205525 001	8222292	Aug 06, 2028	DS DP			
	9345771	Aug 06, 2028	DS DP			
<b><u>DRONEDARONE HYDROCHLORIDE - MULTAQ</u></b>						
N 022425 001	7323493	Jun 19, 2018	DP			
	8318800	Jun 19, 2018	DP			
	8410167	Apr 16, 2029	U-1387			
	8410167	Apr 16, 2029	U-1388			
	8602215	Jun 30, 2031	U-1473			
	9107900	Apr 16, 2029	U-1726			
	9107900	Apr 16, 2029	U-1728			
<b><u>DROSPIRENONE; ESTRADIOL - ANGELIQ</u></b>						
N 021355 001	8906890	Oct 22, 2031	DP			
<b><u>DROSPIRENONE; ESTRADIOL - ANGELIQ</u></b>						
N 021355 002	6933395	Aug 11, 2017	DS			
<b><u>DROSPIRENONE; ETHINYL ESTRADIOL - YASMIN</u></b>						
N 021098 001	6787531	Aug 31, 2020	DP			
	6933395	Aug 11, 2017	DS			
<b><u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u></b>						
N 021676 001	6787531	Aug 31, 2020	DP			
	6933395	Aug 11, 2017	DP			
	6958326	Dec 20, 2021	DP			
	6987101	Dec 22, 2017	U-758			
	7163931	Dec 20, 2021	U-1			
<b><u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLEATE CALCIUM - BEYAZ</u></b>						
N 022532 001	6441168	Jul 30, 2022	DS			
	6958326	Dec 20, 2021	DP			
	7163931	Mar 03, 2022	U-1			
	8617597	Feb 08, 2030	DP			
<b><u>DROSPIRENONE; ETHINYL ESTRADIOL; LEVOMEFOLEATE CALCIUM - SAFYRAL</u></b>						
N 022574 001	6441168	Apr 17, 2020	DS			
	6958326	Dec 20, 2021	DP			
	7163931	Mar 03, 2022	U-1			
	8617597	Feb 08, 2030	DP			
<b><u>DROXIDOPA - NORTHERA</u></b>						
N 203202 001				NCE	Feb 18, 2019	
				ODE-61	Feb 18, 2021	
<b><u>DROXIDOPA - NORTHERA</u></b>						
N 203202 002				NCE	Feb 18, 2019	
				ODE-61	Feb 18, 2021	
<b><u>DROXIDOPA - NORTHERA</u></b>						
N 203202 003				NCE	Feb 18, 2019	
				ODE-61	Feb 18, 2021	

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>DULOXETINE HYDROCHLORIDE - CYMBALTA</b>						
N 021427 001	6596756	Sep 10, 2019	U-882		NPP	Oct 16, 2017
					NPP	Oct 16, 2017
<b>DULOXETINE HYDROCHLORIDE - CYMBALTA</b>						
N 021427 002	6596756	Sep 10, 2019	U-882		NPP	Oct 16, 2017
					NPP	Oct 16, 2017
<b>DULOXETINE HYDROCHLORIDE - CYMBALTA</b>						
N 021427 004	6596756	Sep 10, 2019	U-882		NPP	Oct 16, 2017
					NPP	Oct 16, 2017
<b>ECONAZOLE NITRATE - ECOZA</b>						
N 205175 001	5993830	Jan 16, 2018	DP U-1449			
<b>EDARAVONE - RADICAVA</b>						
N 209176 001	6933310	Nov 13, 2020	U-2013		NCE	May 05, 2022
					ODE-144	May 05, 2024
<b>EDOXABAN TOSYLATE - SAVAYSA</b>						
N 206316 001	7365205	Jun 12, 2023	DS		NCE	Jan 08, 2020
	9149532	Mar 28, 2028	DP			
<b>EDOXABAN TOSYLATE - SAVAYSA</b>						
N 206316 002	7365205	Jun 12, 2023	DS		NCE	Jan 08, 2020
	9149532	Mar 28, 2028	DP			
<b>EDOXABAN TOSYLATE - SAVAYSA</b>						
N 206316 003	7365205	Jun 12, 2023	DS		NCE	Jan 08, 2020
	9149532	Mar 28, 2028	DP			
<b>EFAVIRENZ - SUSTIVA</b>						
N 020972 001	6238695	Apr 06, 2019	DP			
	6238695*PED	Oct 06, 2019				
	6555133	Apr 06, 2019	U-248			
	6555133*PED	Oct 06, 2019				
	6639071	Feb 14, 2018	DS			
	6639071*PED	Aug 14, 2018				
	6939964	Jan 20, 2018	DS			
	6939964*PED	Jul 20, 2018				
<b>EFAVIRENZ - SUSTIVA</b>						
N 020972 002	6238695	Apr 06, 2019	DP			
	6238695*PED	Oct 06, 2019				
	6555133	Apr 06, 2019	U-248			
	6555133*PED	Oct 06, 2019				
	6639071	Feb 14, 2018	DS			
	6639071*PED	Aug 14, 2018				
	6939964	Jan 20, 2018	DS			
	6939964*PED	Jul 20, 2018				
<b>EFAVIRENZ - SUSTIVA</b>						
N 020972 003	6238695	Apr 06, 2019	DP			
	6238695*PED	Oct 06, 2019				
	6555133	Apr 06, 2019	U-248			
	6555133*PED	Oct 06, 2019				
	6639071	Feb 14, 2018	DS			
	6639071*PED	Aug 14, 2018				
	6939964	Jan 20, 2018	DS			
	6939964*PED	Jul 20, 2018				
<b>EFAVIRENZ - SUSTIVA</b>						
N 021360 001	6639071	Feb 14, 2018	DS			
	6639071*PED	Aug 14, 2018				
	6939964	Jan 20, 2018	DS			
	6939964*PED	Jul 20, 2018				
<b>EFAVIRENZ - SUSTIVA</b>						
N 021360 002	6639071	Feb 14, 2018	DS			
	6639071*PED	Aug 14, 2018				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EFAVIRENZ - SUSTIVA</u></b>						
N 021360	002	6939964	Jan 20, 2018	DS		
		6939964*PED	Jul 20, 2018			
<b><u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u></b>						
N 021937	001	5914331	Jul 02, 2017	DS		
		5922695	Jul 25, 2017	DS U-1170		
		5922695	Jul 25, 2017	DS U-750		
		5935946	Jul 25, 2017	DS DP U-1170		
		5935946	Jul 25, 2017	DS DP U-750		
		5977089	Jul 25, 2017	DS DP U-1170		
		5977089	Jul 25, 2017	DS DP U-750		
		6043230	Jul 25, 2017	U-1170		
		6043230	Jul 25, 2017	U-750		
		6639071	Feb 14, 2018	DS		
		6639071*PED	Aug 14, 2018			
		6642245	Nov 04, 2020	U-1170		
		6642245	Nov 04, 2020	U-750		
		6703396	Mar 09, 2021	DS DP		
		6939964	Jan 20, 2018	DS		
		6939964*PED	Jul 20, 2018			
		8592397	Jan 13, 2024	DP U-1170		
		8592397	Jan 13, 2024	DP U-750		
		8598185	Apr 28, 2029	DP		
		8716264	Jan 13, 2024	DP U-257		
		9018192	Jun 13, 2026	U-1170		
		9018192	Jun 13, 2026	U-750		
		9457036	Jan 13, 2024	DP U-257		
		9545414	Jun 13, 2026	DP U-1170		
		9545414	Jun 13, 2026	DP U-750		
		9744181	Jan 13, 2024	DP U-257		
<b><u>EFINACONAZOLE - JUBLIA</u></b>						
N 203567	001	7214506	Oct 05, 2021	U-281		
		8039494	Jul 08, 2030	U-281		
		8486978	Oct 24, 2030	DP		
		9302009	Oct 24, 2030	DP		
		9566272	Jan 03, 2028	U-1969		
		9662394	Oct 02, 2034	DP		
		9861698	Jul 08, 2030	DP		
<b><u>ELBASVIR; GRAZOPREVIR - ZEPATIER</u></b>						
N 208261	001	7973040	Jul 24, 2029	DS DP U-1813		
		8871759	May 04, 2031	DS DP U-1813		
<b><u>ELETRIPTAN HYDROBROMIDE - RELPAX</u></b>						
N 021016	001	6110940	Aug 29, 2017			
<b><u>ELETRIPTAN HYDROBROMIDE - RELPAX</u></b>						
N 021016	002	6110940	Aug 29, 2017			
<b><u>ELIGLUSTAT TARTRATE - CERDELGA</u></b>						
N 205494	001	6916802	Apr 29, 2022	U-1571		
		7196205	Apr 29, 2022	DS		
		7615573	Apr 29, 2022	U-1571		
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291	001	6280959	Oct 30, 2018	DS DP U-1306		
		6280959	Oct 30, 2018	DS DP U-1575		
		6280959	Oct 30, 2018	DS DP U-1714		
		6280959	Oct 30, 2018	DS DP U-930		
		6280959*PED	Apr 30, 2019			
		7160870	Nov 20, 2022	DS DP U-1306		
		7160870	Nov 20, 2022	DS DP U-1575		
		7160870	Nov 20, 2022	DS DP U-1714		
		7160870	Nov 20, 2022	DS DP U-930		
		7160870*PED	May 20, 2023			
		7332481	May 24, 2021	U-1306		
		7332481	May 24, 2021	U-1575		
		7332481	May 24, 2021	U-1714		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 001	7332481	May 24, 2021	U-930			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP U-1714			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1575			
	7473686	May 24, 2021	DS DP U-1714			
	7473686	May 24, 2021	DS DP U-930			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1714			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1575			
	7790704	May 24, 2021	U-1714			
	7790704	May 24, 2021	U-930			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1575			
	7795293	May 21, 2023	U-1714			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	8052993	Aug 01, 2027	DP U-1306			
	8052993	Aug 01, 2027	DP U-1575			
	8052993	Aug 01, 2027	DP U-1714			
	8052993	Aug 01, 2027	DP U-930			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-1619			
	8828430	Aug 01, 2027	DP U-1714			
	8828430*PED	Feb 01, 2028				
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 002	6280959	Oct 30, 2018	DS DP U-1306		D-149	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1575		I-711	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1714		ODE-75	Aug 26, 2021
	6280959	Oct 30, 2018	DS DP U-930		PED	Dec 11, 2018
	6280959*PED	Apr 30, 2019			PED	Dec 11, 2018
	7160870	Nov 20, 2022	DS DP U-1306		PED	Feb 26, 2022
	7160870	Nov 20, 2022	DS DP U-1575			
	7160870	Nov 20, 2022	DS DP U-1714			
	7160870	Nov 20, 2022	DS DP U-930			
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-1575			
	7332481	May 24, 2021	U-1714			
	7332481	May 24, 2021	U-930			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP U-1714			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1575			
	7473686	May 24, 2021	DS DP U-1714			
	7473686	May 24, 2021	DS DP U-930			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1714			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 002	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1575			
	7790704	May 24, 2021	U-1714			
	7790704	May 24, 2021	U-930			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1575			
	7795293	May 21, 2023	U-1714			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1306			
	8052994	Aug 01, 2027	DP U-1575			
	8052994	Aug 01, 2027	DP U-1714			
	8052994	Aug 01, 2027	DP U-930			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-1619			
	8828430	Aug 01, 2027	DP U-1714			
	8828430*PED	Feb 01, 2028				
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 003	6280959	Oct 30, 2018	DS DP U-1306		D-149	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1575		I-711	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1714		ODE-75	Aug 26, 2021
	6280959	Oct 30, 2018	DS DP U-930		PED	Dec 11, 2018
	6280959*PED	Apr 30, 2019			PED	Dec 11, 2018
	7160870	Nov 20, 2022	DS DP U-1306		PED	Feb 26, 2022
	7160870	Nov 20, 2022	DS DP U-1575			
	7160870	Nov 20, 2022	DS DP U-1714			
	7160870	Nov 20, 2022	DS DP U-930			
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-1575			
	7332481	May 24, 2021	U-1714			
	7332481	May 24, 2021	U-930			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP U-1714			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1575			
	7473686	May 24, 2021	DS DP U-1714			
	7473686	May 24, 2021	DS DP U-930			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1714			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1575			
	7790704	May 24, 2021	U-1714			
	7790704	May 24, 2021	U-930			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1575			
	7795293	May 21, 2023	U-1714			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 003	8062665	Aug 01, 2027	DP U-1306			
	8062665	Aug 01, 2027	DP U-1575			
	8062665	Aug 01, 2027	DP U-1714			
	8062665	Aug 01, 2027	DP U-930			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1714			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-1619			
	8828430	Aug 01, 2027	DP U-1714			
	8828430*PED	Feb 01, 2028				
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 004	6280959	Oct 30, 2018	DS DP U-1306		D-149	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1575		I-711	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1714		ODE-75	Aug 26, 2021
	6280959	Oct 30, 2018	DS DP U-930		PED	Dec 11, 2018
	6280959*PED	Apr 30, 2019			PED	Dec 11, 2018
	7160870	Nov 20, 2022	DS DP U-1306		PED	Feb 26, 2022
	7160870	Nov 20, 2022	DS DP U-1575			
	7160870	Nov 20, 2022	DS DP U-1714			
	7160870	Nov 20, 2022	DS DP U-930			
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-1575			
	7332481	May 24, 2021	U-1714			
	7332481	May 24, 2021	U-930			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP U-1714			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1575			
	7473686	May 24, 2021	DS DP U-1714			
	7473686	May 24, 2021	DS DP U-930			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-1714			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1575			
	7790704	May 24, 2021	U-1714			
	7790704	May 24, 2021	U-930			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1575			
	7795293	May 21, 2023	U-1714			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	8052993	Aug 01, 2027	DP U-1714			
	8052993*PED	Feb 01, 2028				
	8052994	Aug 01, 2027	DP U-1714			
	8052994*PED	Feb 01, 2028				
	8062665	Aug 01, 2027	DP U-1714			
	8062665*PED	Feb 01, 2028				
	8071129	Aug 01, 2027	DP U-1306			
	8071129	Aug 01, 2027	DP U-1575			
	8071129	Aug 01, 2027	DP U-1714			
	8071129	Aug 01, 2027	DP U-930			
	8071129*PED	Feb 01, 2028				
	8828430	Aug 01, 2027	DP U-1306			
	8828430	Aug 01, 2027	DP U-1619			
	8828430	Aug 01, 2027	DP U-1714			
	8828430*PED	Feb 01, 2028				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 022291 005	6280959	Oct 30, 2018	DS DP U-1306		D-149	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1575		I-711	Jun 11, 2018
	6280959	Oct 30, 2018	DS DP U-1714		ODE-75	Aug 26, 2021
	6280959	Oct 30, 2018	DS DP U-930		PED	Dec 11, 2018
	6280959*PED	Apr 30, 2019			PED	Dec 11, 2018
	7160870	Nov 20, 2022	DS DP U-1306		PED	Feb 26, 2022
	7160870	Nov 20, 2022	DS DP U-1575			
	7160870	Nov 20, 2022	DS DP U-1714			
	7160870	Nov 20, 2022	DS DP U-930			
	7160870*PED	May 20, 2023				
	7332481	May 24, 2021	U-1306			
	7332481	May 24, 2021	U-1575			
	7332481	May 24, 2021	U-1714			
	7332481	May 24, 2021	U-930			
	7332481*PED	Nov 24, 2021				
	7452874	May 24, 2021	DS DP U-1714			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1306			
	7473686	May 24, 2021	DS DP U-1575			
	7473686	May 24, 2021	DS DP U-1714			
	7473686	May 24, 2021	DS DP U-930			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1306			
	7547719	Jul 13, 2025	DS DP U-1575			
	7547719	Jul 13, 2025	DS DP U-930			
	7547719*PED	Jan 13, 2026				
	7790704	May 24, 2021	U-1306			
	7790704	May 24, 2021	U-1575			
	7790704	May 24, 2021	U-930			
	7790704*PED	Nov 24, 2021				
	7795293	May 21, 2023	U-1306			
	7795293	May 21, 2023	U-1575			
	7795293	May 21, 2023	U-930			
	7795293*PED	Nov 21, 2023				
	8052995	Aug 01, 2027	DP U-1306			
	8052995	Aug 01, 2027	DP U-1575			
	8052995*PED	Feb 01, 2028				
<b><u>ELTROMBOPAG OLAMINE - PROMACTA</u></b>						
N 207027 001	6280959	Oct 30, 2018	DS DP U-1736		D-149	Jun 11, 2018
	6280959*PED	Apr 30, 2019			I-711	Jun 11, 2018
	7160870	Nov 20, 2022	DS DP U-1736		ODE-74	Aug 26, 2021
	7160870*PED	May 20, 2023			PED	Dec 11, 2018
	7332481	May 24, 2021	U-1736		PED	Dec 11, 2018
	7332481*PED	Nov 24, 2021			PED	Feb 26, 2022
	7452874	May 24, 2021	DS DP			
	7452874*PED	Nov 24, 2021				
	7473686	May 24, 2021	DS DP U-1736			
	7473686*PED	Nov 24, 2021				
	7547719	Jul 13, 2025	DS DP U-1736			
	7547719*PED	Jan 13, 2026				
	7790704	May 24, 2021	U-1736			
	7790704*PED	Nov 24, 2021				
	7795293	May 24, 2023	U-1736			
	7795293*PED	Nov 24, 2023				
<b><u>ELUXADOLINE - VIBERZI</u></b>						
N 206940 001	7741356	Mar 25, 2028	DS DP		NCE	May 27, 2020
	7786158	Mar 14, 2025	DS			
	8344011	Mar 14, 2025		U-1709		
	8609709	Mar 14, 2025	DS			
	8691860	Jul 07, 2028	DS	U-1709		
	8772325	Mar 14, 2025		U-1709		
	9115091	Jul 07, 2028	DS DP	U-1738		
	9205076	Mar 14, 2025		U-1709		
	9364489	Jul 07, 2028		U-1709		
	9675587	Mar 14, 2033	DP			
	9700542	Mar 14, 2025	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ELUXADOLINE - VIBERZI</u></b>						
N 206940 001	9789125	Jul 07, 2028	DP U-1709			
	9789125	Jul 07, 2028	DP U-2152			
<b><u>ELUXADOLINE - VIBERZI</u></b>						
N 206940 002	7741356	Mar 25, 2028	DS DP		NCE	May 27, 2020
	7786158	Mar 14, 2025	DS			
	8344011	Mar 14, 2025		U-1709		
	8609709	Mar 14, 2025	DS			
	8691860	Jul 07, 2028	DS	U-1709		
	8772325	Mar 14, 2025		U-1709		
	9115091	Jul 07, 2028	DS DP	U-1738		
	9205076	Mar 14, 2025		U-1709		
	9364489	Jul 07, 2028		U-1709		
	9675587	Mar 14, 2033	DP			
	9700542	Mar 14, 2025	DP			
	9789125	Jul 07, 2028	DP	U-1709		
	9789125	Jul 07, 2028	DP	U-2152		
<b><u>ELVITEGRAVIR - VITEKTA</u></b>						
N 203093 001	7176220	Aug 27, 2026	DS DP	U-257	NCE	Aug 27, 2017
	7635704	Oct 26, 2026	DS DP	U-257	NP	Sep 24, 2017
	8981103	Oct 26, 2026	DS DP			
<b><u>ELVITEGRAVIR - VITEKTA</u></b>						
N 203093 002	7176220	Aug 27, 2026	DS DP	U-257	NCE	Aug 27, 2017
	7635704	Oct 26, 2026	DS DP	U-257	NP	Sep 24, 2017
	8981103	Oct 26, 2026	DS DP			
<b><u>EMPAGLIFLOZIN - JARDIANC</u></b>						
N 204629 001	7579449	Nov 05, 2025	DS		I-739	Dec 02, 2019
	7713938	Apr 15, 2027	DS DP		M-160	Jun 26, 2018
	8551957	Oct 14, 2029		U-1651	M-161	Jun 26, 2018
					M-174	Mar 18, 2019
					NCE	Aug 01, 2019
<b><u>EMPAGLIFLOZIN - JARDIANC</u></b>						
N 204629 002	7579449	Nov 05, 2025	DS		I-739	Dec 02, 2019
	7713938	Apr 15, 2027	DS DP		M-160	Jun 26, 2018
	8551957	Oct 14, 2029		U-1651	M-161	Jun 26, 2018
					M-174	Mar 18, 2019
					NCE	Aug 01, 2019
<b><u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u></b>						
N 206073 001	6890898	Feb 02, 2019		U-1652	I-739	Dec 02, 2019
	7078381	Feb 02, 2019		U-1651	NC	Jan 30, 2018
	7407955	May 02, 2025	DS DP		NCE	Aug 01, 2019
	7459428	Feb 02, 2019		U-1651		
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023		U-1651		
	8178541	Aug 12, 2023	DP	U-1653		
	8178541	Aug 12, 2023	DP	U-1654		
	8551957	Oct 14, 2029	DP	U-1651		
	8673927	May 04, 2027		U-1652		
	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP	U-1772		
<b><u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u></b>						
N 206073 002	6890898	Feb 02, 2019		U-1652	I-739	Dec 02, 2019
	7078381	Feb 02, 2019		U-1651	NC	Jan 30, 2018
	7407955	May 02, 2025	DS DP		NCE	Aug 01, 2019
	7459428	Feb 02, 2019		U-1651		
	7579449	Nov 05, 2025	DS			
	7713938	Apr 15, 2027	DS DP			
	8119648	Aug 12, 2023		U-1651		
	8178541	Aug 12, 2023	DP	U-1653		
	8178541	Aug 12, 2023	DP	U-1654		
	8551957	Oct 14, 2029	DP	U-1651		
	8673927	May 04, 2027		U-1652		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EMPAGLIFLOZIN; LINAGLIPTIN - GLYXAMBI</u></b>						
N 206073 002	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP U-1772			
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u></b>						
N 206111 001	7579449	Nov 05, 2025	DS	I-739	Dec 02, 2019	
	7713938	Apr 15, 2027	DS DP	M-174	Mar 18, 2019	
				NCE	Aug 01, 2019	
				NP	Aug 26, 2018	
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u></b>						
N 206111 002	7579449	Nov 05, 2025	DS	I-739	Dec 02, 2019	
	7713938	Apr 15, 2027	DS DP	M-174	Mar 18, 2019	
				NCE	Aug 01, 2019	
				NP	Aug 26, 2018	
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u></b>						
N 206111 003	7579449	Nov 05, 2025	DS	I-739	Dec 02, 2019	
	7713938	Apr 15, 2027	DS DP	M-174	Mar 18, 2019	
				NCE	Aug 01, 2019	
				NP	Aug 26, 2018	
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY</u></b>						
N 206111 004	7579449	Nov 05, 2025	DS	I-739	Dec 02, 2019	
	7713938	Apr 15, 2027	DS DP	M-174	Mar 18, 2019	
				NCE	Aug 01, 2019	
				NP	Aug 26, 2018	
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u></b>						
N 208658 001	6488962	Jun 20, 2020	DP	I-739	Dec 02, 2019	
	7579449	Nov 05, 2025	DS	NCE	Aug 01, 2019	
	7713938	Apr 15, 2027	DS DP			
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u></b>						
N 208658 002	6488962	Jun 20, 2020	DP	I-739	Dec 02, 2019	
	7579449	Nov 05, 2025	DS	NCE	Aug 01, 2019	
	7713938	Apr 15, 2027	DS DP			
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u></b>						
N 208658 003	6488962	Jun 20, 2020	DP	I-739	Dec 02, 2019	
	7579449	Nov 05, 2025	DS	NCE	Aug 01, 2019	
	7713938	Apr 15, 2027	DS DP			
<b><u>EMPAGLIFLOZIN; METFORMIN HYDROCHLORIDE - SYNJARDY XR</u></b>						
N 208658 004	6488962	Jun 20, 2020	DP	I-739	Dec 02, 2019	
	7579449	Nov 05, 2025	DS	NCE	Aug 01, 2019	
	7713938	Apr 15, 2027	DS DP			
<b><u>EMTRICITABINE - EMTRIVA</u></b>						
N 021500 001	5914331	Jul 02, 2017	DS			
	6642245	Nov 04, 2020	U-257			
	6642245	Nov 04, 2020	U-541			
	6703396	Mar 09, 2021	DS DP			
<b><u>EMTRICITABINE - EMTRIVA</u></b>						
N 021896 001	5914331	Jul 02, 2017	DS			
	6642245	Nov 04, 2020	U-257			
	6703396	Mar 09, 2021	DS DP			
<b><u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u></b>						
N 208351 001	5914331	Jul 02, 2017	DS	M-206	Aug 21, 2020	
	5914331*PED	Jan 02, 2018		M-207	Aug 21, 2020	
	6642245	Nov 04, 2020	U-257	NCE	Nov 05, 2020	
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
	6838464	Feb 26, 2021	DS DP			
	7067522	Dec 20, 2019	DS DP			
	7125879	Apr 21, 2025	DS DP U-257			
	7390791	May 07, 2022	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR ALAFENAMIDE FUMARATE - ODEFSEY</u></b>						
N 208351 001	7803788	Feb 02, 2022		U-257		
	8080551	Apr 11, 2023	DS DP			
	8101629	Aug 09, 2022	DP			
	8754065	Aug 15, 2032	DS DP U-257			
	9296769	Aug 15, 2032	DS DP U-257			
<b><u>EMTRICITABINE; RILPIVIRINE HYDROCHLORIDE; TENOFOVIR DISOPROXIL FUMARATE - COMPLERA</u></b>						
N 202123 001	5914331	Jul 02, 2017	DS			
	5922695	Jul 25, 2017	DS U-257			
	5935946	Jul 25, 2017	DS DP U-257			
	5977089	Jul 25, 2017	DS DP U-257			
	6043230	Jul 25, 2017	U-257			
	6642245	Nov 04, 2020	U-257			
	6703396	Mar 09, 2021	DS DP			
	6838464	Feb 26, 2021	DS DP			
	7067522	Dec 20, 2019	DS DP			
	7125879	Apr 21, 2025	DS DP U-257			
	8080551	Apr 11, 2023	DS DP			
	8101629	Aug 09, 2022	DP			
	8592397	Jan 13, 2024	DP U-257			
	8716264	Jan 13, 2024	DP U-257			
	8841310	Dec 09, 2025	DP U-257			
	9457036	Jan 13, 2024	DP U-257			
	9744181	Jan 13, 2024	DP U-257			
<b><u>EMTRICITABINE; TENOFOVIR ALAFENAMIDE FUMARATE - DESCovy</u></b>						
N 208215 001	5914331	Jul 02, 2017	DS		NCE	Nov 05, 2020
	5914331*PED	Jan 02, 2018			NPP	Sep 25, 2020
	6642245	Nov 04, 2020	U-257			
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
	7390791	May 07, 2022	DS DP			
	7803788	Feb 02, 2022	U-257			
	8754065	Aug 15, 2032	DS DP U-257			
	9296769	Aug 15, 2032	DS DP U-257			
<b><u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u></b>						
N 021752 001	5914331	Jul 02, 2017	DS			
	5914331*PED	Jan 02, 2018				
	5922695	Jul 25, 2017	DS U-1170			
	5922695	Jul 25, 2017	DS U-1259			
	5922695	Jul 25, 2017	DS U-248			
	5922695	Jul 25, 2017	DS U-541			
	5935946	Jul 25, 2017	DS DP U-1170			
	5935946	Jul 25, 2017	DS DP U-1259			
	5935946	Jul 25, 2017	DS DP U-248			
	5935946	Jul 25, 2017	DS DP U-541			
	5977089	Jul 25, 2017	DS DP U-1170			
	5977089	Jul 25, 2017	DS DP U-1259			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089	Jul 25, 2017	DS DP U-541			
	6043230	Jul 25, 2017	U-1170			
	6043230	Jul 25, 2017	U-1259			
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-541			
	6043230*PED	Jan 25, 2018				
	6642245	Nov 04, 2020	U-1170			
	6642245	Nov 04, 2020	U-248			
	6642245	Nov 04, 2020	U-541			
	6703396	Mar 09, 2021	DS DP			
	8592397	Jan 13, 2024	DP U-1170			
	8592397	Jan 13, 2024	DP U-248			
	8592397	Jan 13, 2024	DP U-541			
	8716264	Jan 13, 2024	DP U-257			
	9457036	Jan 13, 2024	DP U-257			
	9744181	Jan 13, 2024	DP U-257			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u></b>						
N 021752 002	5914331	Jul 02, 2017	DS			
	5914331*PED	Jan 02, 2018				
	5922695	Jul 25, 2017	DS U-1170			
	5922695	Jul 25, 2017	DS U-1259			
	5922695	Jul 25, 2017	DS U-248			
	5922695	Jul 25, 2017	DS U-541			
	5935946	Jul 25, 2017	DS DP U-1170			
	5935946	Jul 25, 2017	DS DP U-1259			
	5935946	Jul 25, 2017	DS DP U-248			
	5935946	Jul 25, 2017	DS DP U-541			
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-1170			
	5977089	Jul 25, 2017	DS DP U-1259			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089	Jul 25, 2017	DS DP U-541			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-1170			
	6043230	Jul 25, 2017	U-1259			
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-541			
	6043230*PED	Jan 25, 2018				
	6642245	Nov 04, 2020	U-1170			
	6642245	Nov 04, 2020	U-248			
	6642245	Nov 04, 2020	U-541			
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
<b><u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u></b>						
N 021752 003	5914331	Jul 02, 2017	DS			
	5914331*PED	Jan 02, 2018				
	5922695	Jul 25, 2017	DS U-1170			
	5922695	Jul 25, 2017	DS U-1259			
	5922695	Jul 25, 2017	DS U-248			
	5922695	Jul 25, 2017	DS U-541			
	5922695*PED	Jan 25, 2018				
	5935946	Jul 25, 2017	DS DP U-1170			
	5935946	Jul 25, 2017	DS DP U-1259			
	5935946	Jul 25, 2017	DS DP U-248			
	5935946	Jul 25, 2017	DS DP U-541			
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-1170			
	5977089	Jul 25, 2017	DS DP U-1259			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089	Jul 25, 2017	DS DP U-541			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-1170			
	6043230	Jul 25, 2017	U-1259			
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-541			
	6043230*PED	Jan 25, 2018				
	6642245	Nov 04, 2020	U-1170			
	6642245	Nov 04, 2020	U-248			
	6642245	Nov 04, 2020	U-541			
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
<b><u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u></b>						
N 021752 004	5914331	Jul 02, 2017	DS			
	5914331*PED	Jan 02, 2018				
	5922695	Jul 25, 2017	DS U-1170			
	5922695	Jul 25, 2017	DS U-1259			
	5922695	Jul 25, 2017	DS U-248			
	5922695	Jul 25, 2017	DS U-541			
	5922695*PED	Jan 25, 2018				
	5935946	Jul 25, 2017	DS DP U-1170			
	5935946	Jul 25, 2017	DS DP U-1259			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u></b>						
N 021752 004	5935946	Jul 25, 2017	DS DP U-248			
	5935946	Jul 25, 2017	DS DP U-541			
	5935946*PED	Jan 25, 2018				
	5977089	Jul 25, 2017	DS DP U-1170			
	5977089	Jul 25, 2017	DS DP U-1259			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089	Jul 25, 2017	DS DP U-541			
	5977089*PED	Jan 25, 2018				
	6043230	Jul 25, 2017	U-1170			
	6043230	Jul 25, 2017	U-1259			
	6043230	Jul 25, 2017	U-248			
	6043230	Jul 25, 2017	U-541			
	6043230*PED	Jan 25, 2018				
	6642245	Nov 04, 2020	U-1170			
	6642245	Nov 04, 2020	U-248			
	6642245	Nov 04, 2020	U-541			
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS DP			
	6703396*PED	Sep 09, 2021				
<b><u>ENALAPRIL MALEATE - EPANED KIT</u></b>						
N 204308 001	8568747	Nov 06, 2032	DP			
	8778366	Nov 06, 2032	U-1723			
	8778366	Nov 06, 2032	U-185			
	8778366	Nov 06, 2032	U-1892			
	8778366	Nov 06, 2032	U-3			
	8778366	Nov 06, 2032	U-71			
	9855214	Nov 06, 2032	DP			
<b><u>ENALAPRIL MALEATE - EPANED</u></b>						
N 208686 001	9669008	Mar 25, 2036	DP			
	9808442	Mar 25, 2036	U-1723			
	9808442	Mar 25, 2036	U-185			
	9808442	Mar 25, 2036	U-1892			
	9808442	Mar 25, 2036	U-3			
	9808442	Mar 25, 2036	U-71			
<b><u>ENASIDENIB MESYLATE - IDHIFA</u></b>						
N 209606 001	9512107	Jan 07, 2033	DS DP U-2087		NCE	Aug 01, 2022
	9732062	Sep 16, 2034	DS		ODE-151	Aug 01, 2024
	9738625	Aug 01, 2034	DS			
<b><u>ENASIDENIB MESYLATE - IDHIFA</u></b>						
N 209606 002	9512107	Jan 07, 2033	DS DP U-2087		NCE	Aug 01, 2022
	9732062	Sep 16, 2034	DS		ODE-151	Aug 01, 2024
	9738625	Aug 01, 2034	DS			
<b><u>ENTACAPONE - COMTAN</u></b>						
N 020796 001	6599530	Sep 14, 2018	DP U-219			
<b><u>ENZALUTAMIDE - XTANDI</u></b>						
N 203415 001	7709517	Aug 13, 2027	DS DP		I-693	Sep 10, 2017
	8183274	Aug 24, 2026	U-1281		NCE	Aug 31, 2017
	8183274	Aug 24, 2026	U-1588			
	9126941	May 15, 2026	U-1588			
<b><u>EPINEPHRINE - EPIPEN</u></b>						
N 019430 001	7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
	8870827	Sep 11, 2025	DP			
	9586010	Sep 11, 2025	DP			
<b><u>EPINEPHRINE - EPIPEN JR.</u></b>						
N 019430 002	7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
	8870827	Sep 11, 2025	DP			
	9586010	Sep 11, 2025	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EPINEPHRINE - EPIPEN JR.</u></b>						
N 019430 002	7449012	Sep 11, 2025	DP			
	7794432	Sep 11, 2025	DP			
	8048035	Sep 11, 2025	DP			
	8870827	Sep 11, 2025	DP			
	9586010	Sep 11, 2025	DP			
<b><u>EPINEPHRINE - TWINJECT 0.3</u></b>						
N 020800 001	7297136	Jan 18, 2025	DP			
	7621891	Feb 04, 2025	DP			
<b><u>EPINEPHRINE - TWINJECT 0.15</u></b>						
N 020800 002	7297136	Jan 18, 2025	DP			
	7621891	Feb 04, 2025	DP			
<b><u>EPINEPHRINE - ADRENACCLICK</u></b>						
N 020800 003	7905352	Apr 12, 2027	DP			
<b><u>EPINEPHRINE - ADRENACCLICK</u></b>						
N 020800 004	7905352	Apr 12, 2027	DP			
<b><u>EPINEPHRINE - AUVI-Q</u></b>						
N 201739 001	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8920377	Nov 23, 2024	DP			
	8926594	Mar 31, 2026	DP			
	9056170	Nov 23, 2024	DP			
	9149579	Jul 19, 2025	U-1758			
	9238108	Feb 20, 2027	DP			
	9259539	Feb 01, 2026	DP			
	9278182	Feb 01, 2026	DP			
	9724471	May 23, 2027	DP U-2092			
	9737669	Nov 23, 2024	DP			
<b><u>EPINEPHRINE - AUVI-Q</u></b>						
N 201739 002	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8920377	Nov 23, 2024	DP			
	8926594	Mar 31, 2026	DP			
	9056170	Nov 23, 2024	DP			
	9149579	Jul 19, 2025	U-1758			
	9238108	Feb 20, 2027	DP			
	9259539	Feb 01, 2026	DP			
	9278182	Feb 01, 2026	DP			
	9724471	May 23, 2027	DP U-2092			
	9737669	Nov 23, 2024	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EPINEPHRINE - AUVI-Q</u></b>						
N 201739 002	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8920377	Nov 23, 2024	DP			
	8926594	Mar 31, 2026	DP			
	9056170	Nov 23, 2024	DP			
	9149579	Jul 19, 2025	U-1758			
	9238108	Feb 20, 2027	DP			
	9259539	Feb 01, 2026	DP			
	9278182	Feb 01, 2026	DP			
	9724471	May 23, 2027	DP	U-2092		
	9737669	Nov 23, 2024	DP			
<b><u>EPINEPHRINE - AUVI-Q</u></b>						
N 201739 003	7731686	Jun 01, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8920377	Nov 23, 2024	DP			
	8926594	Mar 31, 2026	DP			
	9056170	Nov 23, 2024	DP			
	9149579	Jul 19, 2025	U-1758			
	9238108	Feb 20, 2027	DP			
	9259539	Feb 01, 2026	DP			
	9278182	Feb 01, 2026	DP			
	9724471	May 23, 2027	DP	U-2092		
	9737669	Nov 23, 2024	DP			
	9833573	Nov 23, 2024	U-2172			
<b><u>EPINEPHRINE - ADRENALIN</u></b>						
N 204200 001	9119876	Mar 13, 2035	DP			
	9295657	Mar 13, 2035	U-1829			
<b><u>EPINEPHRINE - ADRENALIN</u></b>						
N 204640 001	9119876	Mar 13, 2035	DP			
	9295657	Mar 13, 2035	U-1829			
<b><u>EPINEPHRINE - EPINEPHRINE</u></b>						
N 205029 001	9283197	Aug 15, 2034	DP	U-1828		
	9283197	Aug 15, 2034	DP	U-1829		
	9283197	Aug 15, 2034	DP	U-1830		
<b><u>EPINEPHRINE; LIDOCAINE HYDROCHLORIDE - LIDOSITE TOPICAL SYSTEM KIT</u></b>						
N 021504 001	6629968	Jun 30, 2020	DS	DP		
	6635045	Jun 29, 2021	DS	DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>EPLERENONE - INSPRA</b>						
N 021437 001	6410054	Dec 08, 2019	U-3			
	6410054	Dec 08, 2019	U-537			
	6410524	Nov 05, 2019	U-467			
	6495165	Dec 08, 2019	U-3			
	6495165	Dec 08, 2019	U-537			
	6534093	Dec 08, 2019	U-3			
	6534093	Dec 08, 2019	U-537			
	6558707	Dec 08, 2019	DP U-537			
	6747020	Nov 05, 2019	U-587			
	7157101	Dec 08, 2019	DP U-664			
<b>EPLERENONE - INSPRA</b>						
N 021437 002	6410054	Dec 08, 2019	U-3			
	6410054	Dec 08, 2019	U-537			
	6410524	Nov 05, 2019	U-467			
	6495165	Dec 08, 2019	U-3			
	6495165	Dec 08, 2019	U-537			
	6534093	Dec 08, 2019	U-3			
	6534093	Dec 08, 2019	U-537			
	6558707	Dec 08, 2019	DP U-537			
	6747020	Nov 05, 2019	U-587			
	7157101	Dec 08, 2019	DP U-664			
<b>EPLERENONE - INSPRA</b>						
N 021437 003	6410054	Dec 08, 2019	U-3			
	6410054	Dec 08, 2019	U-537			
	6410524	Nov 05, 2019	U-467			
	6495165	Dec 08, 2019	U-3			
	6495165	Dec 08, 2019	U-537			
	6534093	Dec 08, 2019	U-3			
	6534093	Dec 08, 2019	U-537			
	6558707	Dec 08, 2019	DP U-537			
	6747020	Nov 05, 2019	U-587			
	7157101	Dec 08, 2019	DP U-664			
<b>EPOPROSTENOL SODIUM - VELETRI</b>						
N 022260 001	8318802	Mar 15, 2027	DP			
	8598227	Feb 02, 2027				
<b>EPOPROSTENOL SODIUM - VELETRI</b>						
N 022260 002	8318802	Mar 15, 2027	DP			
	8598227	Feb 02, 2027				
<b>ERIBULIN MESYLATE - HALAVEN</b>						
N 201532 001	6214865	Jul 20, 2023	DS		I-721	Jan 28, 2019
	6469182	Jun 16, 2019		U-1096	ODE-107	Jan 28, 2023
	6469182	Jun 16, 2019		U-1812		
	7470720	Jun 16, 2019	DP			
	8097648	Jan 22, 2021		U-1096		
<b>ERLOTINIB HYDROCHLORIDE - TARCEVA</b>						
N 021743 001	5747498	Nov 08, 2018	DS DP U-659		D-164	May 20, 2019
	5747498*PED	May 08, 2019			M-181	Jun 01, 2019
	6900221	Nov 09, 2020	DS DP U-1046		M-190	Oct 18, 2019
	6900221	Nov 09, 2020	DS DP U-1403			
	6900221	Nov 09, 2020	DS DP U-659			
	6900221	Nov 09, 2020	DS DP U-875			
	6900221*PED	May 09, 2021				
	7087613	Nov 09, 2020	U-1045			
	7087613	Nov 09, 2020	U-1403			
	7087613	Nov 09, 2020	U-659			
	7087613*PED	May 09, 2021				
	RE41065	Nov 08, 2018	DS DP			
	RE41065*PED	May 08, 2019				
<b>ERLOTINIB HYDROCHLORIDE - TARCEVA</b>						
N 021743 002	5747498	Nov 08, 2018	DS DP U-659		D-164	May 20, 2019
	5747498*PED	May 08, 2019			M-181	Jun 01, 2019
	6900221	Nov 09, 2020	DS DP U-1046		M-190	Oct 18, 2019

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u></b>						
N 021743 002	6900221	Nov 09, 2020	DS DP U-1403			
	6900221	Nov 09, 2020	DS DP U-659			
	6900221	Nov 09, 2020	DS DP U-875			
	6900221*PED	May 09, 2021				
	7087613	Nov 09, 2020	U-1045			
	7087613	Nov 09, 2020	U-1403			
	7087613	Nov 09, 2020	U-659			
	7087613*PED	May 09, 2021				
	RE41065	Nov 08, 2018	DS DP			
	RE41065*PED	May 08, 2019				
<b><u>ERLOTINIB HYDROCHLORIDE - TARCEVA</u></b>						
N 021743 003	5747498	Nov 08, 2018	DS DP U-659		D-164	May 20, 2019
	5747498*PED	May 08, 2019			M-181	Jun 01, 2019
	6900221	Nov 09, 2020	DS DP U-1046		M-190	Oct 18, 2019
	6900221	Nov 09, 2020	DS DP U-1403			
	6900221	Nov 09, 2020	DS DP U-659			
	6900221	Nov 09, 2020	DS DP U-875			
	6900221*PED	May 09, 2021				
	7087613	Nov 09, 2020	U-1045			
	7087613	Nov 09, 2020	U-1403			
	7087613	Nov 09, 2020	U-659			
	7087613*PED	May 09, 2021				
	RE41065	Nov 08, 2018	DS DP			
	RE41065*PED	May 08, 2019				
<b><u>ERTUGLIFLOZIN - STEGLATRO</u></b>						
N 209803 001					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN - STEGLATRO</u></b>						
N 209803 002					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u></b>						
N 209806 001					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u></b>						
N 209806 002					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u></b>						
N 209806 003					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN; METFORMIN HYDROCHLORIDE - SEGLUROMET</u></b>						
N 209806 004					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u></b>						
N 209805 001					NCE	Dec 19, 2022
<b><u>ERTUGLIFLOZIN; SITAGLIPTIN PHOSPHATE - STEGLUJAN</u></b>						
N 209805 002					NCE	Dec 19, 2022
<b><u>ESCITALOPRAM OXALATE - LEXAPRO</u></b>						
N 021323 001	6916941	Aug 12, 2022	DS DP			
	7420069	Aug 12, 2022	DP			
<b><u>ESCITALOPRAM OXALATE - LEXAPRO</u></b>						
N 021323 002	6916941	Aug 12, 2022	DS DP			
	7420069	Aug 12, 2022	DP			
<b><u>ESCITALOPRAM OXALATE - LEXAPRO</u></b>						
N 021323 003	6916941	Aug 12, 2022	DS DP			
	7420069	Aug 12, 2022	DP			
<b><u>ESLICARBAZEPINE ACETATE - APTIOM</u></b>						
N 022416 001	5753646	Jun 27, 2021	DS DP U-2041		D-150	Aug 27, 2018
	8372431	Apr 17, 2030	DP		I-715	Aug 27, 2018
	9206135	Apr 21, 2026	DS		NCE	Nov 08, 2018
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ESLICARBAZEPINE ACETATE - APTIOM</u></b>						
N 022416 001	9763954	Sep 13, 2028		U-2123		
<b><u>ESLICARBAZEPINE ACETATE - APTIOM</u></b>						
N 022416 002	5753646	Jun 27, 2021	DS DP U-2041		D-150	Aug 27, 2018
	8372431	Apr 17, 2030	DP		I-715	Aug 27, 2018
	9206135	Apr 21, 2026	DS		NCE	Nov 08, 2018
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<b><u>ESLICARBAZEPINE ACETATE - APTIOM</u></b>						
N 022416 003	5753646	Jun 27, 2021	DS DP U-2041		D-150	Aug 27, 2018
	8372431	Apr 17, 2030	DP		I-715	Aug 27, 2018
	9206135	Apr 21, 2026	DS		NCE	Nov 08, 2018
	9566244	Oct 23, 2028	DP			
	9643929	Apr 21, 2026	DP			
	9750747	Aug 24, 2032	U-2041			
	9750747	Aug 24, 2032	U-2121			
	9763954	Sep 13, 2028	U-2123			
<b><u>ESMOLOL HYDROCHLORIDE - REVIBLOC IN PLASTIC CONTAINER</u></b>						
N 019386 004	6310094	Jan 12, 2021			D-150	Aug 27, 2018
	6528540	Jan 12, 2021			I-715	Aug 27, 2018
<b><u>ESMOLOL HYDROCHLORIDE - REVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER</u></b>						
N 019386 005	6310094	Jan 12, 2021			NCE	Nov 08, 2018
	6528540	Jan 12, 2021				
<b><u>ESMOLOL HYDROCHLORIDE - REVIBLOC</u></b>						
N 019386 006	6310094	Jan 12, 2021				
	6528540	Jan 12, 2021				
<b><u>ESMOLOL HYDROCHLORIDE - REVIBLOC</u></b>						
N 019386 007	6310094	Jan 12, 2021				
	6528540	Jan 12, 2021				
<b><u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE IN PLASTIC CONTAINER</u></b>						
N 205703 001	6310094	Jan 12, 2021	DP			
	6528540	Jan 12, 2021	DP			
	8829054	Mar 15, 2033	DP			
	8835505	Mar 15, 2033	DP			
<b><u>ESMOLOL HYDROCHLORIDE - ESMOLOL HYDROCHLORIDE DOUBLE STRENGTH IN PLASTIC CONTAINER</u></b>						
N 205703 002	6310094	Jan 12, 2021	DP			
	6528540	Jan 12, 2021	DP			
	8829054	Mar 15, 2033	DP			
	8835505	Mar 15, 2033	DP			
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021153 001	6147103	Oct 09, 2018				
	6166213	Oct 09, 2018				
	6191148	Oct 09, 2018				
	6369085	May 25, 2018	DS DP U-729			
	6369085	May 25, 2018	DS DP U-770			
	6428810	Nov 03, 2019	DP U-469			
	6428810	Nov 03, 2019	DP U-729			
	6428810	Nov 03, 2019	DP U-770			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021153 001	7411070	May 25, 2018	DS			
	8466175	May 25, 2018		U-1417		
	8466175*PED	Nov 25, 2018				
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021153 002	6147103	Oct 09, 2018				
	6166213	Oct 09, 2018				
	6191148	Oct 09, 2018				
	6369085	May 25, 2018	DS DP	U-729		
	6369085	May 25, 2018	DS DP	U-770		
	6428810	Nov 03, 2019	DP	U-469		
	6428810	Nov 03, 2019	DP	U-729		
	6428810	Nov 03, 2019	DP	U-770		
	7411070	May 25, 2018	DS			
	8466175	May 25, 2018		U-1417		
	8466175*PED	Nov 25, 2018				
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021957 001	6369085	May 25, 2018	DS DP	U-1207		
	6369085	May 25, 2018	DS DP	U-729		
	6369085	May 25, 2018	DS DP	U-773		
	6428810	Nov 03, 2019	DP	U-1207		
	6428810	Nov 03, 2019	DP	U-729		
	6428810	Nov 03, 2019	DP	U-773		
	7411070	May 25, 2018	DS			
	8466175	May 25, 2018		U-1417		
	8466175*PED	Nov 25, 2018				
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021957 002	6369085	May 25, 2018	DS DP	U-1207		
	6369085	May 25, 2018	DS DP	U-729		
	6369085	May 25, 2018	DS DP	U-773		
	6428810	Nov 03, 2019	DP	U-1207		
	6428810	Nov 03, 2019	DP	U-729		
	6428810	Nov 03, 2019	DP	U-773		
	7411070	May 25, 2018	DS			
	8466175	May 25, 2018		U-1417		
	8466175*PED	Nov 25, 2018				
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021957 003	6369085	May 25, 2018	DS DP	U-1207		
	6428810	Nov 03, 2019	DP	U-1207		
	7411070	May 25, 2018	DS			
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 021957 004	6369085	May 25, 2018	DS DP	U-1207		
	6428810	Nov 03, 2019	DP	U-1207		
	7411070	May 25, 2018	DS			
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u></b>						
N 022101 001	6369085	May 25, 2018	DS DP	U-858		
	6428810	Nov 03, 2019	DP	U-858		
	7411070	May 25, 2018	DS			
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u></b>						
N 204655 001	6369085	May 25, 2018	DS DP	U-1509		
	6369085	May 25, 2018	DS DP	U-1875		
	6369085*PED	Nov 25, 2018				
	6428810	Nov 03, 2019	DP	U-1509		
	6428810	Nov 03, 2019	DP	U-1874		
	6428810*PED	May 03, 2020				
	7411070	May 25, 2018	DS			
	7411070*PED	Nov 25, 2018				
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u></b>						
N 207920 001	6369085	May 25, 2018	DS DP	U-1784		
	6369085*PED	Nov 25, 2018				
	6428810	Nov 03, 2019	DP	U-1785		
	6428810*PED	May 03, 2020				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ESOMEPRAZOLE MAGNESIUM - NEXIUM 24HR</u></b>						
N 207920 001	7411070	May 18, 2018	DS			
	7411070*PED	Nov 18, 2018				
<b><u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u></b>						
N 022511 001	6369085	May 25, 2018	DS DP U-1053		NPP	Jul 06, 2020
	6926907	Feb 28, 2023	DP U-1052			
	7411070	May 25, 2018	DS U-1053			
	7745466	Oct 13, 2018	DP U-1053			
	8557285	May 31, 2022	DP			
	8852636	May 31, 2022	DP U-1052			
	8858996	May 31, 2022	DP U-1052			
	8945621	Oct 17, 2031	U-1661			
	9161920	May 31, 2022	U-1760			
	9198888	May 31, 2022	U-1781			
	9220698	Mar 10, 2031	U-1781			
	9345695	May 31, 2022	DP			
	9393208	Sep 03, 2029	U-1781			
	9707181	May 31, 2022	DP			
<b><u>ESOMEPRAZOLE MAGNESIUM; NAPROXEN - VIMOVO</u></b>						
N 022511 002	6369085	May 25, 2018	DS DP U-1053		NPP	Jul 06, 2020
	6926907	Feb 28, 2023	DP U-1052			
	7411070	May 25, 2018	DS U-1053			
	7745466	Oct 13, 2018	DP U-1053			
	8557285	May 31, 2022	DP			
	8852636	May 31, 2022	DP U-1052			
	8858996	May 31, 2022	DP U-1052			
	8945621	Oct 17, 2031	U-1661			
	9161920	May 31, 2022	U-1760			
	9198888	May 31, 2022	U-1781			
	9345695	May 31, 2022	DP			
	9393208	Sep 03, 2029	U-1781			
	9707181	May 31, 2022	DP			
<b><u>ESTRADIOL - VAGIFEM</u></b>						
N 020908 002	5860946	Jul 01, 2017	DP			
	7018992	Sep 17, 2022	U-1023			
<b><u>ESTRADIOL - MENOSTAR</u></b>						
N 021674 001	5891868	Nov 21, 2017	DP U-594			
	6692763	Nov 21, 2017	DP U-594			
<b><u>ESTRADIOL - ELESTRIN</u></b>						
N 021813 001	7198801	Jun 25, 2022	DP			
	7470433	Aug 03, 2021	DP			
<b><u>ESTRADIOL - EVAMIST</u></b>						
N 022014 001	6978945	Jul 31, 2022	DP			
<b><u>ESTRADIOL - MINIVELLE</u></b>						
N 203752 001	6841716	Apr 27, 2020	DP			
	8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028		U-2086		
	9833419	Jul 10, 2028	DP			
<b><u>ESTRADIOL - MINIVELLE</u></b>						
N 203752 002	6841716	Apr 27, 2020	DP			
	8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028		U-2086		
	9833419	Jul 10, 2028	DP			
<b><u>ESTRADIOL - MINIVELLE</u></b>						
N 203752 003	6841716	Apr 27, 2020	DP			
	8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028		U-2086		
	9833419	Jul 10, 2028	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ESTRADIOL - MINIVELLE</u></b>						
N 203752 004	6841716	Apr 27, 2020	DP			
	8231906	Jul 04, 2030	DS DP			
	9730900	Jul 10, 2028		U-2086		
	9833419	Jul 10, 2028	DP			
<b><u>ESTRADIOL - MINIVELLE</u></b>						
N 203752 005	6841716	Apr 27, 2020	DP			
	8231906	Jul 04, 2030	DS DP			
	9724310	Jul 10, 2028	DS DP			
	9730900	Jul 10, 2028	DP	U-2086		
	9833419	Jul 10, 2028	DP			
<b><u>ESTRADIOL ACETATE - FEMTRACE</u></b>						
N 021633 001	6962908	Dec 21, 2021	DP			
	7572779	Oct 02, 2025		U-904		
	7799771	Dec 21, 2021	DP			
<b><u>ESTRADIOL ACETATE - FEMTRACE</u></b>						
N 021633 002	6962908	Dec 21, 2021	DP			
	7572779	Oct 02, 2025		U-904		
	7799771	Dec 21, 2021	DP			
<b><u>ESTRADIOL ACETATE - FEMTRACE</u></b>						
N 021633 003	6962908	Dec 21, 2021	DP			
	7572779	Oct 02, 2025		U-904		
	7799771	Dec 21, 2021	DP			
<b><u>ESTRADIOL; ESTRADIOL; NORGESTIMATE - PREFEST</u></b>						
N 021040 001	6747019	Mar 20, 2020		U-311		
	7320970	Mar 30, 2020	DP	U-844		
<b><u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u></b>						
N 021443 001	6660726	Mar 08, 2021	DS DP	U-904		
	6660726	Mar 08, 2021	DS DP	U-905		
	6855703	Feb 12, 2021	DS DP	U-904		
	6855703	Feb 12, 2021	DS DP	U-905		
<b><u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u></b>						
N 021443 002	6660726	Mar 08, 2021	DS DP	U-904		
	6660726	Mar 08, 2021	DS DP	U-905		
	6855703	Feb 12, 2021	DS DP	U-904		
	6855703	Feb 12, 2021	DS DP	U-905		
<b><u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u></b>						
N 021443 003	6660726	Mar 08, 2021	DS DP	U-904		
	6660726	Mar 08, 2021	DS DP	U-905		
	6855703	Feb 12, 2021	DS DP	U-904		
	6855703	Feb 12, 2021	DS DP	U-905		
<b><u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u></b>						
N 021443 004	6660726	Mar 08, 2021	DS DP	U-904		
	6660726	Mar 08, 2021	DS DP	U-905		
	6855703	Feb 12, 2021	DS DP	U-904		
	6855703	Feb 12, 2021	DS DP	U-905		
<b><u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVIA</u></b>						
N 021443 005	6660726	Mar 08, 2021	DS DP	U-904		
	6660726	Mar 08, 2021	DS DP	U-905		
	6855703	Feb 12, 2021	DS DP	U-904		
	6855703	Feb 12, 2021	DS DP	U-905		
<b><u>ETELCALCETIDE - PARSABIV</u></b>						
N 208325 001	8377880	Jul 29, 2030	DS DP		NCE	Feb 07, 2022
	8999932	Jul 29, 2030	DS DP	U-2014		
	9278995	Jul 29, 2030	DS			
	9701712	Jul 29, 2030	DS DP	U-2014		
	9820938	Jun 27, 2034	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ETELCALCETIDE - PARSABIV</u></b>						
N 208325 002	8377880	Jul 29, 2030	DS DP		NCE	Feb 07, 2022
	89999932	Jul 29, 2030	DS DP	U-2014		
	9278995	Jul 29, 2030	DS			
	9701712	Jul 29, 2030	DS DP	U-2014		
	9820938	Jun 27, 2034	DP			
<b><u>ETELCALCETIDE - PARSABIV</u></b>						
N 208325 003	8377880	Jul 29, 2030	DS DP		NCE	Feb 07, 2022
	89999932	Jul 29, 2030	DS DP	U-2014		
	9278995	Jul 29, 2030	DS			
	9701712	Jul 29, 2030	DS DP	U-2014		
	9820938	Jun 27, 2034	DP			
<b><u>ETEPLIRSEN - EXONDYS 51</u></b>						
N 206488 001	8486907	Jun 28, 2025		U-1904	Y	NCE
	9018368	Jun 28, 2025	DS DP		ODE-122	Sep 19, 2023
	9243245	Oct 27, 2028	DS	U-2097		
	9243245	Oct 27, 2028	DS	U-2098		
	9416361	May 04, 2021	DS			
	9506058	Mar 14, 2034		U-1918		
	9506058	Mar 14, 2034		U-1919		
<b><u>ETEPLIRSEN - EXONDYS 51</u></b>						
N 206488 002	8486907	Jun 28, 2025		U-1904	Y	NCE
	9018368	Jun 28, 2025	DS DP		ODE-122	Sep 19, 2023
	9243245	Oct 27, 2028	DS	U-2097		
	9243245	Oct 27, 2028	DS	U-2098		
	9416361	May 04, 2021	DS			
	9506058	Mar 14, 2034		U-1918		
	9506058	Mar 14, 2034		U-1919		
<b><u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; NORGESTIMATE; NORGESTIMATE; NORGESTIMATE - ORTHO TRI-CYCLEN LO</u></b>						
N 021241 001	6214815	Jun 09, 2019		U-112		
<b><u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u></b>						
N 021840 001	7320969	Jan 30, 2024		U-828		
	7615545	Jun 15, 2023		U-1		
	7855190	Dec 05, 2028		U-1		
	7858605	Jun 23, 2023	DP			
<b><u>ETHINYL ESTRADIOL; ETHINYL ESTRADIOL; LEVONORGESTREL - LOSEASONIQUE</u></b>						
N 022262 001	7615545	Jun 15, 2023		U-1		
	7855190	Dec 05, 2028		U-1		
	7858605	Jun 23, 2023	DP			
<b><u>ETHINYL ESTRADIOL; ETONOGESTREL - NUVARING</u></b>						
N 021187 001	5989581	Apr 08, 2018				
<b><u>ETHINYL ESTRADIOL; LEVONORGESTREL - FAYOSIM</u></b>						
A 205943 001					PC	Sep 30, 2017
<b><u>ETHINYL ESTRADIOL; LEVONORGESTREL - PREVEN EMERGENCY CONTRACEPTIVE KIT</u></b>						
N 020946 001	6156742	Dec 05, 2020		U-374		
<b><u>ETHINYL ESTRADIOL; LEVONORGESTREL - LYBREL</u></b>						
N 021864 001	6500814	Sep 03, 2018		U-1		
<b><u>ETHINYL ESTRADIOL; LEVONORGESTREL - QUARTETTE</u></b>						
N 204061 001	8415332	Mar 11, 2029	DP			
	8450299	Oct 07, 2025		U-1		
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE - FEMCON FE</u></b>						
N 021490 001	6667050	Apr 06, 2019	DP	U-1		
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE - NORETHINDRONE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u></b>						
N 022573 001	6667050	Apr 06, 2019	DP	U-828		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - MIBELAS 24 FE</u></b>						
A 206287 001					PC	Sep 11, 2017
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO LOESTRIN FE</u></b>						
N 022501 001	7704984	Feb 02, 2029	U-1090			
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - MINASTRIN 24 FE</u></b>						
N 203667 001	6667050	Apr 06, 2019	DP U-1			
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - TAYTULLA</u></b>						
N 204426 001	6652880	Mar 29, 2020	DP			
<b><u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LO MINASTRIN FE</u></b>						
N 204654 001	6667050	Apr 06, 2019	DP U-1			
	7704984	Feb 02, 2029	U-1			
<b><u>ETHIODIZED OIL - LIPIODOL</u></b>						
N 009190 001					ODE-64	Apr 04, 2021
<b><u>ETONOGESTREL - IMPLANON</u></b>						
N 021529 001	9757552	Jul 28, 2030	DP U-1			
<b><u>ETONOGESTREL - NEXPLANON</u></b>						
N 021529 002	8722037	Sep 28, 2027	DP			
	8888745	Aug 28, 2026	DP			
	9757552	Jul 28, 2030	DP U-1			
<b><u>ETRAVIRINE - INTELENCE</u></b>						
N 022187 001	6878717	Nov 05, 2019	U-1016			
	6878717	Nov 05, 2019	U-1237			
	6878717	Nov 05, 2019	U-256			
	7037917	Dec 13, 2020	DS DP U-1016			
	7037917	Dec 13, 2020	DS DP U-1237			
	7037917	Dec 13, 2020	DS DP U-256			
	7887845	Mar 25, 2019	DP			
	8003789	Nov 01, 2019	DS DP			
<b><u>ETRAVIRINE - INTELENCE</u></b>						
N 022187 002	6878717	Nov 05, 2019	U-1016			
	6878717	Nov 05, 2019	U-1237			
	6878717	Nov 05, 2019	U-256			
	7037917	Dec 13, 2020	DS DP U-1016			
	7037917	Dec 13, 2020	DS DP U-1237			
	7037917	Dec 13, 2020	DS DP U-256			
	7887845	Mar 25, 2019	DP			
	8003789	Nov 01, 2019	DS DP			
<b><u>EVEROLIMUS - ZORTRESS</u></b>						
N 021560 001	5665772	Sep 09, 2019	DS DP U-1049			
	5665772	Sep 09, 2019	DS DP U-1365			
	5665772*PED	Mar 09, 2020				
	6239124	Jul 29, 2017	U-1049			
	6239124*PED	Jan 29, 2018				
	6455518	Jul 29, 2017	U-1049			
	6455518	Jul 29, 2017	U-1365			
	6455518*PED	Jan 29, 2018				
<b><u>EVEROLIMUS - ZORTRESS</u></b>						
N 021560 002	5665772	Sep 09, 2019	DS DP U-1049			
	5665772	Sep 09, 2019	DS DP U-1365			
	5665772*PED	Mar 09, 2020				
	6239124	Jul 29, 2017	U-1049			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EVEROLIMUS - ZORTRESS</u></b>						
N 021560 002	6239124*PED	Jan 29, 2018				
	6455518	Jul 29, 2017		U-1049		
	6455518	Jul 29, 2017		U-1365		
	6455518*PED	Jan 29, 2018				
<b><u>EVEROLIMUS - ZORTRESS</u></b>						
N 021560 003	5665772	Sep 09, 2019	DS DP	U-1049		
	5665772	Sep 09, 2019	DS DP	U-1365		
	5665772*PED	Mar 09, 2020				
	6239124	Jul 29, 2017		U-1049		
	6239124*PED	Jan 29, 2018				
	6455518	Jul 29, 2017		U-1049		
	6455518	Jul 29, 2017		U-1365		
	6455518*PED	Jan 29, 2018				
<b><u>EVEROLIMUS - AFINITOR</u></b>						
N 022334 001	5665772	Sep 09, 2019	DS DP		I-724	Feb 26, 2019
	7297703	Dec 06, 2019	DP		ODE-108	Feb 26, 2023
	7741338	Dec 06, 2019	DP		ODE-11	May 05, 2018
	8410131	Nov 01, 2025		U-1368	ODE-24	Apr 26, 2019
	8410131*PED	May 01, 2026			ODE-4	Oct 29, 2017
	8436010	Feb 22, 2022		U-1396	PED	Apr 29, 2018
	8436010*PED	Aug 22, 2022			PED	Nov 05, 2018
	8778962	Feb 18, 2022		U-1541	PED	Oct 26, 2019
	8778962*PED	Aug 18, 2022			PED	
	9006224	Jul 01, 2028		U-1681		
<b><u>EVEROLIMUS - AFINITOR</u></b>						
N 022334 002	5665772	Sep 09, 2019	DS DP		I-724	Feb 26, 2019
	7297703	Dec 06, 2019	DP		ODE-108	Feb 26, 2023
	7741338	Dec 06, 2019	DP		ODE-11	May 05, 2018
	8410131	Nov 01, 2025		U-1368	ODE-24	Apr 26, 2019
	8410131*PED	May 01, 2026			ODE-4	Oct 29, 2017
	8436010	Feb 22, 2022		U-1396	PED	Apr 29, 2018
	8436010*PED	Aug 22, 2022			PED	Nov 05, 2018
	8778962	Feb 18, 2022		U-1541	PED	Oct 26, 2019
	8778962*PED	Aug 18, 2022			PED	
	9006224	Jul 01, 2028		U-1681		
<b><u>EVEROLIMUS - AFINITOR</u></b>						
N 022334 003	5665772	Sep 09, 2019	DS DP		I-724	Feb 26, 2019
	7297703	Dec 06, 2019	DP		ODE-108	Feb 26, 2023
	7741338	Dec 06, 2019	DP		ODE-11	May 05, 2018
	8410131	Nov 01, 2025		U-1368	ODE-24	Apr 26, 2019
	8410131*PED	May 01, 2026			ODE-4	Oct 29, 2017
	8436010	Feb 22, 2022		U-1396	PED	Apr 29, 2018
	8436010*PED	Aug 22, 2022			PED	Nov 05, 2018
	8778962	Feb 18, 2022		U-1541	PED	Oct 26, 2019
	8778962*PED	Aug 18, 2022			PED	
	9006224	Jul 01, 2028		U-1681		
<b><u>EVEROLIMUS - AFINITOR</u></b>						
N 022334 004	5665772	Sep 09, 2019	DS DP		I-724	Feb 26, 2019
	7297703	Dec 06, 2019	DP		ODE-108	Feb 26, 2023
	7741338	Dec 06, 2019	DP		ODE-11	May 05, 2018
	8410131	Nov 01, 2025		U-1368	ODE-24	Apr 26, 2019
	8410131*PED	May 01, 2026			ODE-4	Oct 29, 2017
	8436010	Feb 22, 2022		U-1396	PED	Apr 29, 2018
	8436010*PED	Aug 22, 2022			PED	Nov 05, 2018
	8778962	Feb 18, 2022		U-1541	PED	Oct 26, 2019
	8778962*PED	Aug 18, 2022			PED	
	9006224	Jul 01, 2028		U-1681		
<b><u>EVEROLIMUS - AFINITOR DISPERZ</u></b>						
N 203985 001	5665772	Sep 09, 2019	DS DP		ODE-4	Oct 29, 2017
	7297703	Dec 06, 2019	DP		PED	Apr 29, 2018
	8617598	Sep 27, 2022	DP			
	8617598*PED	Mar 27, 2023				
	8778962	Feb 18, 2022		U-1541		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EVEROLIMUS - AFINITOR DISPERZ</u></b>						
N 203985 001	8778962*PED	Aug 18, 2022				
<b><u>EVEROLIMUS - AFINITOR DISPERZ</u></b>						
N 203985 002	5665772 7297703 8617598 8617598*PED 8778962 8778962*PED	Sep 09, 2019 Dec 06, 2019 Sep 27, 2022 Mar 27, 2023 Feb 18, 2022 Aug 18, 2022	DS DP DP DP U-1541		ODE-4 PED	Oct 29, 2017 Apr 29, 2018
<b><u>EVEROLIMUS - AFINITOR DISPERZ</u></b>						
N 203985 003	5665772 7297703 8617598 8617598*PED 8778962 8778962*PED	Sep 09, 2019 Dec 06, 2019 Sep 27, 2022 Mar 27, 2023 Feb 18, 2022 Aug 18, 2022	DS DP DP DP U-1541		ODE-4 PED	Oct 29, 2017 Apr 29, 2018
<b><u>EXENATIDE - BYDUREON BCISE</u></b>						
N 209210 001	6479065 6667061 6824822 6872700 6956026 7223440 7456254 7563871 7612176 7741269 8329648 8329648 8329648 8329648 8431685 8461105 8895033 8895033 8895033 8906851 9238076	Aug 10, 2020 May 25, 2020 Oct 09, 2022 Jan 14, 2020 Jan 07, 2018 Aug 31, 2021 Jun 30, 2025 Apr 15, 2024 Apr 13, 2025 Jan 07, 2018 Aug 18, 2026 Aug 18, 2026 Aug 18, 2026 Aug 18, 2026 Apr 13, 2025 Apr 13, 2025 Oct 04, 2030 Oct 04, 2030 Oct 04, 2030 Aug 18, 2026 Apr 15, 2024	DP DP DP U-654 U-687 DP DP U-1223 DP DP U-1223 U-1224 U-1313 U-2154 U-2155 U-2156 DP U-412 DP U-412 DP U-1313 DP U-2157 DP U-2158 U-1313 DP U-412		NP	Oct 20, 2020
<b><u>EXENATIDE SYNTHETIC - BYETTA</u></b>						
N 021773 001	6872700 6902744 6956026 6956026 6956026 7297761 7521423 7741269 7741269 7741269	Jan 14, 2020 Jan 14, 2020 Jan 07, 2018 Jan 07, 2018 Jan 07, 2018 Oct 15, 2017 Oct 15, 2017 Jan 07, 2018 Jan 07, 2018 Jan 07, 2018	U-654 DP U-1074 U-1623 U-687 DP DP U-1074 U-1108 U-653		M-148	Nov 24, 2017
<b><u>EXENATIDE SYNTHETIC - BYETTA</u></b>						
N 021773 002	6872700 6902744 6956026 6956026 6956026 7297761 7521423 7741269 7741269 7741269	Jan 14, 2020 Jan 14, 2020 Jan 07, 2018 Jan 07, 2018 Jan 07, 2018 Oct 15, 2017 Oct 15, 2017 Jan 07, 2018 Jan 07, 2018 Jan 07, 2018	U-654 DP U-1074 U-1623 U-687 DP DP U-1074 U-1108 U-653		M-148	Nov 24, 2017
<b><u>EXENATIDE SYNTHETIC - BYDUREON</u></b>						
N 022200 001	6414126 6414126 6479065	Oct 04, 2020 Oct 04, 2020 Aug 10, 2020	DS DP U-2139 DS DP U-493 DP		M-162 M-212	Sep 24, 2018 Oct 20, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EXENATIDE SYNTHETIC - BYDUREON</u></b>						
N 022200 001	6495164	May 25, 2020	DP			
	6515117	Oct 04, 2020	DS DP U-2139			
	6515117	Oct 04, 2020	DS DP U-493			
	6667061	May 25, 2020	DP			
	6824822	Oct 09, 2022	DP			
	6872700	Jan 14, 2020	U-654			
	6936590	Oct 04, 2020	U-493			
	6956026	Jan 07, 2018	U-687			
	7223440	Aug 31, 2021	DP			
	7456254	Jun 30, 2025	DP U-1223			
	7563871	Apr 15, 2024	DP			
	7612176	Apr 13, 2025	DP U-1223			
	7741269	Jan 07, 2018	U-1224			
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-1313			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-412			
	8461105	Apr 13, 2025	DP U-412			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8716251	Mar 21, 2028	DP			
	8906851	Aug 18, 2026	U-1313			
	9198925	Oct 04, 2020	U-2139			
	9198925	Oct 04, 2020	U-493			
	9238076	Apr 15, 2024	DP U-412			
<b><u>EXENATIDE SYNTHETIC - BYDUREON PEN</u></b>						
N 022200 002	6414126	Oct 04, 2020	DS DP U-2139			
	6414126	Oct 04, 2020	DS DP U-493			
	6479065	Aug 10, 2020	DP			
	6495164	May 25, 2020	DP			
	6515117	Oct 04, 2020	DS DP U-2139			
	6515117	Oct 04, 2020	DS DP U-493			
	6667061	May 25, 2020	DP			
	6824822	Oct 09, 2022	DP			
	6872700	Jan 14, 2020	U-654			
	6936590	Oct 04, 2020	U-493			
	6956026	Jan 07, 2018	U-687			
	7223440	Aug 31, 2021	DP			
	7456254	Jun 30, 2025	DP U-1223			
	7563871	Apr 15, 2024	DP			
	7612176	Apr 13, 2025	DP U-1223			
	7741269	Jan 07, 2018	U-1224			
	7851502	Aug 19, 2028	DP			
	7919598	Dec 16, 2029	DS			
	8216180	Jan 12, 2028	DP			
	8221786	Mar 21, 2028	DP			
	8329648	Aug 18, 2026	U-1313			
	8361972	Mar 21, 2028	U-2139			
	8361972	Mar 21, 2028	U-493			
	8431685	Apr 13, 2025	DP U-412			
	8439864	Mar 25, 2028	DP			
	8461105	Apr 13, 2025	DP U-412			
	8501698	Jun 20, 2027	DP U-493			
	8685934	May 26, 2030	U-1522			
	8690837	May 19, 2029	DP			
	8716251	Mar 21, 2028	DP			
	8721615	Jan 18, 2030	DP			
	8758292	Nov 12, 2027	DP			
	8827963	Feb 04, 2029	DP			
	8906851	Aug 18, 2026	U-1313			
	8998876	Jan 07, 2030	DP			
	9198925	Oct 04, 2020	U-2139			
	9198925	Oct 04, 2020	U-493			
	9238076	Apr 15, 2024	DP U-412			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>EXENATIDE SYNTHETIC - BYDUREON PEN</u></b>						
N 022200 002	9320853	Mar 25, 2028	DP			
<b><u>EZETIMIBE - ZETIA</u></b>						
N 021445 001	7030106	Jan 25, 2022	DP			
	7612058	Oct 30, 2025		U-1027		
	7612058	Oct 30, 2025		U-1173		
	7612058*PED	Apr 30, 2026				
<b><u>FAMOTIDINE - PEPCID AC</u></b>						
N 020801 002	6814978	Aug 26, 2021	DP			
<b><u>FAMOTIDINE - FLUXID</u></b>						
N 021712 001	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>FAMOTIDINE - FLUXID</u></b>						
N 021712 002	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>FAMOTIDINE; IBUPROFEN - DUEXIS</u></b>						
N 022519 001	8067033	Jul 18, 2026	DP			
	8067451	Jul 18, 2026		DP U-1196		
	8309127	Jul 18, 2026		DP		
	8318202	Jul 18, 2026		DP		
	8449910	Jul 18, 2026		DP		
	8501228	Jul 18, 2026		U-1196		
<b><u>FEBUXOSTAT - ULORIC</u></b>						
N 021856 001	5614520	Mar 25, 2019	DS DP	U-954	M-205	Aug 15, 2020
	6225474	Jun 18, 2019	DS			
	7361676	Mar 08, 2024		DP		
	8372872	Sep 08, 2031		U-1346		
	9107912	Sep 08, 2031		U-1346		
<b><u>FEBUXOSTAT - ULORIC</u></b>						
N 021856 002	5614520	Mar 25, 2019	DS DP	U-954	M-205	Aug 15, 2020
	6225474	Jun 18, 2019	DS			
	7361676	Mar 08, 2024		DP		
	8372872	Sep 08, 2031		U-1346		
	9107912	Sep 08, 2031		U-1346		
<b><u>FENOFIBRATE - TRICOR</u></b>						
N 021203 001	6074670	Jan 09, 2018				
	6277405	Jan 09, 2018				
	6589552	Jan 09, 2018				
	6652881	Jan 09, 2018		DP		
	7037529	Jan 09, 2018		DP		
	7041319	Jan 09, 2018		DP		
<b><u>FENOFIBRATE - TRICOR</u></b>						
N 021203 003	6074670	Jan 09, 2018				
	6277405	Jan 09, 2018				
	6589552	Jan 09, 2018				
	6652881	Jan 09, 2018		DP		
	7037529	Jan 09, 2018		DP		
	7041319	Jan 09, 2018		DP		
<b><u>FENOFIBRATE - TRIGLIDE</u></b>						
N 021350 001	6696084	Sep 11, 2021	DS DP	U-680		
<b><u>FENOFIBRATE - TRIGLIDE</u></b>						
N 021350 002	6696084	Sep 11, 2021	DS DP	U-680		
<b><u>FENOFIBRATE - TRICOR</u></b>						
N 021656 001	6277405	Jan 09, 2018	DS			
	6375986	Sep 21, 2020		DP U-615		
	6652881	Jan 09, 2018	DS			
	7037529	Jan 09, 2018		DP		
	7041319	Jan 09, 2018		DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>FENOFIBRATE - TRICOR</b>						
N 021656 001	7276249	Feb 21, 2023	DP			
	7320802	Feb 21, 2023	U-847			
<b>FENOFIBRATE - TRICOR</b>						
N 021656 002	6277405	Jan 09, 2018	DS			
	6375986	Sep 21, 2020	DP U-615			
	6652881	Jan 09, 2018	DS			
	7037529	Jan 09, 2018	DP			
	7041319	Jan 09, 2018	DP			
	7276249	Feb 21, 2023	DP			
	7320802	Feb 21, 2023	U-847			
<b>FENOFIBRATE - ANTARA (MICRONIZED)</b>						
N 021695 001	7101574	Aug 20, 2020	DS DP			
	7863331	Aug 08, 2020	U-1106			
	7863331	Aug 08, 2020	U-1107			
<b>FENOFIBRATE - ANTARA (MICRONIZED)</b>						
N 021695 003	7101574	Aug 20, 2020	DS DP			
	7863331	Aug 08, 2020	U-1106			
	7863331	Aug 08, 2020	U-1107			
<b>FENOFIBRATE - ANTARA (MICRONIZED)</b>						
N 021695 004	8026281	Apr 22, 2025	U-1447			
	8026281	Apr 22, 2025	U-1448			
<b>FENOFIBRATE - ANTARA (MICRONIZED)</b>						
N 021695 005	8026281	Apr 22, 2025	U-1447			
	8026281	Apr 22, 2025	U-1448			
	9314447	May 31, 2033	DP U-1447			
	9314447	May 31, 2033	DP U-1448			
<b>FENOFIBRATE - FENOGLIDE</b>						
N 022118 001	7658944	Dec 09, 2024	DP			
	8124125	Oct 01, 2024	DP U-1234			
	8481078	Oct 01, 2024	DP U-1416			
	9173847	Oct 01, 2024	DP			
<b>FENOFIBRATE - FENOGLIDE</b>						
N 022118 002	7658944	Dec 09, 2024	DP			
	8124125	Oct 01, 2024	DP U-1234			
	8481078	Oct 01, 2024	DP U-1416			
	9173847	Oct 01, 2024	DP			
<b>FENOFIBRIC ACID - FIBRICOR</b>						
N 022418 001	7569612	Aug 20, 2027	U-1000			
	7741373	Aug 20, 2027	U-1059			
	7741374	Aug 20, 2027	U-1060			
	7741374	Aug 20, 2027	U-1061			
	7915247	Aug 20, 2027	U-1000			
	7915247	Aug 20, 2027	U-1059			
	7915247	Aug 20, 2027	U-1061			
<b>FENOFIBRIC ACID - FIBRICOR</b>						
N 022418 002	7569612	Aug 20, 2027	U-1000			
	7741373	Aug 20, 2027	U-1059			
	7741374	Aug 20, 2027	U-1060			
	7741374	Aug 20, 2027	U-1061			
	7915247	Aug 20, 2027	U-1000			
	7915247	Aug 20, 2027	U-1059			
	7915247	Aug 20, 2027	U-1061			
<b>FENTANYL - SUBSYS</b>						
N 202788 001	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030	U-55			
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP U-55			
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP U-55			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 001	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 002	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030		U-55		
	8835460	Jan 25, 2027	DP	U-55		
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP	U-55		
	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 003	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030		U-55		
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP	U-55		
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP	U-55		
	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 004	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030		U-55		
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP	U-55		
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP	U-55		
	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 005	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030		U-55		
	8835460	Jan 25, 2027	DP	U-55		
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP	U-55		
	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 006	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030		U-55		
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP	U-55		
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP	U-55		
	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL - SUBSYS</u></b>						
N 202788 007	8486972	Apr 27, 2030	DP			
	8486973	Apr 27, 2030		U-55		
	8835459	Jan 25, 2027	DP			
	8835460	Jan 25, 2027	DP	U-55		
	9241935	Jan 25, 2027	DP			
	9289387	Jan 25, 2027	DP	U-55		
	9642797	Jan 25, 2027	DP	U-55		
	9642844	Jan 25, 2027	DP			
<b><u>FENTANYL CITRATE - FENTORA</u></b>						
N 021947 001	6200604	Mar 26, 2019	U-767			
	6974590	Mar 26, 2019		U-767		
	7862832	Jun 15, 2028	DP			
	7862833	Jun 15, 2028	DP			
	8092832	Dec 30, 2024	DP			
	8728441	Mar 26, 2019		U-1514		
	8753611	Mar 26, 2019	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>FENTANYL CITRATE - FENTORA</b>						
N 021947 001	8765100	Mar 26, 2019	DP			
<b>FENTANYL CITRATE - FENTORA</b>						
N 021947 002	6200604 6974590 7862832 7862833 8092832 8119158 8728441 8753611 8765100	Mar 26, 2019 Mar 26, 2019 Jun 15, 2028 Jun 15, 2028 Dec 30, 2024 Dec 30, 2024 Mar 26, 2019 Mar 26, 2019 Mar 26, 2019	U-767 U-767 DP DP DP DP U-1514 U-1514 DP			
<b>FENTANYL CITRATE - FENTORA</b>						
N 021947 003	6200604 6974590 7862832 7862833 8092832 8119158 8728441 8753611 8765100	Mar 26, 2019 Mar 26, 2019 Jun 15, 2028 Jun 15, 2028 Dec 30, 2024 Dec 30, 2024 Mar 26, 2019 Mar 26, 2019 Mar 26, 2019	U-767 U-767 DP DP DP DP U-1514 U-1514 DP			
<b>FENTANYL CITRATE - FENTORA</b>						
N 021947 004	6200604 6974590 7862832 7862833 8092832 8119158 8728441 8753611 8765100	Mar 26, 2019 Mar 26, 2019 Jun 15, 2028 Jun 15, 2028 Dec 30, 2024 Dec 30, 2024 Mar 26, 2019 Mar 26, 2019 Mar 26, 2019	U-767 U-767 DP DP DP DP U-1514 U-1514 DP			
<b>FENTANYL CITRATE - FENTORA</b>						
N 021947 005	6200604 6974590 7862832 7862833 8092832 8119158 8728441 8753611 8765100	Mar 26, 2019 Mar 26, 2019 Jun 15, 2028 Jun 15, 2028 Dec 30, 2024 Dec 30, 2024 Mar 26, 2019 Mar 26, 2019 Mar 26, 2019	U-767 U-767 DP DP DP DP U-1514 U-1514 DP			
<b>FENTANYL CITRATE - FENTORA</b>						
N 021947 006	6200604 6974590	Mar 26, 2019 Mar 26, 2019	U-767 U-767			
<b>FENTANYL CITRATE - ONSOLIS</b>						
N 022266 001	7579019 9597288	Jan 22, 2020 Jul 23, 2027	U-767 DP U-767			
<b>FENTANYL CITRATE - ONSOLIS</b>						
N 022266 002	7579019 9597288	Jan 22, 2020 Jul 23, 2027	U-767 DP U-767			
<b>FENTANYL CITRATE - ONSOLIS</b>						
N 022266 003	7579019 9597288	Jan 22, 2020 Jul 23, 2027	U-767 DP U-767			
<b>FENTANYL CITRATE - ONSOLIS</b>						
N 022266 004	7579019 9597288	Jan 22, 2020 Jul 23, 2027	U-767 DP U-767			
<b>FENTANYL CITRATE - ONSOLIS</b>						
N 022266 005	7579019 9597288	Jan 22, 2020 Jul 23, 2027	U-767 DP U-767			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>FENTANYL CITRATE - ONSOLIS</b>						
N 022266 005	7579019	Jan 22, 2020	U-767			
	9597288	Jul 23, 2027	DP U-767			
<b>FENTANYL CITRATE - ABSTRAL</b>						
N 022510 001	6759059	Sep 24, 2019	DP U-767			
	6761910	Sep 24, 2019	DP U-767			
	7910132	Sep 24, 2019	DP U-767			
<b>FENTANYL CITRATE - ABSTRAL</b>						
N 022510 002	6759059	Sep 24, 2019	DP U-767			
	6761910	Sep 24, 2019	DP U-767			
	7910132	Sep 24, 2019	DP U-767			
<b>FENTANYL CITRATE - ABSTRAL</b>						
N 022510 003	6759059	Sep 24, 2019	DP U-767			
	6761910	Sep 24, 2019	DP U-767			
	7910132	Sep 24, 2019	DP U-767			
<b>FENTANYL CITRATE - ABSTRAL</b>						
N 022510 004	6759059	Sep 24, 2019	DP U-767			
	6761910	Sep 24, 2019	DP U-767			
	7910132	Sep 24, 2019	DP U-767			
<b>FENTANYL CITRATE - ABSTRAL</b>						
N 022510 005	6759059	Sep 24, 2019	DP U-767			
	6761910	Sep 24, 2019	DP U-767			
	7910132	Sep 24, 2019	DP U-767			
<b>FENTANYL CITRATE - ABSTRAL</b>						
N 022510 006	6759059	Sep 24, 2019	DP U-767			
	6761910	Sep 24, 2019	DP U-767			
	7910132	Sep 24, 2019	DP U-767			
<b>FENTANYL CITRATE - LAZANDA</b>						
N 022569 001	6432440	Apr 20, 2018	DP U-1169			
	8216604	Oct 03, 2024	U-767			
	8889176	Jan 16, 2024	U-767			
	9078814	Jan 08, 2024	DP			
	9731869	Jan 26, 2032	DP			
<b>FENTANYL CITRATE - LAZANDA</b>						
N 022569 002	6432440	Apr 20, 2018	DP U-1169			
	8216604	Oct 03, 2024	U-767			
	8889176	Jan 16, 2024	U-767			
	9078814	Jan 08, 2024	DP			
	9731869	Jan 26, 2032	DP			
<b>FENTANYL CITRATE - LAZANDA</b>						
N 022569 003	9731869	Jan 26, 2032	DP			
<b>FENTANYL HYDROCHLORIDE - IONSYS</b>						
N 021338 001	6169920	Jan 02, 2018	DP U-736			
	6181963	Nov 02, 2019	DP			
	6195582	Jan 28, 2019	DP U-736			
	6881208	Apr 19, 2022	U-736			
	6975902	Apr 01, 2024	DP			
	8301238	Sep 30, 2031	DP			
	8428708	May 21, 2032	U-736			
	8428709	Jun 11, 2032	DP U-736			
	8781571	Mar 31, 2032	DP U-736			
	9095706	Feb 03, 2033	DP			
	9364656	Sep 30, 2031	U-736			
	9731121	Oct 17, 2031	DP			
<b>FERRIC CARBOXYMALTOSE - INJECTAFER</b>						
N 203565 001	7612109	Feb 05, 2024	DS DP			
	7754702	Feb 13, 2027	DP U-1432			
	8895612	Jan 08, 2027	DP U-1620			
	9376505	Oct 20, 2023	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FERRIC CITRATE - AURYXTA</u></b>						
N 205874 001	7767851	Feb 18, 2024	DS DP			
	8093423	Apr 21, 2026		U-1577		
	8299298	Feb 18, 2024		DP		
	8338642	Feb 18, 2024	DS DP	U-1577		
	8609896	Feb 18, 2024		DP		
	8754257	Feb 18, 2024		DP		
	8754258	Feb 18, 2024		DP		
	8846976	Feb 18, 2024		U-1577		
	8901349	Feb 18, 2024		U-1577		
	9050316	Feb 18, 2024		U-1577		
	9328133	Feb 18, 2024	DS DP	U-1577		
	9387191	Jul 21, 2030		DP		
	9757416	Feb 18, 2024	DS DP	U-1577		
<b><u>FERRIC PYROPHOSPHATE CITRATE - TRIFERIC</u></b>						
N 206317 001	7816404	Apr 17, 2029		DP U-1656		
<b><u>FERUMOXYTOL - FERAHEM</u></b>						
N 022180 001	6599498	Jun 30, 2023	DS DP			
	7553479	Mar 08, 2020	DS DP			
	7871597	Mar 08, 2020	DS DP			
	8501158	Mar 08, 2020		U-1422		
	8591864	Mar 08, 2020		DP		
	8926947	Mar 08, 2020	DS DP			
<b><u>FESOTERODINE FUMARATE - TOVIAZ</u></b>						
N 022030 001	6858650	Jul 03, 2022	DS	U-913		
	7384980	May 11, 2019	DS DP	U-913		
	7807715	Jun 07, 2027		DP U-913		
	7855230	May 11, 2019		U-913		
	7985772	May 11, 2019	DS DP	U-913		
	8088398	Jun 07, 2027		DP U-913		
	8338478	May 11, 2019	DS DP	U-913		
	8501723	Jun 07, 2027		DP		
<b><u>FESOTERODINE FUMARATE - TOVIAZ</u></b>						
N 022030 002	6858650	Jul 03, 2022	DS	U-913		
	7384980	May 11, 2019	DS DP	U-913		
	7807715	Jun 07, 2027		DP U-913		
	7855230	May 11, 2019		U-913		
	7985772	May 11, 2019	DS DP	U-913		
	8088398	Jun 07, 2027		DP U-913		
	8338478	May 11, 2019	DS DP	U-913		
	8501723	Jun 07, 2027		DP		
<b><u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u></b>						
N 021909 002	6723348	Nov 26, 2021		DP U-1466		
<b><u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u></b>						
N 021909 003	6723348	Nov 26, 2021		DP		
<b><u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA ALLERGY</u></b>						
N 201373 001	8933097	Aug 16, 2032		DP		
<b><u>FEXOFENADINE HYDROCHLORIDE - CHILDREN'S ALLEGRA HIVES</u></b>						
N 201373 002	8933097	Aug 16, 2032		DP		
<b><u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 12 HOUR ALLERGY AND CONGESTION</u></b>						
N 020786 002	6039974	Jul 31, 2018		DP		
<b><u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 24 HOUR ALLERGY AND CONGESTION</u></b>						
N 021704 002	6613357	Dec 25, 2020		DP U-1159		
	RE39069	May 29, 2018		DP		
<b><u>FIDAXOMICIN - DIFICID</u></b>						
N 201699 001	7378508	Jul 31, 2027	DS DP			
	7863249	Jul 31, 2027	DS DP			
	7906489	Mar 04, 2027		U-319		
	8586551	Jul 15, 2023	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FIDAXOMICIN - DIFCICID</u></b>						
N 201699 001	8859510	Jul 31, 2027		U-319		
<b><u>FINAFLOXACIN - XTORO</u></b>						
N 206307 001	8536167	Aug 08, 2031	U-1679		NCE	Dec 17, 2019
	9119859	Jul 02, 2030	U-1679		PED	Jun 17, 2020
	9504691	Nov 21, 2033	DP U-1679			
<b><u>FINASTERIDE - PROSCAR</u></b>						
N 020180 001	5942519	Oct 23, 2018		U-280		
<b><u>FINGOLIMOD - GILENYA</u></b>						
N 022527 001	5604229	Feb 18, 2019	DS	U-1086		
	6004565	Sep 23, 2017		U-1086		
	8324283	Mar 29, 2026				
	9187405	Jun 25, 2027		U-1086		
<b><u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u></b>						
N 207648 001					NCE	Jul 13, 2021
<b><u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u></b>						
N 207648 002					NCE	Jul 13, 2021
<b><u>FISH OIL; MEDIUM CHAIN TRIGLYCERIDES; OLIVE OIL; SOYBEAN OIL - SMOFLIPID 20%</u></b>						
N 207648 003					NCE	Jul 13, 2021
<b><u>FLIBANSERIN - ADDYI</u></b>						
N 022526 001	7151103	May 09, 2023		U-1734		
	7420057	Aug 01, 2022	DS DP		NCE	Aug 18, 2020
	8227471	May 09, 2023		U-1734		
	9468639	Oct 16, 2022		U-1734		
<b><u>FLORBETABEN F-18 - NEURACEQ</u></b>						
N 204677 001	7807135	Mar 18, 2029	DS DP	U-1497	NCE	Mar 21, 2019
<b><u>FLORBETAPIR F-18 - AMYVID</u></b>						
N 202008 001	7687052	Apr 30, 2027	DS DP			
	8506929	Apr 30, 2027	DS DP	U-1423		
<b><u>FLORBETAPIR F-18 - AMYVID</u></b>						
N 202008 002	7687052	Apr 30, 2027	DS DP			
	8506929	Apr 30, 2027	DS DP	U-1423		
<b><u>FLORBETAPIR F-18 - AMYVID</u></b>						
N 202008 003	7687052	Apr 30, 2027	DS DP			
	8506929	Apr 30, 2027	DS DP	U-1423		
<b><u>FLUCICLOVINE F-18 - AXUMIN</u></b>						
N 208054 001	5808146	Nov 09, 2018	DS			
	9387266	Nov 28, 2026		U-1879	NCE	May 27, 2021
<b><u>FLUDARABINE PHOSPHATE - OFORTA</u></b>						
N 022273 001	7148207	Dec 20, 2022		DP U-944		
	7547776	Dec 10, 2018	DS			
<b><u>FLUOCINOLONE ACETONIDE - RETISERT</u></b>						
N 021737 001	6217895	Mar 22, 2019		DP U-708		
	6548078	Mar 22, 2019		DP U-708		
<b><u>FLUOCINOLONE ACETONIDE - ILUVIEN</u></b>						
N 201923 001	6217895	Mar 22, 2019		DP U-1597		
	6375972	Apr 26, 2020		DP U-1597	NP	Sep 26, 2017
	6548078	Mar 22, 2019		DP U-1597		
	8252307	Jun 27, 2019		DP		
	8871241	Aug 12, 2027		DP		
<b><u>FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN - TRI-LUMA</u></b>						
N 021112 001	7915243	Mar 22, 2026	DP			
	7939516	May 04, 2025	DP			
	8247395	Oct 22, 2022	DP			
	8653053	Oct 25, 2022	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN - TRI-LUMA</u></b>						
N 021112 001	7915243	Mar 22, 2026	DP			
	7939516	May 04, 2025	DP			
	8247395	Oct 22, 2022	DP			
	8653053	Oct 25, 2022	DP			
<b><u>FLUOCINONIDE - VANOS</u></b>						
N 021758 001	6765001	Dec 21, 2021	DP			
	7220424	Jan 07, 2023	U-861			
	7794738	Sep 11, 2022	U-1084			
	8232264	Mar 09, 2023	DP			
<b><u>FLUOROURACIL - CARAC</u></b>						
N 020985 001	6670335	Jun 02, 2021	DP U-68			
<b><u>FLUOROURACIL - TOLAK</u></b>						
N 022259 001	7169401	Jul 18, 2023	DP		NP	Sep 18, 2018
<b><u>FLUOXETINE HYDROCHLORIDE - PROZAC</u></b>						
N 018936 001	6960577	Nov 01, 2017	U-963			
<b><u>FLUOXETINE HYDROCHLORIDE - PROZAC</u></b>						
N 018936 003	6960577	Nov 01, 2017	U-963			
<b><u>FLUOXETINE HYDROCHLORIDE - PROZAC</u></b>						
N 018936 006	6960577	Nov 01, 2017	U-963			
<b><u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u></b>						
N 021520 001	6960577	Nov 01, 2017	U-962		M-142	Oct 10, 2017
<b><u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u></b>						
N 021520 002	6960577	Nov 01, 2017	U-962		M-142	Oct 10, 2017
<b><u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u></b>						
N 021520 003	6960577	Nov 01, 2017	U-962		M-142	Oct 10, 2017
<b><u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u></b>						
N 021520 004	6960577	Nov 01, 2017	U-962		M-142	Oct 10, 2017
<b><u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u></b>						
N 021520 005	6960577	Nov 01, 2017	U-962		M-142	Oct 10, 2017
<b><u>FLUTEMETAMOL F-18 - VIZAMYL</u></b>						
N 203137 001	7270800	Sep 03, 2025	DS DP U-336		NCE	Oct 25, 2018
	7351401	Jan 24, 2023	DS DP U-336			
	8236282	May 21, 2024	DS DP			
	8691185	Jan 24, 2023	U-336			
	8916131	Sep 16, 2028	DP			
<b><u>FLUTEMETAMOL F-18 - VIZAMYL</u></b>						
N 203137 002	7270800	Sep 03, 2025	DS DP U-336		NCE	Oct 25, 2018
	7351401	Jan 24, 2023	DS DP U-336			
	8236282	May 21, 2024	DS DP			
	8691185	Jan 24, 2023	U-336			
	8916131	Sep 16, 2028	DP			
<b><u>FLUTICASONE FUROATE - FLONASE SENSI-MIST ALLERGY RELIEF</u></b>						
N 022051 002	6858596	Aug 03, 2021	DP U-1890			
	7101866	Aug 03, 2021	DS DP U-1890			
	7541350	Aug 03, 2021	DP U-1890			
	8062264	Apr 05, 2026	DP			
	8147461	Oct 15, 2028	DP			
	8347879	Jul 15, 2028	DP			
	8752543	Apr 05, 2026	DP			
	9320862	Nov 06, 2024	DP			
<b><u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u></b>						
N 205625 001	7101866	Aug 03, 2021	DS DP U-1559		NP	Aug 20, 2017
	7629335	Aug 03, 2021	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u></b>						
N 205625 001	8201556	Feb 05, 2029	DP			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
<b><u>FLUTICASONE FUROATE - ARNUITY ELLIPTA</u></b>						
N 205625 002	7101866	Aug 03, 2021	DS DP U-1559		NP	Aug 20, 2017
	7629335	Aug 03, 2021	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8201556	Feb 05, 2029	DP			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
<b><u>FLUTICASONE FUROATE; UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - TRELEGY ELLIPTA</u></b>						
N 209482 001	6537983	Aug 03, 2021	DP U-2125		NCE	May 10, 2018
	6759398	Aug 03, 2021	DP U-2125		NCE	Dec 18, 2018
	6878698	Aug 03, 2021	U-2134			
	7101866	Aug 03, 2021	DS DP U-2126			
	7439393	May 21, 2025	DS DP U-2127			
	7488827	Dec 18, 2027	DS DP			
	7498440	Apr 27, 2025	DS DP			
	7629335	Aug 03, 2021	DP			
	7776895	Sep 11, 2022	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8183257	Jul 27, 2025	U-2128			
	8309572	Apr 27, 2025	U-2129			
	8511304	Jun 14, 2027	DP			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	9750762	Nov 29, 2030	DP			
	RE44874	Mar 23, 2023	DS DP U-2127			
<b><u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u></b>						
N 204275 001	6537983	Aug 03, 2021	DP U-1401		I-708	Apr 30, 2018
	6537983	Aug 03, 2021	DP U-1691		M-202	May 15, 2020
	6759398	Aug 03, 2021	DP U-1401		NCE	May 10, 2018
	6759398	Aug 03, 2021	DP U-1691			
	6878698	Aug 03, 2021	U-1401			
	7101866	Aug 03, 2021	DS DP U-1401			
	7101866	Aug 03, 2021	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-1401			
	7439393	May 21, 2025	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-2099			
	7439393	May 21, 2025	DS DP U-2100			
	7629335	Aug 03, 2021	DP			
	7776895	Sep 11, 2022	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8511304	Jun 14, 2027	DP U-1424			
	8511304	Jun 14, 2027	DP U-1691			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	RE44874	Mar 23, 2023	DS DP U-1548			
	RE44874	Mar 23, 2023	DS DP U-1691			
<b><u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u></b>						
N 204275 002	6537983	Aug 03, 2021	DP U-1691		M-202	May 15, 2020
	6759398	Aug 03, 2021	DP U-1691		NCE	May 10, 2018
	7101866	Aug 03, 2021	DS DP U-1691		NP	Apr 30, 2018
	7439393	May 21, 2025	DS DP U-1691			
	7439393	May 21, 2025	DS DP U-2099			
	7439393	May 21, 2025	DS DP U-2100			
	7629335	Aug 03, 2021	DP			
	7776895	Sep 11, 2022	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FLUTICASONE FUROATE; VILANTEROL TRIFENATATE - BREO ELLIPTA</u></b>						
N 204275 002	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8511304	Jun 14, 2027	DP U-1691			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	RE44874	Mar 23, 2023	DS DP U-1691			
<b><u>FLUTICASONE PROPIONATE - CUTIVATE</u></b>						
N 021152 001	7300669	Oct 20, 2019	DP U-835		NPP	Jan 16, 2018
<b><u>FLUTICASONE PROPIONATE - FLOVENT HFA</u></b>						
N 021433 001	6161724	Jan 16, 2018	DP			
	6170717	Dec 23, 2017	DP			
	6315173	Dec 23, 2017	DP			
	6431168	Jun 08, 2018	DP			
	6435372	Jan 16, 2018	DP			
	6510969	Dec 23, 2017	DP			
	6938796	Jan 16, 2018	DP			
	6966467	Dec 23, 2017	DP			
	6997349	Jan 16, 2018	DP			
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2018	DP			
	7143908	Jan 16, 2018	DP			
	7350676	Aug 24, 2018	DP			
	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026	DP			
	7832351	Jun 19, 2023	DP			
<b><u>FLUTICASONE PROPIONATE - FLOVENT HFA</u></b>						
N 021433 002	6161724	Jan 16, 2018	DP			
	6170717	Dec 23, 2017	DP			
	6315173	Dec 23, 2017	DP			
	6431168	Jun 08, 2018	DP			
	6435372	Jan 16, 2018	DP			
	6510969	Dec 23, 2017	DP			
	6743413	Jun 01, 2021	U-581		Y	
	6938796	Jan 16, 2018	DP			
	6966467	Dec 23, 2017	DP			
	6997349	Jan 16, 2018	DP			
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2018	DP			
	7143908	Jan 16, 2018	DP			
	7350676	Aug 24, 2018	DP			
	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026	DP			
	7832351	Jun 19, 2023	DP			
<b><u>FLUTICASONE PROPIONATE - FLOVENT HFA</u></b>						
N 021433 003	6161724	Jan 16, 2018	DP			
	6170717	Dec 23, 2017	DP			
	6315173	Dec 23, 2017	DP			
	6431168	Jun 08, 2018	DP			
	6435372	Jan 16, 2018	DP			
	6510969	Dec 23, 2017	DP			
	6938796	Jan 16, 2018	DP			
	6966467	Dec 23, 2017	DP			
	6997349	Jan 16, 2018	DP			
	7107986	Jun 08, 2018	DP			
	7107986*PED	Dec 08, 2018	DP			
	7143908	Jan 16, 2018	DP			
	7350676	Aug 24, 2018	DP			
	7500444	Feb 26, 2026	DP			
	7500444*PED	Aug 26, 2026	DP			
	7832351	Jun 19, 2023	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FLUTICASONE PROPIONATE - FLONASE ALLERGY RELIEF</u></b>						
N 205434 001					M-147	Jul 23, 2017
<b><u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u></b>						
N 208798 001	6446627	Dec 18, 2017	DP		NP	Jan 27, 2020
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			
<b><u>FLUTICASONE PROPIONATE - ARMONAIR RESPICLICK</u></b>						
N 208798 002	6446627	Dec 18, 2017	DP		NP	Jan 27, 2020
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			
<b><u>FLUTICASONE PROPIONATE - XHANCE</u></b>						
N 209022 001	6715485	Mar 03, 2020	DP		NP	Sep 18, 2020
	7975690	Dec 29, 2025	U-2133			
	8327844	Oct 08, 2023	U-2133			
	8522778	May 11, 2022	DP			
	8550073	Oct 22, 2029	DP			
	8555878	Mar 20, 2020	DP			
	8978647	Aug 06, 2030	DP			
	9072857	Apr 10, 2021	DP			
	9468727	Jul 30, 2020	DP			
<b><u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50</u></b>						
N 021077 001					M-214	Dec 20, 2020
<b><u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50</u></b>						
N 021077 002					M-214	Dec 20, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50</u>						
N 021077	003				M-214	Dec 20, 2020
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 021254	001	6161724 6170717 6315173 6431168 6435372 6510969 6938796 6966467 6997349 7107986 7143908 7350676 7500444 7500444*PED 7832351	Jan 16, 2018 Dec 23, 2017 Dec 23, 2017 Jun 08, 2018 Jan 16, 2018 Dec 23, 2017 Jan 16, 2018 Dec 23, 2017 Jan 16, 2018 Jun 08, 2018 Jan 16, 2018 Aug 24, 2018 Feb 26, 2026 Aug 26, 2026 Jun 19, 2023	DP DP DP DP DP DP DP DP DP DP DP DP DP DP DP DP		
N 021254	002	6161724 6170717 6315173 6431168 6435372 6510969 6938796 6966467 6997349 7107986 7143908 7350676 7500444 7500444*PED 7832351	Jan 16, 2018 Dec 23, 2017 Dec 23, 2017 Jun 08, 2018 Jan 16, 2018 Dec 23, 2017 Jan 16, 2018 Dec 23, 2017 Jan 16, 2018 Jun 08, 2018 Jan 16, 2018 Aug 24, 2018 Feb 26, 2026 Aug 26, 2026 Jun 19, 2023	DP DP DP DP DP DP DP DP DP DP DP DP DP DP DP		
N 021254	003	6161724 6170717 6315173 6431168 6435372 6510969 6938796 6966467 6997349 7107986 7143908 7350676 7500444 7500444*PED 7832351	Jan 16, 2018 Dec 23, 2017 Dec 23, 2017 Jun 08, 2018 Jan 16, 2018 Dec 23, 2017 Jan 16, 2018 Dec 23, 2017 Jan 16, 2018 Jun 08, 2018 Jan 16, 2018 Aug 24, 2018 Feb 26, 2026 Aug 26, 2026 Jun 19, 2023	DP DP DP DP DP DP DP DP DP DP DP DP DP DP DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>						
N 208799	001	6446627 6701917 6718972 6748947 6871646 7540282 8006690 8651103 8714149 8978966 9066957 9216260 9415008 9463288 9616024	Dec 18, 2017 Jun 23, 2021 Jun 23, 2021 Jun 23, 2021 Jun 23, 2021 May 06, 2023 Jun 23, 2021 Mar 26, 2028 Feb 25, 2032 Jan 13, 2032 Oct 06, 2034 Jun 28, 2031 Oct 06, 2034 May 19, 2025 Sep 01, 2024	DP DP DP DP DP DP DP DP DP DP DP DP DP DP DP DP	NP	Jan 27, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u></b>						
N 208799 001	9731087	May 18, 2031	DP			
<b><u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u></b>						
N 208799 002	6446627	Dec 18, 2017	DP		NP	Jan 27, 2020
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9066957	Oct 06, 2034	DP U-645			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			
<b><u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - AIRDUO RESPICLICK</u></b>						
N 208799 003	6446627	Dec 18, 2017	DP		NP	Jan 27, 2020
	6701917	Jun 23, 2021	DP			
	6718972	Jun 23, 2021	DP			
	6748947	Jun 23, 2021	DP			
	6871646	Jun 23, 2021	DP			
	7540282	May 06, 2023	DP			
	8006690	Jun 23, 2021	DP			
	8651103	Mar 26, 2028	DP			
	8714149	Feb 25, 2032	DP			
	8978966	Jan 13, 2032	DP			
	9066957	Oct 06, 2034	DP U-645			
	9216260	Jun 28, 2031	DP			
	9463288	May 19, 2025	DP			
	9616024	Sep 01, 2024	DP			
	9731087	May 18, 2031	DP			
<b><u>FLUVASTATIN SODIUM - LESCOL XL</u></b>						
N 021192 001	6242003	Apr 13, 2020				
<b><u>FLUVOXAMINE MALEATE - LUVOX CR</u></b>						
N 022033 001	7465462	May 10, 2020	DP U-929			
<b><u>FLUVOXAMINE MALEATE - LUVOX CR</u></b>						
N 022033 002	7465462	May 10, 2020	DP U-929			
<b><u>FOLLITROPIN ALFA/BETA - GONAL-F</u></b>						
N 020378 004	7563763	Aug 23, 2019	DP			
<b><u>FOLLITROPIN ALFA/BETA - GONAL-F</u></b>						
N 020378 005	7563763	Aug 23, 2019	DP			
<b><u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u></b>						
N 021211 001	5929028	Jan 14, 2018	DP U-1366			
	5929028	Jan 14, 2018	DP U-567			
	7446090	Aug 23, 2019	DP			
	7563763	Aug 23, 2019	U-1183			
	7563763	Aug 23, 2019	U-1367			
	7563763	Aug 23, 2019	U-993			
<b><u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u></b>						
N 021211 002	5929028	Jan 14, 2018	DP U-1366			
	5929028	Jan 14, 2018	DP U-567			
	7446090	Aug 23, 2019	DP			
	7563763	Aug 23, 2019	U-1183			
	7563763	Aug 23, 2019	U-1367			
	7563763	Aug 23, 2019	U-993			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u></b>						
N 021211 003	5929028	Jan 14, 2018	DP U-1366			
	5929028	Jan 14, 2018	DP U-567			
	7446090	Aug 23, 2019	DP			
	7563763	Aug 23, 2019	U-1183			
	7563763	Aug 23, 2019	U-1367			
	7563763	Aug 23, 2019	U-993			
<b><u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u></b>						
N 021211 004	5929028	Jan 14, 2018	DP U-1366			
	5929028	Jan 14, 2018	DP U-567			
	7446090	Aug 23, 2019	DP			
	7563763	Aug 23, 2019	U-1183			
	7563763	Aug 23, 2019	U-1367			
	7563763	Aug 23, 2019	U-993			
<b><u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u></b>						
N 021273 001	5929028	Jan 14, 2018	DP U-1366			
<b><u>FOLLITROPIN ALFA/BETA - FOLLISTIM AQ</u></b>						
N 021273 002	5929028	Jan 14, 2018	DP U-1366			
<b><u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u></b>						
N 021684 001	7446090	Aug 23, 2019	DP			
	7741268	Apr 02, 2024	DP			
<b><u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u></b>						
N 021684 002	7446090	Aug 23, 2019	DP			
	7741268	Apr 02, 2024	DP			
<b><u>FOLLITROPIN ALFA/BETA - GONAL-F RFF REDI-JECT</u></b>						
N 021684 003	7446090	Aug 23, 2019	DP			
	7741268	Apr 02, 2024	DP			
<b><u>FOMEPIZOLE - ANTIZOL</u></b>						
N 020696 001	7553863	Jun 30, 2027	DS DP			
<b><u>FORMOTEROL FUMARATE - FORADIL</u></b>						
N 020831 001	6488027	Mar 08, 2019				
	6887459	Nov 28, 2020	U-762			
<b><u>FORMOTEROL FUMARATE - PERFOROMIST</u></b>						
N 022007 001	6667344	Jun 22, 2021	DP			
	6814953	Jun 22, 2021	DP U-813			
	7348362	Jun 22, 2021	DP			
	7462645	Jun 22, 2021	DP U-813			
	8623922	Jun 22, 2021	DP			
<b><u>FORMOTEROL FUMARATE; GLYCOPYRROLATE - BEVESPI AEROSPHERE</u></b>						
N 208294 001	8324266	May 28, 2030	U-1841		NP	Apr 25, 2019
	8703806	May 28, 2030	U-1841			
	8808713	May 28, 2030	DP U-1841			
	8815258	Mar 17, 2031	U-1841			
	9415009	May 28, 2030	U-1841			
	9463161	May 28, 2030	DP U-1841			
<b><u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u></b>						
N 022518 001	6068832	Aug 27, 2017	DP U-1068		M-214	Dec 20, 2020
	7067502	May 21, 2020	DP U-1068			
	7566705	May 21, 2020	DP U-1068			
<b><u>FORMOTEROL FUMARATE; MOMETASONE FUROATE - DULERA</u></b>						
N 022518 002	6068832	Aug 27, 2017	DP U-1068		M-214	Dec 20, 2020
	7067502	May 21, 2020	DP U-1068			
	7566705	May 21, 2020	DP U-1068			
<b><u>FOSAMPRENAVIR CALCIUM - LEXIVA</u></b>						
N 021548 001	6436989	Dec 24, 2017	DS DP U-257			
	6514953	Jul 15, 2019	DS DP U-257			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>FOSAMPRENAVIR CALCIUM - LEXIVA</u></b>						
N 022116 001	6436989	Dec 24, 2017	DS DP U-257			
<b><u>FOSAPREPITANT DIMEGLUMINE - EMEND</u></b>						
N 022023 001	5691336	Mar 04, 2019	DS DP			
<b><u>FOSAPREPITANT DIMEGLUMINE - EMEND</u></b>						
N 022023 002	5691336	Mar 04, 2019	DS DP		D-155	Feb 01, 2019
<b><u>FOSPROPOFOL DISODIUM - LUSEDRA</u></b>						
N 022244 001	6204257	Jul 01, 2022	DS DP U-945			
	6872838	Aug 07, 2018	DS			
<b><u>FULVESTRANT - FASLODEX</u></b>						
N 021344 001	6774122	Jan 09, 2021	U-1826		I-725	Feb 19, 2019
	6774122	Jan 09, 2021	U-2108		I-749	Aug 25, 2020
	6774122	Jan 09, 2021	U-2163			
	6774122	Jan 09, 2021	U-596			
	6774122*PED	Jul 09, 2021				
	7456160	Jan 09, 2021	U-1826			
	7456160	Jan 09, 2021	U-2108			
	7456160	Jan 09, 2021	U-2163			
	7456160	Jan 09, 2021	U-596			
	7456160*PED	Jul 09, 2021				
	8329680	Jan 09, 2021	U-1826			
	8329680	Jan 09, 2021	U-2108			
	8329680	Jan 09, 2021	U-2163			
	8329680	Jan 09, 2021	U-596			
	8329680*PED	Jul 09, 2021				
	8466139	Jan 09, 2021	U-1826			
	8466139	Jan 09, 2021	U-2108			
	8466139	Jan 09, 2021	U-2163			
	8466139	Jan 09, 2021	U-596			
	8466139*PED	Jul 09, 2021				
<b><u>GABAPENTIN - NEURONTIN</u></b>						
N 021129 001	7256216	May 28, 2022	DP			
<b><u>GABAPENTIN - GRALISE</u></b>						
N 022544 001	6488962	Jun 20, 2020	DP		ODE-6	Jan 28, 2018
	6723340	Oct 25, 2021	DP			
	7438927	Feb 26, 2024	U-1114			
	7731989	Oct 25, 2022	DP			
	8192756	Oct 25, 2022	DP U-1114			
	8252332	Oct 25, 2022	DP U-1114			
	8333992	Oct 25, 2022	DP U-1114			
<b><u>GABAPENTIN - GRALISE</u></b>						
N 022544 002	6488962	Jun 20, 2020	DP		ODE-6	Jan 28, 2018
	6723340	Oct 25, 2021	DP			
	7438927	Feb 26, 2024	U-1114			
	7731989	Oct 25, 2022	DP			
	8192756	Oct 25, 2022	DP U-1114			
	8252332	Oct 25, 2022	DP U-1114			
	8333992	Oct 25, 2022	DP U-1114			
<b><u>GABAPENTIN ENACARBIL - HORIZANT</u></b>						
N 022399 001	6818787	Nov 06, 2022	DS DP		ODE-25	Jun 06, 2019
	8026279	Nov 10, 2026	DS DP			
	8048917	Nov 06, 2022	DS DP U-1247			
	8114909	Apr 11, 2026	U-1231			
	8686034	Jan 24, 2025	U-1231			
	8686034	Jan 24, 2025	U-1247			
	8795725	Jun 10, 2029	DP U-1231			
	8795725	Jun 10, 2029	DP U-1247			
<b><u>GABAPENTIN ENACARBIL - HORIZANT</u></b>						
N 022399 002	6818787	Nov 06, 2022	DS DP		ODE-25	Jun 06, 2019
	8026279	Nov 10, 2026	DS DP			
	8048917	Nov 06, 2022	DS DP U-1247			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>GABAPENTIN ENACARBIL - HORIZANT</b>						
N 022399 002	8114909	Apr 11, 2026	U-1231			
	8686034	Jan 24, 2025	U-1231			
	8686034	Jan 24, 2025	U-1247			
	8795725	Jun 10, 2029	DP U-1231			
	8795725	Jun 10, 2029	DP U-1247			
<b>GADOBUTROL - GADAVIST</b>						
N 201277 002				I-731		Apr 27, 2019
<b>GADOBUTROL - GADAVIST</b>						
N 201277 006	5980864	Nov 09, 2021	DS DP U-1119			
<b>GADOFOSVESET TRISODIUM - ABLAVAR</b>						
N 021711 001	6676929	May 04, 2020	DP			
<b>GADOFOSVESET TRISODIUM - ABLAVAR</b>						
N 021711 002	6676929	May 04, 2020	DP			
<b>GADOTERATE MEGLUMINE - DOTAREM</b>						
N 204781 001				NCE		Mar 20, 2018
				NPP		Aug 25, 2020
<b>GADOTERATE MEGLUMINE - DOTAREM</b>						
N 204781 002				NCE		Mar 20, 2018
				NPP		Aug 25, 2020
<b>GADOTERATE MEGLUMINE - DOTAREM</b>						
N 204781 003				NCE		Mar 20, 2018
				NPP		Aug 25, 2020
<b>GADOTERATE MEGLUMINE - DOTAREM</b>						
N 204781 004				NCE		Mar 20, 2018
				NPP		Aug 25, 2020
<b>GADOTERATE MEGLUMINE - DOTAREM</b>						
N 204781 005				NCE		Mar 20, 2018
				NPP		Aug 25, 2020
<b>GADOXETATE DISODIUM - EOVIST</b>						
N 022090 001	6039931	Nov 13, 2021	U-1239		M-155	Mar 27, 2018
<b>GADOXETATE DISODIUM - EOVIST</b>						
N 022090 002					M-155	Mar 27, 2018
<b>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</b>						
N 021615 001	7160559	Dec 20, 2019	DP			
<b>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</b>						
N 021615 002	7160559	Dec 20, 2019	DP			
<b>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</b>						
N 021615 003	7160559	Dec 20, 2019	DP			
<b>GALLIUM DOTATATE GA-68 - NETSPOT</b>						
N 208547 001					NCE	Jun 01, 2021
					ODE-120	Jun 01, 2023
<b>GANCICLOVIR - GANCICLOVIR</b>						
N 209347 001	9486530	Sep 02, 2034	DP			
<b>GANIRELIX ACETATE - GANIRELIX ACETATE</b>						
N 021057 001	6653286	Jun 16, 2018	U-1354			
<b>GATIFLOXACIN - ZYMAR</b>						
N 021493 001	6333045	Aug 20, 2019	DP		Y	
	6333045*PED	Feb 20, 2020				
<b>GEFITINIB - IRESSA</b>						
N 206995 001				NP		Jul 13, 2018
				ODE-95		Jul 13, 2022

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>GEFITINIB - IRESSA</u></b>						
N 206995	001				NP ODE-95	Jul 13, 2018 Jul 13, 2022
<b><u>GEMIFLOXACIN MESYLATE - FACTIVE</u></b>						
N 021158	001	6262071 6331550 6340689 6455540 6723734 6803376 6803376	Sep 21, 2019 Sep 21, 2019 Sep 14, 2019 Sep 21, 2019 Mar 20, 2018 Sep 21, 2019 Sep 21, 2019	U-513 U-511 U-512 U-511 DS DP DS DP U-608 DS DP U-609		
<b><u>GLATIRAMER ACETATE - COPAXONE</u></b>						
N 020622	003	8232250 8399413 8969302 9155776 9402874	Aug 19, 2030 Aug 19, 2030 Aug 19, 2030 Aug 19, 2030 Aug 19, 2030	U-441 U-441 U-441 U-441 U-441		
<b><u>GLECAPREVIR; PIBRENTASVIR - MAVYRET</u></b>						
N 209394	001	8648037 8937150 9321807 9586978	Jan 19, 2032 May 18, 2032 Jun 05, 2035 Jun 10, 2030	DS DP U-2141 DS DP DS U-2141	NCE	Aug 03, 2022
<b><u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u></b>						
N 021925	001	7700128 8071130	Jan 30, 2027 Jun 08, 2028	DP DP		
<b><u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u></b>						
N 021925	002	7700128 8071130	Jan 30, 2027 Jun 08, 2028	DP DP		
<b><u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u></b>						
N 021700	001	7358366	Apr 19, 2020	DS		
<b><u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u></b>						
N 021700	002	7358366	Apr 19, 2020	DS		
<b><u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u></b>						
N 021700	003	7358366	Apr 19, 2020	DS		
<b><u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u></b>						
N 021700	004	7358366	Apr 19, 2020	DS		
<b><u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u></b>						
N 021700	005	7358366	Apr 19, 2020	DS		
<b><u>GLIPIZIDE - GLUCOTROL XL</u></b>						
N 020329	001	RE44459	Mar 26, 2019	U-1431		
<b><u>GLIPIZIDE - GLUCOTROL XL</u></b>						
N 020329	002	RE44459	Mar 26, 2019	U-1431		
<b><u>GLIPIZIDE - GLUCOTROL XL</u></b>						
N 020329	003	RE44459	Mar 26, 2019	U-1431		
<b><u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u></b>						
N 021178	001	6303146	Jul 14, 2019	U-412		
<b><u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u></b>						
N 021178	002	6303146	Jul 14, 2019	U-412		
<b><u>GLYBURIDE; METFORMIN HYDROCHLORIDE - GLUCOVANCE</u></b>						
N 021178	003	6303146	Jul 14, 2019	U-412		
<b><u>GLYCEROL PHENYLBUTYRATE - RAVICTI</u></b>						
N 203284	001	5968979 8404215 8642012	Jul 28, 2018 Mar 09, 2032 Sep 22, 2030	DS DP U-1378 U-1383 U-1383	NPP ODE ODE-42	Apr 28, 2020 Apr 28, 2024 Feb 01, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>GLYCEROL PHENYLBUTYRATE - RAVICTI</u></b>						
N 203284 001	9095559	Mar 09, 2032	U-1383			
	9254278	Mar 09, 2032	U-1816			
	9326966	Mar 09, 2032	U-1816			
	9561197	Sep 22, 2030	U-1383			
<b><u>GLYCOPYRROLATE - CUVPOSA</u></b>						
N 022571 001	7638552	Aug 20, 2023	U-1076		ODE-1	Jul 28, 2017
	7816396	Aug 20, 2023	U-1076			
<b><u>GLYCOPYRROLATE - SEEBRI</u></b>						
N 207923 001	7229607	Apr 09, 2021	U-1773		NP	Oct 29, 2018
	7736670	Jun 27, 2021	DP			
	8029768	Apr 09, 2021	U-1773			
	8048451	Jun 27, 2021	DP			
	8182838	Oct 20, 2028	DP			
	8303991	Jun 27, 2021	DP			
	8435567	Jun 27, 2021	DP			
	8479730	Oct 11, 2028	DP			
	8580306	Jun 27, 2021	DP			
	8956661	Jun 27, 2021	DP			
<b><u>GLYCOPYRROLATE - LONHALA MAGNAIR KIT</u></b>						
N 208437 001	6962151	Oct 27, 2020	DP		NP	Dec 05, 2020
	7316067	Sep 06, 2022	DP			
	7458372	Nov 18, 2024	DP			
	7931212	Nov 25, 2025	DP			
	8511581	Nov 08, 2023	DP			
	9168556	Sep 01, 2032	DP			
	9265900	Dec 07, 2028	DP			
	9604018	May 16, 2033	DP			
	9789270	Oct 30, 2030	DP			
<b><u>GLYCOPYRROLATE ; INDACATEROL MALEATE - UTIBRON</u></b>						
N 207930 001	6878721	Feb 25, 2025	DS DP U-1773		NP	Oct 29, 2018
	7229607	Apr 09, 2021	U-1773			
	7736670	Jun 27, 2021	DP			
	7820694	Jun 02, 2020	DP U-1773			
	8029768	Apr 09, 2021	U-1773			
	8048451	Jun 27, 2021	DP			
	8067437	Jun 02, 2020	U-1773			
	8182838	Oct 20, 2028	DP			
	8283362	Jun 02, 2020	DP U-1773			
	8303991	Jun 27, 2021	DP			
	8435567	Jun 27, 2021	DP			
	8479730	Oct 11, 2028	DP			
	8580306	Jun 27, 2021	DP			
	8658673	Jun 02, 2020	DP U-1773			
	8796307	Jun 02, 2020	DP			
	8956661	Jun 27, 2021	DP			
<b><u>GOSERELIN ACETATE - ZOLADEX</u></b>						
N 019726 001	7118552	Apr 13, 2022	DP			
	7220247	Apr 09, 2022	DP			
	7500964	Feb 26, 2021	DP			
<b><u>GOSERELIN ACETATE - ZOLADEX</u></b>						
N 020578 001	7118552	Apr 13, 2022	DP			
	7220247	Apr 09, 2022	DP			
	7500964	Feb 26, 2021	DP			
<b><u>GRANISETRON - SANCUSO</u></b>						
N 022198 001	7608282	Jan 22, 2025	DP U-1011			
<b><u>GRANISETRON - SUSTOL</u></b>						
N 022445 001	6613355	Jun 28, 2021	DP		NDF	Aug 09, 2019
	6790458	May 11, 2021	DP			
	8252304	Sep 28, 2024	DP			
	8252305	Sep 28, 2024	U-1891			
	8715710	Sep 28, 2024	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>GRANISETRON - SUSTOL</u></b>						
N 022445 001	6613355	Jun 28, 2021	DP		NDF	Aug 09, 2019
	6790458	May 11, 2021	DP			
	8252304	Sep 28, 2024	DP			
	8252305	Sep 28, 2024	U-1891			
	8715710	Sep 28, 2024	DP			
<b><u>GUAIFENESIN - MUCINEX</u></b>						
N 021282 001	6372252	Apr 28, 2020	U-489			
	6955821	Apr 28, 2020	DP U-489			
	7838032	Apr 28, 2020	DP			
<b><u>GUAIFENESIN - MUCINEX</u></b>						
N 021282 002	6372252	Apr 28, 2020	U-489			
	6955821	Apr 28, 2020	DP U-489			
	7838032	Apr 28, 2020	DP			
<b><u>GUAIFENESIN; HYDROCODONE BITARTRATE - OBREDON</u></b>						
N 205474 001	9549907	Nov 13, 2035	DS DP	U-2023		
<b><u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u></b>						
N 021585 001	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP U-686			
	7838032	Apr 28, 2020	DP			
<b><u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE - MUCINEX D</u></b>						
N 021585 002	6372252	Apr 28, 2020	DP			
	6955821	Apr 28, 2020	DP U-686			
	7838032	Apr 28, 2020	DP			
<b><u>GUANFACINE HYDROCHLORIDE - INTUNIV</u></b>						
N 022037 001	6287599	Dec 20, 2020	DP		D-145	Nov 19, 2017
	6287599*PED	Jun 20, 2021			M-154	Mar 18, 2018
	6811794	Jul 04, 2022	DP U-494		PED	May 19, 2018
	6811794*PED	Jan 04, 2023				
<b><u>GUANFACINE HYDROCHLORIDE - INTUNIV</u></b>						
N 022037 002	6287599	Dec 20, 2020	DP		D-145	Nov 19, 2017
	6287599*PED	Jun 20, 2021			M-154	Mar 18, 2018
	6811794	Jul 04, 2022	DP U-494		PED	May 19, 2018
	6811794*PED	Jan 04, 2023				
<b><u>GUANFACINE HYDROCHLORIDE - INTUNIV</u></b>						
N 022037 003	6287599	Dec 20, 2020	DP		D-145	Nov 19, 2017
	6287599*PED	Jun 20, 2021			M-154	Mar 18, 2018
	6811794	Jul 04, 2022	DP U-494		PED	May 19, 2018
	6811794*PED	Jan 04, 2023				
<b><u>GUANFACINE HYDROCHLORIDE - INTUNIV</u></b>						
N 022037 004	6287599	Dec 20, 2020	DP		D-145	Nov 19, 2017
	6287599*PED	Jun 20, 2021			M-154	Mar 18, 2018
	6811794	Jul 04, 2022	DP U-494		PED	May 19, 2018
	6811794*PED	Jan 04, 2023				
<b><u>HALOBETASOL PROPIONATE - ULTRAVATE</u></b>						
N 208183 001	8962028	Jun 19, 2033	DP U-1775		NP	Nov 06, 2018
<b><u>HEXAMINOLEVULINATE HYDROCHLORIDE - CYSVIEW KIT</u></b>						
N 022555 001	7348361	Nov 06, 2020	DP U-1087			
<b><u>HISTRELIN ACETATE - SUPPRELIN LA</u></b>						
N 022058 001	8062652	Jun 16, 2026		U-1197		
<b><u>HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT</u></b>						
N 021859 001	7767429	Sep 23, 2027	DS DP			
<b><u>HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE - BIDIL</u></b>						
N 020727 001	6465463	Sep 08, 2020	U-71			
	6784177	Sep 08, 2020	U-71			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u></b>						
N 021162 001	6358986	Jan 10, 2020				
<b><u>HYDROCHLOROTHIAZIDE; TELMISARTAN - MICARDIS HCT</u></b>						
N 021162 002	6358986	Jan 10, 2020				
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 001	6228398	Nov 01, 2019	DP			
	6902742	Nov 01, 2019	DP			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 002	6228398	Nov 01, 2019	DP			
	6902742	Nov 01, 2019	DP			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 003	6228398	Nov 01, 2019	DP			
	6902742	Nov 01, 2019	DP			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 004	6228398	Nov 01, 2019	DP			
	6902742	Nov 01, 2019	DP			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 005	6228398	Nov 01, 2019	DP			
	6902742	Nov 01, 2019	DP			
	9132096	Sep 12, 2034	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 005	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<b><u>HYDROCODONE BITARTRATE - ZOHYDRO ER</u></b>						
N 202880 006	6228398	Nov 01, 2019	DP			
	6902742	Nov 01, 2019	DP			
	9132096	Sep 12, 2034	DP			
	9265760	Jul 25, 2033	U-1810			
	9326982	Jul 25, 2033	U-1810			
	9333201	Jul 25, 2033	U-1810			
	9339499	Jul 25, 2033	U-1810			
	9421200	Jul 25, 2033	U-1810			
	9433619	Jul 25, 2033	U-1810			
	9452163	Sep 12, 2034	U-55			
	9486451	Sep 12, 2034	U-55			
	9610286	Jul 25, 2033	U-1810			
	9713611	Sep 12, 2034	DP U-55			
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 001	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027	U-1556			
	9492391	Aug 24, 2027	U-1556			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020	U-1556			
	9682077	Oct 30, 2020	U-1556			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 002	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 002	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020		U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027		U-1556		
	9492391	Aug 24, 2027		U-1556		
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020		U-1556		
	9682077	Oct 30, 2020		U-1556		
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 003	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020		U-1556		
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027		U-1556		
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027		U-1556		
	9492391	Aug 24, 2027		U-1556		
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027		U-1556		
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020		U-1556		
	9682077	Oct 30, 2020		U-1556		
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 004	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9023401	Oct 30, 2020	DP			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027	U-1556			
	9492391	Aug 24, 2027	U-1556			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020	U-1556			
	9682077	Oct 30, 2020	U-1556			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 005	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027	U-1556			
	9492391	Aug 24, 2027	U-1556			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020	U-1556			
	9682077	Oct 30, 2020	U-1556			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 005	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 006	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027	U-1556			
	9492391	Aug 24, 2027	U-1556			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2020	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			
	9675611	Oct 30, 2020	U-1556			
	9682077	Oct 30, 2020	U-1556			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
<b><u>HYDROCODONE BITARTRATE - HYSINGLA</u></b>						
N 206627 007	6733783	Oct 30, 2021	DP U-1556			
	8309060	Nov 20, 2023	DP U-1556			
	8361499	Oct 30, 2021	DP			
	8529948	Aug 06, 2022	DP			
	8551520	Oct 30, 2021	DP			
	8647667	Oct 30, 2021	DP			
	8808740	Dec 21, 2031	DP U-1556			
	9056052	Oct 30, 2020	DP			
	9060940	Oct 30, 2020	U-1556			
	9084816	Aug 24, 2027	DP			
	9095614	Aug 24, 2027	U-1556			
	9095615	Aug 24, 2027	DP			
	9198863	Oct 30, 2020	DP			
	9205056	Oct 30, 2020	DP			
	9289391	Oct 30, 2020	DP			
	9486412	Aug 24, 2027	DP			
	9486413	Aug 24, 2027	DP			
	9492389	Aug 24, 2027	DP			
	9492390	Aug 24, 2027	U-1556			
	9492391	Aug 24, 2027	U-1556			
	9517236	Oct 30, 2020	DP			
	9545380	Aug 24, 2027	U-1556			
	9572779	Dec 21, 2031	DP			
	9572804	Oct 30, 2021	DP			
	9669023	Oct 30, 2020	DP			
	9669024	Oct 30, 2020	DP			
	9675610	Jun 16, 2023	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROCODONE BITARTRATE - HYSTINGLA</u></b>						
N 206627 007	9675611	Oct 30, 2020	U-1556			
	9682077	Oct 30, 2020	U-1556			
	9750703	Dec 21, 2031	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775809	Aug 24, 2027	DP			
	9861584	Dec 21, 2031	DP			
<b><u>HYDROCODONE BITARTRATE - VANTRELA ER</u></b>						
N 207975 001	8445018	Jul 31, 2029	DP			
	9216176	Sep 13, 2027	DP			
	9572803	Sep 13, 2027	DP			
<b><u>HYDROCODONE BITARTRATE - VANTRELA ER</u></b>						
N 207975 002	8445018	Jul 31, 2029	DP			
	9216176	Sep 13, 2027	DP			
	9572803	Sep 13, 2027	DP			
<b><u>HYDROCODONE BITARTRATE - VANTRELA ER</u></b>						
N 207975 003	8445018	Jul 31, 2029	DP			
	9216176	Sep 13, 2027	DP			
	9572803	Sep 13, 2027	DP			
<b><u>HYDROCODONE BITARTRATE - VANTRELA ER</u></b>						
N 207975 004	8445018	Jul 31, 2029	DP			
	9216176	Sep 13, 2027	DP			
	9572803	Sep 13, 2027	DP			
<b><u>HYDROCODONE BITARTRATE - VANTRELA ER</u></b>						
N 207975 005	8445018	Jul 31, 2029	DP			
	9216176	Sep 13, 2027	DP			
	9572803	Sep 13, 2027	DP			
<b><u>HYDROCORTISONE BUTYRATE - LOCOID</u></b>						
N 022076 001	7378405	Dec 19, 2026	DP			
	7981877	Jan 23, 2025	DP			
<b><u>HYDROGEN PEROXIDE - ESKATA</u></b>						
N 209305 001	7381427	Jun 08, 2022	U-2205		NP	
	9675639	Jul 04, 2035	DP U-2205			Dec 14, 2020
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u></b>						
N 019034 001	6589960	Nov 09, 2020	DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID-HP</u></b>						
N 019034 002	6589960	Nov 09, 2020	DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019034 003	6589960	Nov 09, 2020	DS DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019034 004	6589960	Nov 09, 2020	DS DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019034 005	6589960	Nov 09, 2020	DS DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019891 001	6589960	Nov 09, 2020	DS DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019892 001	6589960	Nov 09, 2020	DS DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019892 002	6589960	Nov 09, 2020	DS DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - DILAUDID</u></b>						
N 019892 003	6589960	Nov 09, 2020	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u></b>						
N 021044 001	6589960	Nov 09, 2020	DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u></b>						
N 021044 002	6589960	Nov 09, 2020	DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u></b>						
N 021044 003	6589960	Nov 09, 2020	DP			
<b><u>HYDROMORPHONE HYDROCHLORIDE - PALLADONE</u></b>						
N 021044 004	6589960	Nov 09, 2020	DP			
<b><u>HYDROXYPROGESTERONE CAPROATE - MAKENA</u></b>						
N 021945 001					ODE-7	Feb 03, 2018
<b><u>HYDROXYPROGESTERONE CAPROATE - MAKENA PRESERVATIVE FREE</u></b>						
N 021945 002					ODE-7	Feb 03, 2018
<b><u>HYDROXYUREA - SIKLOS</u></b>						
N 208843 001					NP	Dec 21, 2020
<b><u>HYDROXYUREA - SIKLOS</u></b>						
N 208843 002					NP	Dec 21, 2020
<b><u>IBANDRONATE SODIUM - BONIVA</u></b>						
N 021455 001	6294196	Oct 07, 2019	DP			
<b><u>IBANDRONATE SODIUM - BONIVA</u></b>						
N 021455 002	6294196	Oct 07, 2019	DP			
7192938		May 06, 2023	U-798			
7410957		May 06, 2023	U-887			
7718634		May 06, 2023	U-642			
<b><u>IBRUTINIB - IMBRUVICA</u></b>						
N 205552 001	7514444	Dec 28, 2026	DS DP		D-165	May 06, 2019
8008309		Dec 28, 2026	DS DP		I-689	Jul 28, 2017
8476284		Dec 28, 2026	U-1456		I-702	Jan 29, 2018
8476284		Dec 28, 2026	U-1650		I-729	Mar 04, 2019
8476284		Dec 28, 2026	U-1946		I-736	May 06, 2019
8476284		Dec 28, 2026	U-1947		I-737	May 06, 2019
8497277		Dec 28, 2026	U-1456		I-741	Jan 18, 2020
8497277		Dec 28, 2026	U-1491		I-753	Aug 02, 2020
8497277		Dec 28, 2026	U-1650		NCE	Nov 13, 2018
8497277		Dec 28, 2026	U-1946		ODE-109	Mar 04, 2023
8497277		Dec 28, 2026	U-1947		ODE-117	May 06, 2023
8697711		Dec 28, 2026	DS DP		ODE-128	Jan 18, 2024
8703780		Dec 28, 2026	U-1491		ODE-152	Aug 02, 2024
8735403		Dec 28, 2026	DS DP		ODE-55	Nov 13, 2020
8754090		Jun 03, 2031	U-1456		ODE-60	Feb 12, 2021
8754091		Dec 28, 2026	DP		ODE-72	Jul 28, 2021
8952015		Dec 28, 2026	U-1456		ODE-86	Jan 29, 2022
8952015		Dec 28, 2026	U-1491			
8952015		Dec 28, 2026	U-1650			
8952015		Dec 28, 2026	U-1946			
8952015		Dec 28, 2026	U-1947			
8957079		Dec 28, 2026	DS DP			
8999999		Jun 03, 2031	U-1683			
8999999		Jun 03, 2031	U-1684			
9125889		Jun 03, 2031	U-1745			
9181257		Dec 28, 2026	DS DP			
9296753		Oct 30, 2033	DS DP			
9540382		Aug 18, 2033	U-1456			
9540382		Aug 18, 2033	U-1650			
9540382		Aug 18, 2033	U-1684			
9540382		Aug 18, 2033	U-1946			
9540382		Aug 18, 2033	U-1947			
9713617		Jun 03, 2033	DP			
9725455		Jun 03, 2033	DS			
9795604		Oct 24, 2034	U-2150			
9801881		Jun 03, 2031	U-1491			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>IBRUTINIB - IMBRUVICA</u></b>						
N 205552 001	9801883	Jun 03, 2031	U-2159			
	9814721	Jun 03, 2031	U-1947			
<b><u>IBUPROFEN - MIDOL LIQUID GELS</u></b>						
N 021472 001	6251426	Jun 25, 2018				
<b><u>IBUPROFEN - CALDOLOR</u></b>						
N 022348 001	6727286	Nov 27, 2021	DP U-981			
	8735452	Sep 30, 2029	U-981		D-152	Nov 20, 2018
	8871810	Sep 30, 2029	U-1599		NPP	Nov 20, 2018
	9012508	Sep 14, 2030	U-981			
	9114068	Sep 30, 2029	U-1735			
	9138404	Sep 30, 2029	U-1756			
	9295639	Sep 30, 2029	U-1756			
	9649284	Sep 30, 2029	U-2018			
<b><u>IBUPROFEN LYSINE - NEOPROFEN</u></b>						
N 021903 001	6342530	Nov 14, 2020	DP U-1127			
	6342530	Nov 14, 2020	DP U-794			
	6344479	Mar 20, 2021	DS DP U-794	Y		
	8415337	Mar 02, 2032	DS DP			
<b><u>IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE - CHILDREN'S MOTRIN COLD</u></b>						
N 021128 001	6211246	Jun 10, 2019				
<b><u>ICATIBANT ACETATE - FIRAZYR</u></b>						
N 022150 001	5648333	Jul 15, 2019	DS DP U-1187		ODE-14	Aug 25, 2018
<b><u>ICODEXTRIN - EXTRANEAL</u></b>						
N 021321 001	6248726	Jun 19, 2018	U-495			
<b><u>ICOSAPENT ETHYL - VASCEPA</u></b>						
N 202057 001	8188146	Jan 27, 2020	DS DP		NCE	Jul 26, 2017
	8293727	Feb 09, 2030	U-1287			
	8293728	Feb 09, 2030	U-1287			
	8298554	Apr 29, 2030	DP			
	8314086	Feb 09, 2030	U-1287			
	8318715	Feb 09, 2030	U-1287			
	8357677	Feb 09, 2030	U-1287			
	8367652	Feb 09, 2030	U-1287			
	8377920	Feb 09, 2030	U-1287			
	8399446	Feb 09, 2030	U-1287			
	8415335	Feb 09, 2030	U-1287			
	8426399	Feb 09, 2030	U-1287			
	8431560	Feb 09, 2030	U-1287			
	8440650	Feb 09, 2030	U-1287			
	8445003	Apr 29, 2030	U-1287			
	8445013	Apr 29, 2030	U-1287			
	8501225	Apr 29, 2030	U-1287			
	8518929	Apr 29, 2030	U-1287			
	8524698	Apr 29, 2030	U-1287			
	8546372	Apr 29, 2030	U-1287			
	8551521	Apr 29, 2030	U-1287			
	8563608	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-1478			
	8617594	Apr 29, 2030	U-1287			
	8623406	Apr 29, 2030	U-1478			
<b><u>ICOSAPENT ETHYL - VASCEPA</u></b>						
N 202057 002	8188146	Jan 27, 2020	DS DP		NCE	Jul 26, 2017
	8293727	Feb 09, 2030	U-1287			
	8293728	Feb 09, 2030	U-1287			
	8298554	Apr 29, 2030	DP			
	8314086	Feb 09, 2030	U-1287			
	8318715	Feb 09, 2030	U-1287			
	8357677	Feb 09, 2030	U-1287			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ICOSAPENT ETHYL - VASCEPA</u></b>						
N 202057 002	8367652	Feb 09, 2030	U-1287			
	8377920	Feb 09, 2030	U-1287			
	8399446	Feb 09, 2030	U-1287			
	8415335	Feb 09, 2030	U-1287			
	8426399	Feb 09, 2030	U-1287			
	8440650	Feb 09, 2030	U-1287			
	8445003	Apr 29, 2030	U-1287			
	8445013	Apr 29, 2030	U-1287			
	8501225	Apr 29, 2030	U-1287			
	8518929	Apr 29, 2030	U-1287			
	8524698	Apr 29, 2030	U-1287			
	8546372	Apr 29, 2030	U-1287			
	8551521	Apr 29, 2030	U-1287			
	8563608	Apr 29, 2030	U-1287			
	8617593	Apr 29, 2030	U-1287			
	8617594	Apr 29, 2030	U-1287			
	8623406	Apr 29, 2030	U-1287			
<b><u>IDEALALISIB - ZYDELIG</u></b>						
N 205858 001	6800620	Apr 24, 2021	DS	U-1560	NCE	Jul 23, 2019
	6949535	Apr 24, 2021	DS	U-1560	ODE-70	Jul 23, 2021
	8138195	Apr 24, 2021	DS DP	U-1549	ODE-71	Jul 23, 2021
	8492389	Apr 24, 2021	DS DP			
	8637533	Apr 24, 2021	DS DP			
	8865730	Mar 05, 2033	DS DP	U-1615		
	8980901	May 12, 2025		U-1678		
	9149477	May 12, 2025		U-1757		
	9469643	Sep 02, 2033	DS			
	9492449	Mar 11, 2030		U-1914		
	RE44599	Jul 21, 2025		U-1558		
	RE44599	Jul 21, 2025		U-1615		
	RE44638	Aug 05, 2025	DS DP			
<b><u>IDEALALISIB - ZYDELIG</u></b>						
N 205858 002	6800620	Apr 24, 2021	DS	U-1560	NCE	Jul 23, 2019
	6949535	Apr 24, 2021	DS	U-1560	ODE-70	Jul 23, 2021
	8138195	Apr 24, 2021	DS DP	U-1549	ODE-71	Jul 23, 2021
	8492389	Apr 24, 2021	DS DP			
	8637533	Apr 24, 2021	DS DP			
	8865730	Mar 05, 2033	DS DP	U-1615		
	8980901	May 12, 2025		U-1678		
	9149477	May 12, 2025		U-1757		
	9469643	Sep 02, 2033	DS			
	9492449	Mar 11, 2030		U-1914		
	RE44599	Jul 21, 2025		U-1558		
	RE44599	Jul 21, 2025		U-1615		
	RE44638	Aug 05, 2025	DS DP			
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 001	8586610	Nov 02, 2027		U-1625	M-180	May 26, 2019
	8652776	Aug 31, 2030		U-1685		
	8999638	Oct 28, 2030		U-1674		
	9072742	Jan 16, 2031		U-1674		
	9074254	Dec 28, 2031		U-1674		
	9074255	Dec 17, 2030		U-1674		
	9074256	Feb 10, 2031		U-1674		
	9138432	Sep 30, 2025		U-1737		
	9157121	Apr 05, 2030		U-1674		
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 002	8586610	Nov 02, 2027		U-1625	M-180	May 26, 2019
	8652776	Aug 31, 2030		U-1685		
	8999638	Oct 28, 2030		U-1674		
	9072742	Jan 16, 2031		U-1674		
	9074254	Dec 28, 2031		U-1674		
	9074255	Dec 17, 2030		U-1674		
	9074256	Feb 10, 2031		U-1674		
	9138432	Sep 30, 2025		U-1737		
	9157121	Apr 05, 2030		U-1674		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 002	8586610	Nov 02, 2027	U-1625		M-180	May 26, 2019
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 003	8586610	Nov 02, 2027	U-1625		M-180	May 26, 2019
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 004	8586610	Nov 02, 2027	U-1625		M-180	May 26, 2019
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 005	8586610	Nov 02, 2027	U-1625		M-180	May 26, 2019
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 006	8586610	Nov 02, 2027	U-1625		M-180	May 26, 2019
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<b><u>ILOPERIDONE - FANAPT</u></b>						
N 022192 007	8586610	Nov 02, 2027	U-1625		M-180	May 26, 2019
	8652776	Aug 31, 2030	U-1685			
	8999638	Oct 28, 2030	U-1674			
	9072742	Jan 16, 2031	U-1674			
	9074254	Dec 28, 2031	U-1674			
	9074255	Dec 17, 2030	U-1674			
	9074256	Feb 10, 2031	U-1674			
	9138432	Sep 30, 2025	U-1737			
	9157121	Apr 05, 2030	U-1674			
<b><u>IMATINIB MESYLATE - GLEEVEC</u></b>						
N 021335 001	6894051	May 23, 2019	DS DP	U-649		
	6958335	Dec 19, 2021		U-791		
	RE43932	Jul 16, 2018	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>IMATINIB MESYLATE - GLEEVEC</u></b>						
N 021335 001	RE43932*PED	Jan 16, 2019				
<b><u>IMATINIB MESYLATE - GLEEVEC</u></b>						
N 021335 002	6894051 6958335 RE43932 RE43932*PED	May 23, 2019 Dec 19, 2021 Jul 16, 2018 Jan 16, 2019	DS DP U-649 U-791 DS DP DS DP			
<b><u>IMATINIB MESYLATE - GLEEVEC</u></b>						
N 021588 001	6894051 6958335 6958335 6958335*PED 7544799 7544799*PED RE43932 RE43932*PED	May 23, 2019 Dec 19, 2021 Dec 19, 2021 Jun 19, 2022 Jul 16, 2018 Jan 16, 2019 Jul 16, 2018 Jan 16, 2019	DS DP U-649 U-1883 U-791 DS DP Y DS DP DS DP		ODE-40	Jan 25, 2020
<b><u>IMATINIB MESYLATE - GLEEVEC</u></b>						
N 021588 002	6894051 6958335 6958335 6958335*PED 7544799 7544799*PED RE43932 RE43932*PED	May 23, 2019 Dec 19, 2021 Dec 19, 2021 Jun 19, 2022 Jul 16, 2018 Jan 16, 2019 Jul 16, 2018 Jan 16, 2019	DS DP U-649 U-1883 U-791 DS DP Y DS DP DS DP		ODE-40	Jan 25, 2020
<b><u>IMIQUIMOD - ALDARA</u></b>						
N 020723 001	7696159 7696159	Apr 01, 2024 Apr 01, 2024	DS DS	U-1047 U-1048		
<b><u>IMIQUIMOD - ZYCLARA</u></b>						
N 022483 001	8236816 8299109 8598196 8598196	Dec 11, 2029 Dec 11, 2029 Aug 18, 2029 Aug 18, 2029		U-68 U-68 U-1455 U-172		
<b><u>IMIQUIMOD - ZYCLARA</u></b>						
N 022483 002	8222270	Dec 11, 2029		U-68		
<b><u>INDACATEROL MALEATE - ARCAPTA NEOHALER</u></b>						
N 022383 001	6878721 8067437 8479730 8658673 8796307	Feb 25, 2025 Jun 02, 2020 Oct 11, 2028 Jun 02, 2020 Jun 02, 2020	DS DP U-1168 DP DS DP DS DP	U-1168 U-1168 U-1455 U-172		
<b><u>INDINAVIR SULFATE - CRIXIVAN</u></b>						
N 020685 001	6645961 6689761	Mar 04, 2018 Feb 10, 2021		DP U-554		
<b><u>INDINAVIR SULFATE - CRIXIVAN</u></b>						
N 020685 003	6645961 6689761	Mar 04, 2018 Feb 10, 2021		DP U-554		
<b><u>INDINAVIR SULFATE - CRIXIVAN</u></b>						
N 020685 005	6645961 6689761	Mar 04, 2018 Feb 10, 2021		DP U-554		
<b><u>INDINAVIR SULFATE - CRIXIVAN</u></b>						
N 020685 006	6645961 6689761	Mar 04, 2018 Feb 10, 2021		DP U-554		
<b><u>INDIUM IN-111 PENTETREOTIDE KIT - OCTREOSCAN</u></b>						
N 020314 001	6123916	Sep 26, 2017		U-1125		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INDOMETHACIN - TIVORBEX</u></b>						
N 204768 001	8734847	Apr 23, 2030	DP			
	8992982	Apr 23, 2030	DP			
	9089471	Apr 23, 2030	U-55			
<b><u>INDOMETHACIN - TIVORBEX</u></b>						
N 204768 002	8734847	Apr 23, 2030	DP			
	8992982	Apr 23, 2030	DP			
	9089471	Apr 23, 2030	U-55			
<b><u>INGENOL MEBUTATE - PICATO</u></b>						
N 202833 001	6432452	Aug 19, 2018	U-68		M-169	
	6787161	Aug 19, 2018	U-68			
	6844013	Dec 13, 2018	U-1221			
	7410656	Oct 10, 2020	U-1222			
	8278292	Jul 06, 2027	DP			
	8372827	Dec 18, 2026	DP			
	8372828	Dec 18, 2026	DP			
	8377919	Dec 18, 2026	DP			
	8536163	Dec 18, 2026	U-1440			
	8716271	Dec 18, 2026	U-1440			
	8735375	Dec 18, 2026	U-1440			
	9789078	May 15, 2033	U-2138			
	9820959	Dec 18, 2026	DP U-1440			
	9833428	Dec 18, 2026	DP			
	9833429	Dec 18, 2026	DP			
<b><u>INGENOL MEBUTATE - PICATO</u></b>						
N 202833 002	6432452	Aug 19, 2018	U-68			
	6787161	Aug 19, 2018	U-68			
	6844013	Dec 13, 2018	U-1221			
	7410656	Oct 10, 2020	U-1222			
	8278292	Jul 06, 2027	DP			
	8372827	Dec 18, 2026	DP			
	8372828	Dec 18, 2026	DP			
	8377919	Dec 18, 2026	DP			
	8536163	Dec 18, 2026	U-1440			
	8716271	Dec 18, 2026	U-1440			
	8735375	Dec 18, 2026	U-1440			
	9820959	Dec 18, 2026	DP U-1440			
	9833428	Dec 18, 2026	DP			
	9833429	Dec 18, 2026	DP			
<b><u>INSULIN ASPART - FIASP</u></b>						
N 208751 001	8324157	Jun 25, 2030	DP		NP	
<b><u>INSULIN ASPART - FIASP FLEXTOUCH</u></b>						
N 208751 002	6899699	Jan 02, 2022	DP		NP	
	7686786	Aug 03, 2026	DP			
	8324157	Jun 25, 2030	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Jan 20, 2026	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
<b><u>INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT - NOVOLOG MIX 70/30 FLEXPEN</u></b>						
N 021172 004	6004297	Jan 28, 2019	DP			
	9265893	Sep 23, 2032	DP			
	RE41956	Jan 21, 2021	DP			
	RE43834	Jan 28, 2019	DP			
<b><u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXPEN</u></b>						
N 020986 003	6004297	Jan 28, 2019	DP			
	9265893	Sep 23, 2032	DP			
	RE41956	Jan 21, 2021	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXPEN</u></b>						
N 020986 003	RE43834	Jan 28, 2019	DP			
<b><u>INSULIN ASPART RECOMBINANT - NOVOLOG INNOLET</u></b>						
N 020986 004	RE41956	Jan 21, 2021	DP			
<b><u>INSULIN ASPART RECOMBINANT - NOVOLOG FLEXTOUCH</u></b>						
N 020986 005	5866538*PED 6899699 8672898 8684969 8920383 9108002 9132239 9457154 9486588 9687611 9775953 RE46363	Dec 20, 2017 Jan 02, 2022 Jan 02, 2022 Oct 20, 2025 Jul 17, 2026 Jan 20, 2026 Feb 01, 2032 Sep 27, 2027 Jan 02, 2022 Feb 27, 2027 Jul 17, 2026 Aug 03, 2026	DP DP DP DP DP DP DP DP DP DP DP			
<b><u>INSULIN ASPART; INSULIN DEGLUDEC - RYZODEG 70/30</u></b>						
N 203313 001	6899699 7615532 8672898 8684969 8920383 9108002 9132239 9457154 9486588 9687611 9775953 RE46363	Jan 02, 2022 May 25, 2025 Jan 02, 2022 Oct 20, 2025 Jul 17, 2026 Jan 20, 2026 Feb 01, 2032 Sep 27, 2027 Jan 02, 2022 Feb 27, 2027 Jul 17, 2026 Aug 03, 2026	DP DS DP DP DP DP DP DP DP DP DP DP		NCE NPP	Sep 25, 2020 Dec 16, 2019
<b><u>INSULIN DEGLUDEC - TRESIBA</u></b>						
N 203314 001	6899699 7615532 8672898 8684969 8920383 9108002 9132239 9457154 9486588 9687611 9775953 RE46363	Jan 02, 2022 May 25, 2025 Jan 02, 2022 Oct 20, 2025 Jul 17, 2026 Jan 20, 2026 Feb 01, 2032 Sep 27, 2027 Jan 02, 2022 Feb 27, 2027 Jul 17, 2026 Aug 03, 2026	DP DS DP DP DP DP DP DP DP DP DP DP		NCE NPP	Sep 25, 2020 Dec 16, 2019
<b><u>INSULIN DEGLUDEC - TRESIBA</u></b>						
N 203314 002	6899699 7615532 8672898 8684969 8920383 9108002 9132239 9457154 9486588 9687611 9775953 RE46363	Jan 02, 2022 May 25, 2025 Jan 02, 2022 Oct 20, 2025 Jul 17, 2026 Jan 20, 2026 Feb 01, 2032 Sep 27, 2027 Jan 02, 2022 Feb 27, 2027 Jul 17, 2026 Aug 03, 2026	DP DS DP DP DP DP DP DP DP DP DP DP		NCE NPP	Sep 25, 2020 Dec 16, 2019
<b><u>INSULIN DEGLUDEC; LIRAGLUTIDE - XULTOPHY 100/3.6</u></b>						
N 208583 001	6268343 6458924 6899699 7235627 7615532 8672898 8684969	Aug 22, 2022 Aug 22, 2017 Jan 02, 2022 Aug 22, 2017 May 25, 2025 Jan 02, 2022 Oct 20, 2025	DS DP DS DP DP DS DS DP DP DP		NC NCE	Nov 21, 2019 Sep 25, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INSULIN DEGLUDEC; LIRAGLUTIDE - XULTOPHY 100/3.6</u></b>						
N 208583 001	8846618	Jun 27, 2022	DS DP			
	8920383	Jul 17, 2026	DP			
	8937042	May 05, 2029	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b><u>INSULIN DETEMIR RECOMBINANT - LEVEMIR</u></b>						
N 021536 001	5750497	Jun 16, 2019	DS DP U-668			
<b><u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXPEN</u></b>						
N 021536 002	5750497	Jun 16, 2019	DS DP U-668			
	6004297	Jan 28, 2019	DP			
	9265893	Sep 23, 2032	DP			
	RE41956	Jan 21, 2021	DP			
	RE43834	Jan 28, 2019	DP			
<b><u>INSULIN DETEMIR RECOMBINANT - LEVEMIR INNOLET</u></b>						
N 021536 003	5750497	Jun 16, 2019	DS DP U-668			
<b><u>INSULIN DETEMIR RECOMBINANT - LEVEMIR PENFILL</u></b>						
N 021536 004	5750497	Jun 16, 2019	DS DP U-668			
<b><u>INSULIN DETEMIR RECOMBINANT - LEVEMIR FLEXTOUCH</u></b>						
N 021536 005	5750497	Jun 16, 2019	DS DP U-668			
	6899699	Jan 02, 2022	DP			
	7686786	Aug 03, 2026	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b><u>INSULIN GLARGINE - BASAGLAR</u></b>						
N 205692 001					NP	Dec 16, 2018
<b><u>INSULIN GLARGINE RECOMBINANT - LANTUS</u></b>						
N 021081 001	7476652	Jul 23, 2023	DP			
	7713930	Jun 13, 2023	DP			
	7918833	Sep 23, 2027	DP			
<b><u>INSULIN GLARGINE RECOMBINANT - LANTUS SOLOSTAR</u></b>						
N 021081 002	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024	U-1832			
	9233211	Mar 02, 2024	DP			
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9775954	Mar 02, 2024	DP			
	9827379	Mar 02, 2024	DP U-2146			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INSULIN GLARGINE RECOMBINANT - TOUJEO SOLOSTAR</u></b>						
N 206538 001	7918833	Sep 23, 2027	DP		NP	Feb 25, 2018
	7918833*PED	Mar 23, 2028				
	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024	U-1832			
	9233211	Mar 02, 2024	DP			
	9345750	May 18, 2031	DP U-1855			
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9775954	Mar 02, 2024	DP			
	9827379	Mar 02, 2024	DP U-2146			
<b><u>INSULIN GLARGINE; LIXISENATIDE - SOLIQUA 100/33</u></b>						
N 208673 001	7918833	Sep 23, 2027	DP		NC	Nov 21, 2019
	8512297	Sep 15, 2024	DP		NCE	Jul 27, 2021
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024	U-1923			
	9233211	Mar 02, 2024	DP			
	9408979	Mar 02, 2024	DP			
	9526764	Oct 09, 2029	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9707176	Nov 11, 2030	DP			
	9775954	Mar 02, 2024	DP			
	9821032	May 09, 2032	U-2182			
	9827379	Mar 02, 2024	DP U-2146			
	RE45313	Jul 12, 2020	DS DP			
<b><u>INSULIN GLULISINE RECOMBINANT - APIDRA</u></b>						
N 021629 001	6221633	Jun 18, 2018	DS DP U-471			
	6960561	Jan 25, 2023	DP U-471			
	7452860	Mar 22, 2022	DP			
	7696162	Mar 22, 2022	DP U-471			
<b><u>INSULIN GLULISINE RECOMBINANT - APIDRA</u></b>						
N 021629 002	6221633	Jun 18, 2018	DS DP U-471			
	6960561	Jan 25, 2023	DP U-471			
	7452860	Mar 22, 2022	DP			
	7696162	Mar 22, 2022	DP U-471			
<b><u>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u></b>						
N 021629 003	6221633	Jun 18, 2018	DS DP U-471			
	6960561	Jan 25, 2023	DP U-471			
	7452860	Mar 22, 2022	DP			
	7696162	Mar 22, 2022	DP U-471			
	7918833	Sep 23, 2027	DP			
	8512297	Sep 15, 2024	DP			
	8556864	Mar 03, 2024	DP			
	8603044	Mar 02, 2024	DP			
	8679069	Apr 12, 2025	DP			
	8992486	Jun 05, 2024	DP			
	9011391	Mar 26, 2024	U-1832			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INSULIN GLULISINE RECOMBINANT - APIIDRA SOLOSTAR</u></b>						
N 021629 003	9233211	Mar 02, 2024	DP			
	9408979	Mar 02, 2024	DP			
	9526844	Mar 02, 2024	DP			
	9533105	Aug 17, 2024	DP			
	9561331	Aug 28, 2024	DP			
	9604008	Mar 02, 2024	DP			
	9604009	Aug 16, 2024	DP			
	9610409	Mar 02, 2024	DP			
	9623189	Aug 19, 2024	DP			
	9775954	Mar 02, 2024	DP			
	9827379	Mar 02, 2024	DP U-2146			
<b><u>INSULIN HUMAN - HUMULIN R</u></b>						
N 018780 004	7291132	Aug 09, 2024	DP			
<b><u>INSULIN LISPRO - ADMELOG</u></b>						
N 209196 001					NP	Dec 11, 2020
<b><u>INSULIN LISPRO - ADMELOG SOLOSTAR</u></b>						
N 209196 002					NP	Dec 11, 2020
<b><u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 75/25 KWIKPEN</u></b>						
N 021017 002	7291132	Aug 09, 2024	DP			
<b><u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT - HUMALOG MIX 50/50 KWIKPEN</u></b>						
N 021018 002	7291132	Aug 09, 2024	DP			
<b><u>INSULIN LISPRO RECOMBINANT - HUMALOG KWIKPEN</u></b>						
N 020563 003	7291132	Aug 09, 2024	DP			
<b><u>INSULIN LISPRO RECOMBINANT - HUMALOG KWIKPEN</u></b>						
N 205747 001	6034054	Jun 11, 2018	DP U-1707			
	6034054	Jun 11, 2018	DP U-1708			
	6551992	Jun 11, 2018	DP U-1707			
	6551992	Jun 11, 2018	DP U-1708			
	7291132	Aug 09, 2024	DP			
<b><u>INSULIN RECOMBINANT HUMAN - EXUBERA</u></b>						
N 021868 001	6257233	May 14, 2019	U-704			
	6546929	May 14, 2019	U-704			
	6582728	Jun 24, 2020	DP			
	6685967	Sep 11, 2018	DP			
<b><u>INSULIN RECOMBINANT HUMAN - EXUBERA</u></b>						
N 021868 002	6257233	May 14, 2019	U-704			
	6546929	May 14, 2019	U-704			
	6582728	Jun 24, 2020	DP			
	6685967	Sep 11, 2018	DP			
<b><u>INSULIN RECOMBINANT HUMAN - AFREZZA</u></b>						
N 022472 001	6444226	Jun 29, 2020	DP U-1534			
	6652885	Jun 29, 2020	U-1535			
	7305986	Jan 16, 2023	DP			
	7464706	Mar 02, 2023	DP			
	7648960	Jun 29, 2020	U-1535			
	7943178	Jun 29, 2020	DP U-1535			
	7943572	Aug 10, 2026	U-1539			
	8119593	Aug 11, 2029	U-1537			
	8146588	Apr 24, 2023	DP			
	8156936	Jan 16, 2023	DP			
	8215300	Nov 24, 2022	DP			
	8258095	Aug 11, 2029	U-1537			
	8389470	Jun 29, 2020	DP U-1621			
	8424518	Oct 17, 2031	DP			
	8485180	Mar 25, 2030	DP			
	8499757	Feb 19, 2032	DP			
	8551528	Jun 11, 2030	DP			
	8623817	Sep 18, 2029	U-1537			
	8636001	Jul 12, 2032	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INSULIN RECOMBINANT HUMAN - AFREZZA</u></b>						
N 022472 001	8729019	Dec 26, 2028	DP			
	8734845	Jun 11, 2030	DP			
	8778403	Jun 11, 2030	DP U-1538			
	8889099	Jun 29, 2020	DP U-1621			
	8912193	Jun 12, 2029	DP U-1538			
	8950397	Jul 20, 2021	DP			
	9192675	Jun 12, 2029	DP U-1788			
	9283193	Sep 14, 2026	DP			
	9339615	Oct 20, 2029	DP			
	9358352	Feb 15, 2031	DP U-1861			
	9393372	Jul 04, 2029	DP			
	9446133	Jun 12, 2029	DP U-1861			
	9511198	Feb 16, 2030	U-1929			
	9511198	Feb 16, 2030	U-1930			
	9597374	Oct 08, 2031	U-1987			
	9662461	Jun 12, 2029	DP U-2019			
	9717689	Sep 14, 2026	DP			
<b><u>INSULIN RECOMBINANT HUMAN - AFREZZA</u></b>						
N 022472 002	6444226	Jun 29, 2020	DP U-1534			
	6652885	Jun 29, 2020	U-1535			
	7305986	Jan 16, 2023	DP			
	7464706	Mar 02, 2023	DP			
	7648960	Jun 29, 2020	U-1535			
	7943178	Jun 29, 2020	DP U-1535			
	7943572	Aug 10, 2026	U-1539			
	8119593	Aug 11, 2029	U-1537			
	8146588	Apr 24, 2023	DP			
	8156936	Jan 16, 2023	DP			
	8215300	Nov 24, 2022	DP			
	8258095	Aug 11, 2029	U-1537			
	8389470	Jun 29, 2020	DP U-1621			
	8424518	Oct 17, 2031	DP			
	8485180	Mar 25, 2030	DP			
	8499757	Feb 19, 2032	DP			
	8551528	Jun 11, 2030	DP			
	8623817	Sep 18, 2029	U-1537			
	8636001	Jul 12, 2032	DP			
	8729019	Dec 26, 2028	DP			
	8734845	Jun 11, 2030	DP			
	8778403	Jun 11, 2030	DP U-1538			
	8889099	Jun 29, 2020	DP U-1621			
	8912193	Jun 12, 2029	DP U-1538			
	8950397	Jul 20, 2021	DP			
	9192675	Jun 12, 2029	DP U-1788			
	9283193	Sep 14, 2026	DP			
	9339615	Oct 20, 2029	DP			
	9358352	Feb 15, 2031	DP U-1861			
	9393372	Jul 04, 2029	DP			
	9446133	Jun 12, 2029	DP U-1861			
	9511198	Feb 16, 2030	U-1929			
	9511198	Feb 16, 2030	U-1930			
	9597374	Oct 08, 2031	U-1987			
	9662461	Jun 12, 2029	DP U-2019			
	9717689	Sep 14, 2026	DP			
<b><u>INSULIN RECOMBINANT HUMAN - AFREZZA</u></b>						
N 022472 003	6444226	Jun 29, 2020	DP U-1534			
	6652885	Jun 29, 2020	U-1535			
	7305986	Jan 16, 2023	DP			
	7464706	Mar 02, 2023	DP			
	7648960	Jun 29, 2020	U-1535			
	7943178	Jun 29, 2020	DP U-1535			
	7943572	Aug 10, 2026	U-1539			
	8119593	Aug 11, 2029	U-1537			
	8146588	Apr 24, 2023	DP			
	8156936	Jan 16, 2023	DP			
	8215300	Nov 24, 2022	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>INSULIN RECOMBINANT HUMAN - AFREZZA</u></b>						
N 022472 003	8258095	Aug 11, 2029		U-1537		
	8389470	Jun 29, 2020		DP U-1621		
	8424518	Oct 17, 2031		DP		
	8485180	Mar 25, 2030		DP		
	8499757	Feb 19, 2032		DP		
	8551528	Jun 11, 2030		DP		
	8623817	Sep 18, 2029		U-1537		
	8636001	Jul 12, 2032		DP		
	8729019	Dec 26, 2028		DP		
	8734845	Jun 11, 2030		DP		
	8778403	Jun 11, 2030		DP U-1538		
	8889099	Jun 29, 2020		DP U-1621		
	8912193	Jun 12, 2029		DP U-1538		
	8950397	Jul 20, 2021		DP		
	9192675	Jun 12, 2029		DP U-1788		
	9283193	Sep 14, 2026		DP		
	9339615	Oct 20, 2029		DP		
	9358352	Feb 15, 2031		DP U-1861		
	9393372	Jul 04, 2029		DP		
	9446133	Jun 12, 2029		DP U-1861		
	9511198	Feb 16, 2030		U-1929		
	9511198	Feb 16, 2030		U-1930		
	9597374	Oct 08, 2031		U-1987		
	9662461	Jun 12, 2029		DP U-2019		
	9717689	Sep 14, 2026		DP		
<b><u>INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN 70/30</u></b>						
N 019717 001	7291132	Aug 09, 2024		DP		
<b><u>INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN 70/30 PEN</u></b>						
N 019717 002	7291132	Aug 09, 2024		DP		
<b><u>INSULIN SUSP ISOPHANE RECOMBINANT HUMAN - HUMULIN N</u></b>						
N 018781 001	7291132	Aug 09, 2024		DP		
<b><u>IODIXANOL - VISIPAQUE 320</u></b>						
N 020351 002					I-752	Apr 05, 2020
<b><u>IODIXANOL - VISIPAQUE 320</u></b>						
N 020808 002					I-752	Apr 05, 2020
<b><u>IPRATROPIUM BROMIDE - ATROVENT HFA</u></b>						
N 021527 001	6739333	May 26, 2020		DP		
	6983743	May 26, 2020		DP		
	8474447	Jan 17, 2030		DP		
<b><u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u></b>						
N 020571 001	6403569	Apr 28, 2020		U-449		
	6794370	May 01, 2020		U-606		
<b><u>IRINOTECAN HYDROCHLORIDE - CAMPTOSAR</u></b>						
N 020571 002	6403569	Apr 28, 2020		U-449		
	6794370	May 01, 2020		U-606		
<b><u>IRINOTECAN HYDROCHLORIDE - ONIVYDE</u></b>						
N 207793 001	8147867	Aug 29, 2028	DS DP		NP	Oct 22, 2018
	8329213	May 02, 2025	DS DP		ODE-99	Oct 22, 2022
	8703181	May 02, 2025		U-1434		
	8992970	May 02, 2025	DS DP			
	9339497	Jun 12, 2033		U-1848		
	9364473	Jun 12, 2033		U-1856		
	9452162	Jun 12, 2033		U-1899		
	9492442	Jun 12, 2033		U-1848		
	9492442	Jun 12, 2033		U-1899		
	9492442	Jun 12, 2033		U-1917		
	9717724	Jun 12, 2033		U-1848		
	9717724	Jun 12, 2033		U-2091		
	9724303	May 02, 2025	DS DP			
	9730891	May 02, 2025		U-1848		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>IRINOTECAN HYDROCHLORIDE - ONTIVYDE</u></b>						
N 207793 001	9782349	May 02, 2025	DS DP			
<b><u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u></b>						
N 207500 001	6812238	Oct 31, 2020	DS		NCE	Mar 06, 2020
	7459561	Oct 31, 2020	DS		ODE-90	Mar 06, 2022
					GAIN	Mar 06, 2025
					GAIN	Mar 06, 2027
<b><u>ISAVUCONAZONIUM SULFATE - CRESEMBA</u></b>						
N 207501 001	6812238	Oct 31, 2020	DS		NCE	Mar 06, 2020
	7459561	Oct 31, 2020	DS		ODE-90	Mar 06, 2022
					GAIN	Mar 06, 2025
					GAIN	Mar 06, 2027
<b><u>ISOTRETINOIN - ABSORICA</u></b>						
N 021951 001	7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021		U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<b><u>ISOTRETINOIN - ABSORICA</u></b>						
N 021951 002	7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021		U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<b><u>ISOTRETINOIN - ABSORICA</u></b>						
N 021951 003	7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021		U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<b><u>ISOTRETINOIN - ABSORICA</u></b>						
N 021951 004	7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021		U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<b><u>ISOTRETINOIN - ABSORICA</u></b>						
N 021951 005	7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021		U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<b><u>ISOTRETINOIN - ABSORICA</u></b>						
N 021951 006	7435427	Sep 21, 2021	DP			
	8367102	Sep 21, 2021		U-1347		
	8952064	Sep 21, 2021	DP			
	9078925	Sep 21, 2021	DP			
	9089534	Sep 21, 2021	DP			
<b><u>ITRACONAZOLE - SPORANOX</u></b>						
N 020657 001	6407079		Jun 18, 2019			
<b><u>ITRACONAZOLE - SPORANOX</u></b>						
N 020966 001	6407079		Jun 18, 2019			
<b><u>ITRACONAZOLE - ONMEL</u></b>						
N 022484 001	8486456		Oct 03, 2028	DP U-1054		
<b><u>IVABRADINE HYDROCHLORIDE - CORLANOR</u></b>						
N 206143 001	7361649	Apr 17, 2026	DS DP	U-1694		
	7361650	Apr 14, 2026	DS DP	U-1694		
	7867996	Feb 22, 2026	DS DP	U-1694		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>IVABRADINE HYDROCHLORIDE - CORLANOR</u></b>						
N 206143 001	7879842	Feb 22, 2026	DS DP U-1694			
<b><u>IVABRADINE HYDROCHLORIDE - CORLANOR</u></b>						
N 206143 002	7361649	Apr 17, 2026	DS DP U-1694		NCE	Apr 15, 2020
	7361650	Apr 14, 2026	DS DP U-1694			
	7867996	Feb 22, 2026	DS DP U-1694			
	7879842	Feb 22, 2026	DS DP U-1694			
<b><u>IVACAFTOR - KALYDECO</u></b>						
N 203188 001	7495103	May 20, 2027	DS DP		I-705	Dec 30, 2017
	8324242	Aug 05, 2027	U-1311		NPP	Jul 31, 2020
	8324242	Aug 05, 2027	U-1906		ODE-20	Jan 31, 2019
	8354427	Jul 06, 2026	U-1311			
	8354427	Jul 06, 2026	U-1905			
	8410274	Dec 28, 2026	DP			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-1311			
<b><u>IVACAFTOR - KALYDECO</u></b>						
N 207925 001	7495103	May 20, 2027	DS DP		NPP	Jul 31, 2020
	8324242	Aug 05, 2027	U-1311		ODE-20	Jan 31, 2019
	8324242	Aug 05, 2027	U-1906			
	8354427	Jul 06, 2026	U-1311			
	8354427	Jul 06, 2026	U-1905			
	8410274	Dec 28, 2026	DP			
	8754224	Dec 28, 2026	DS DP			
	8883206	Feb 27, 2033	DP			
	9670163	Dec 28, 2026	DP U-1311			
<b><u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u></b>						
N 206038 001	7495103	May 20, 2027	DS DP		NCE	Jul 02, 2020
	7973038	Nov 08, 2026	U-1973		ODE-123	Sep 28, 2023
	8324242	Aug 05, 2027	U-1311		ODE-93	Jul 02, 2022
	8324242	Aug 05, 2027	U-1911			
	8410274	Dec 28, 2026	DP			
	8507534	Sep 20, 2030	DS DP			
	8653103	Dec 04, 2028	DP			
	8716338	Sep 20, 2030	DP U-1718			
	8716338	Sep 20, 2030	DP U-1910			
	8741933	Nov 08, 2026	U-1717			
	8741933	Nov 08, 2026	U-1909			
	8754224	Dec 28, 2026	DS DP			
	8846718	Dec 04, 2028	U-1717			
	8846718	Dec 04, 2028	U-1908			
	8993600	Dec 11, 2030	DP			
	9150552	Dec 04, 2028	U-1908			
	9192606	Sep 29, 2029	DP U-1912			
	9216969	Nov 08, 2026	DS DP			
	9670163	Dec 28, 2026	DP U-1911			
<b><u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u></b>						
N 206038 002	7495103	May 20, 2027	DS DP		NCE	Jul 02, 2020
	7973038	Nov 08, 2026	U-1973		NPP	Sep 28, 2019
	8324242	Aug 05, 2027	U-1911		ODE-123	Sep 28, 2023
	8410274	Dec 28, 2026	DP		ODE-93	Jul 02, 2022
	8507534	Sep 20, 2030	DS DP			
	8653103	Dec 04, 2028	DP			
	8716338	Sep 20, 2030	DP U-1910			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>IVACAFTOR; LUMACAFTOR - ORKAMBI</u></b>						
N 206038 002	8741933	Nov 08, 2026	U-1909			
	8754224	Dec 28, 2026	DS DP			
	8846718	Dec 04, 2028		U-1908		
	8993600	Dec 11, 2030	DP			
	9150552	Dec 04, 2028		U-1908		
	9192606	Sep 29, 2029	DP	U-1912		
	9216969	Nov 08, 2026	DP			
	9670163	Dec 28, 2026	DP	U-1911		
<b><u>IVERMECTIN - SKLINE</u></b>						
N 202736 001	6103248	May 22, 2018	DP			
	8791153	Oct 12, 2027	DP			
	8927595	Oct 12, 2027		U-1782		
<b><u>IVERMECTIN - SOOLANTRA</u></b>						
N 206255 001	5952372	Sep 18, 2018	U-1631		NP	
	6133310	Apr 26, 2019	U-1631			
	7550440	Apr 22, 2024	DP	U-1631		
	8080530	Apr 22, 2024	DP	U-1631		
	8093219	Apr 22, 2024	DP	U-1631		
	8415311	Apr 22, 2024	DP	U-1631		
	8470788	Apr 22, 2024	DP	U-1631		
	8815816	Apr 22, 2024	DP	U-1631		
	9089587	Mar 13, 2034		U-1631		
	9233117	Mar 13, 2034		U-1631		
	9233118	Mar 13, 2034		U-1631		
	9782425	Mar 13, 2034		U-1631		
<b><u>IXABEPILONE - IXEMPRA KIT</u></b>						
N 022065 001	6670384	Jan 23, 2022	DP	U-959		
	6670384	Jan 23, 2022	DP	U-960		
	7022330	Jan 23, 2022	DP	U-958		
	7125899	May 26, 2018	DS	DP U-957		
	7312237	Aug 21, 2024		U-965		
	RE41393	Feb 08, 2022		U-961		
	RE41911	Sep 28, 2020	DS	DP U-961		
<b><u>IXABEPILONE - IXEMPRA KIT</u></b>						
N 022065 002	6670384	Jan 23, 2022	DP	U-959		
	6670384	Jan 23, 2022	DP	U-960		
	7022330	Jan 23, 2022	DP	U-958		
	7125899	May 26, 2018	DS	DP U-957		
	7312237	Aug 21, 2024		U-965		
	RE41393	Feb 08, 2022		U-961		
	RE41911	Sep 28, 2020	DS	DP U-961		
<b><u>IXAZOMIB CITRATE - NINLARO</u></b>						
N 208462 001	7442830	Aug 06, 2027	DS	DP U-1780		
	7687662	Aug 06, 2027	DS	DP		
	8003819	Aug 06, 2027	DS	DP U-1780		
	8530694	Aug 06, 2027	DS	DP U-1780		
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS	DP		
	8871745	Aug 06, 2027		U-1779		
	9175017	Jun 16, 2029		U-1778		
	9233115	Aug 12, 2024		U-1778		
<b><u>IXAZOMIB CITRATE - NINLARO</u></b>						
N 208462 002	7442830	Aug 06, 2027	DS	DP U-1780		
	7687662	Aug 06, 2027	DS	DP		
	8003819	Aug 06, 2027	DS	DP U-1780		
	8530694	Aug 06, 2027	DS	DP U-1780		
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS	DP		
	8871745	Aug 06, 2027		U-1779		
	9175017	Jun 16, 2029		U-1778		
	9233115	Aug 12, 2024		U-1778		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>IXAZOMIB CITRATE - NINLARO</u></b>						
N 208462 003	7442830	Aug 06, 2027	DS DP U-1780		NCE	Nov 20, 2020
	7687662	Aug 06, 2027	DS DP		ODE-103	Nov 20, 2022
	8003819	Aug 06, 2027	DS DP U-1780			
	8530694	Aug 06, 2027	DS DP U-1780			
	8546608	Aug 12, 2024	DS			
	8859504	Jun 16, 2029	DS DP			
	8871745	Aug 06, 2027	U-1779			
	9175017	Jun 16, 2029	U-1778			
	9233115	Aug 12, 2024	U-1778			
<b><u>KETOCONAZOLE - EXTINA</u></b>						
N 021738 001	7553835	Oct 19, 2018	DP U-245			
	8026238	Oct 19, 2018	DP U-1213			
<b><u>KETOCONAZOLE - XOLEGEL</u></b>						
N 021946 001	7179475	Dec 04, 2018	DP U-792			
	8232276	Nov 24, 2020	DP			
	8735393	Dec 04, 2018	DP			
<b><u>KETOROLAC TROMETHAMINE - ACULAR LS</u></b>						
N 021528 001	8008338	May 24, 2027	DS DP U-1181			
	8207215	May 28, 2024	U-1251			
	8377982	May 28, 2024	U-1363			
	8377982*PED	Nov 28, 2024				
	8541463	May 28, 2024	U-1441			
	8541463*PED	Nov 28, 2024				
	8648107	May 28, 2024	DP			
	8906950	May 28, 2024	U-1626			
	8946281	May 28, 2024	U-1662			
	9216167	May 28, 2024	U-1800			
<b><u>KETOROLAC TROMETHAMINE - SPRIX</u></b>						
N 022382 001	6333044	Dec 25, 2018	DP U-1057			
<b><u>KETOROLAC TROMETHAMINE - ACUVAIL</u></b>						
N 022427 001	7842714	Aug 15, 2029	DS DP			
	8512717	Mar 07, 2028	DP			
	8992952	Aug 05, 2024	DP			
	9192571	Mar 07, 2028	DP			
<b><u>KETOROLAC TROMETHAMINE; PHENYLEPHRINE HYDROCHLORIDE - OMIDRIA</u></b>						
N 205388 001	8173707	Jul 30, 2023	U-1518		NPP	Dec 08, 2020
	8173707*PED	Jan 30, 2024			PED	Jun 08, 2021
	8586633	Jul 30, 2023	DP			
	8586633*PED	Jan 30, 2024				
	9066856	Oct 23, 2033	DP			
	9066856*PED	Apr 23, 2034				
	9278101	Jul 30, 2023	U-1518			
	9278101*PED	Jan 30, 2024				
	9399040	Jul 30, 2023	DP			
	9399040*PED	Jan 30, 2024				
	9486406	Oct 23, 2033	DP			
	9486406*PED	Apr 23, 2034				
	9855246	Oct 23, 2033	DP			
<b><u>L-GLUTAMINE - ENDARI</u></b>						
N 208587 001				I-748	Jul 07, 2020	
				ODE-150	Jul 07, 2024	
<b><u>LACOSAMIDE - VIMPAT</u></b>						
N 022253 001	RE38551	Mar 17, 2022	DS U-1567		D-143	Aug 29, 2017
	RE38551	Mar 17, 2022	DS U-2140		D-144	Aug 29, 2017
					I-696	Aug 29, 2017
					NPP	Nov 03, 2020
<b><u>LACOSAMIDE - VIMPAT</u></b>						
N 022253 002	RE38551	Mar 17, 2022	DS U-1567		D-143	Aug 29, 2017
	RE38551	Mar 17, 2022	DS U-2140		D-144	Aug 29, 2017
					I-696	Aug 29, 2017

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LACOSAMIDE - VIMPAT</b>						
N 022253 002					NPP	Nov 03, 2020
<b>LACOSAMIDE - VIMPAT</b>						
N 022253 003	RE38551	Mar 17, 2022	DS U-1567	D-143	Aug 29, 2017	
	RE38551	Mar 17, 2022	DS U-2140	D-144	Aug 29, 2017	
				I-696	Aug 29, 2017	
				NPP	Nov 03, 2020	
<b>LACOSAMIDE - VIMPAT</b>						
N 022253 004	RE38551	Mar 17, 2022	DS U-1567	D-143	Aug 29, 2017	
	RE38551	Mar 17, 2022	DS U-2140	D-144	Aug 29, 2017	
				I-696	Aug 29, 2017	
				NPP	Nov 03, 2020	
<b>LACOSAMIDE - VIMPAT</b>						
N 022254 001	RE38551	Mar 17, 2022	DS DP U-1565	D-143	Aug 29, 2017	
	RE38551	Mar 17, 2022	DS DP U-1568	D-144	Aug 29, 2017	
				I-696	Aug 29, 2017	
				M-217	Nov 03, 2020	
<b>LACOSAMIDE - VIMPAT</b>						
N 022255 001	RE38551	Mar 17, 2022	DS U-1567	D-143	Aug 29, 2017	
	RE38551	Mar 17, 2022	DS U-2140	D-144	Aug 29, 2017	
				I-696	Aug 29, 2017	
				NPP	Nov 03, 2020	
<b>LAMIVUDINE - EPIVIR</b>						
N 020596 001	6004968	Mar 20, 2018	DP U-248			
	6004968*PED	Sep 20, 2018				
<b>LAMIVUDINE - EPIVIR-HBV</b>						
N 021004 001	6004968	Mar 20, 2018				
<b>LAMIVUDINE; RALTEGRAVIR POTASSIUM - DUTREBIS</b>						
N 206510 001	7169780	Oct 03, 2023	DS DP			
	7169780*PED	Apr 03, 2024				
	7217713	Oct 21, 2022	U-1663			
	7217713*PED	Apr 21, 2023				
	7435734	Oct 21, 2022	U-1663			
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP U-1663			
	7754731*PED	Sep 11, 2029				
	7820660	Apr 25, 2023	DS			
<b>LAMOTRIGINE - LAMICTAL</b>						
N 020241 001				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL</b>						
N 020241 002				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL</b>						
N 020241 003				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL</b>						
N 020241 004				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL</b>						
N 020241 005				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL</b>						
N 020241 006				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL CD</b>						
N 020764 001				M-159	May 18, 2018	
<b>LAMOTRIGINE - LAMICTAL CD</b>						
N 020764 002				M-159	May 18, 2018	

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LAMOTRIGINE - LAMICTAL CD</u>	N 020764 003				M-159	May 18, 2018
<u>LAMOTRIGINE - LAMICTAL CD</u>	N 020764 004				M-159	May 18, 2018
<u>LAMOTRIGINE - LAMICTAL XR</u>	N 022115 001 8637512 9144547	Jun 14, 2028 Sep 22, 2023	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>	N 022115 002 8637512 9144547	Jun 14, 2028 Sep 22, 2023	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>	N 022115 003 8637512 9144547	Jun 14, 2028 Sep 22, 2023	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>	N 022115 004 8637512 9144547	Jun 14, 2028 Sep 22, 2023	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>	N 022115 005 8637512 9144547	Jun 14, 2028 Sep 22, 2023	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>	N 022115 006 8637512 9144547	Jun 14, 2028 Sep 22, 2023	DP			
<u>LAMOTRIGINE - LAMICTAL ODT</u>	N 022251 001 7919115 8840925 9339504	Jan 04, 2029 Jul 02, 2028 Jul 02, 2028	DS DP DP U-1596 DP U-1596		M-159	May 18, 2018
<u>LAMOTRIGINE - LAMICTAL ODT</u>	N 022251 002 7919115 8840925 9339504	Jan 04, 2029 Jul 02, 2028 Jul 02, 2028	DS DP DP U-1596 DP U-1596		M-159	May 18, 2018
<u>LAMOTRIGINE - LAMICTAL ODT</u>	N 022251 003 7919115 8840925 9339504	Jan 04, 2029 Jul 02, 2028 Jul 02, 2028	DS DP DP U-1596 DP U-1596		M-159	May 18, 2018
<u>LAMOTRIGINE - LAMICTAL ODT</u>	N 022251 004 7919115 8840925 9339504	Jan 04, 2029 Jul 02, 2028 Jul 02, 2028	DS DP DP U-1596 DP U-1596		M-159	May 18, 2018
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>	N 022074 001 5595760	Mar 08, 2020	DP U-831		I-701 I-754 ODE-156 ODE-82	Dec 19, 2017 Sep 15, 2020 Sep 15, 2024 Dec 16, 2021
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>	N 022074 002 5595760	Mar 08, 2020	DP U-831		I-701 I-754 ODE-156 ODE-82	Dec 19, 2017 Sep 15, 2020 Sep 15, 2024 Dec 16, 2021
<u>LANREOTIDE ACETATE - SOMATULINE DEPOT</u>	N 022074 003 5595760	Mar 08, 2020	DP U-831		I-701 I-754 ODE-156 ODE-82	Dec 19, 2017 Sep 15, 2020 Sep 15, 2024 Dec 16, 2021

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LANSOPRAZOLE - PREVACID</u></b>						
N 021428 001	6328994	May 17, 2019				
	7399485	May 26, 2018	DP			
	7431942	May 17, 2019	DP			
	7875292	May 17, 2019	DP			
<b><u>LANSOPRAZOLE - PREVACID</u></b>						
N 021428 002	6328994	May 17, 2019				
	7399485	May 26, 2018	DP			
	7431942	May 17, 2019	DP			
	7875292	May 17, 2019	DP			
<b><u>LANSOPRAZOLE - PREVACID IV</u></b>						
N 021566 001	7396841	Aug 17, 2021	DP U-947			
<b><u>LANTHANUM CARBONATE - FOSRENOL</u></b>						
N 021468 001	5968976	Oct 26, 2018	DP U-613			
	7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<b><u>LANTHANUM CARBONATE - FOSRENOL</u></b>						
N 021468 002	5968976	Oct 26, 2018	DP U-613			
	7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<b><u>LANTHANUM CARBONATE - FOSRENOI</u></b>						
N 021468 003	5968976	Oct 26, 2018	DP U-613			
	7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<b><u>LANTHANUM CARBONATE - FOSRENOI</u></b>						
N 021468 004	5968976	Oct 26, 2018	DP U-613			
	7381428	Aug 26, 2024	U-890			
	7465465	Aug 26, 2024	DP			
<b><u>LANTHANUM CARBONATE - FOSRENOI</u></b>						
N 204734 001	5968976	Oct 26, 2018	DP U-1592			
	7465465	Aug 26, 2024	DP			
	8980327	Dec 01, 2030	DP			
	9023397	Dec 01, 2030	DP			
<b><u>LANTHANUM CARBONATE - FOSRENOI</u></b>						
N 204734 002	5968976	Oct 26, 2018	DP U-1592			
	7465465	Aug 26, 2024	DP			
	8980327	Dec 01, 2030	DP			
	9023397	Dec 01, 2030	DP			
<b><u>LAPATINIB DITOSYLATE - TYKERB</u></b>						
N 022059 001	6391874	Jul 11, 2017	DS DP U-1429			
	6391874	Jul 11, 2017	DS DP U-800			
	6713485	Sep 29, 2020	DS DP U-1429			
	6713485	Sep 29, 2020	DS DP U-800			
	6727256	Jan 08, 2019	DS DP U-1429			
	6727256	Jan 08, 2019	DS DP U-800			
	6828320	Jul 11, 2017	U-1429			
	6828320	Jul 11, 2017	U-800			
	7157466	Nov 19, 2021	DS DP			
	8513262	Jan 08, 2019	DS DP			
	8821927	Sep 18, 2029	DS DP			
<b><u>LATANOPROSTENE BUNOD - VYZULTA</u></b>						
N 207795 001	6211233	Jun 17, 2018	DS DP			
	7273946	Oct 03, 2025	DS DP U-2144			
	7629345	Jan 05, 2025	DP U-2144			
	7910767	Jan 05, 2025	DS DP U-2144			
	8058467	Jan 05, 2025	DS U-2144			
<b><u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u></b>						
N 205834 001	7964580	Mar 26, 2029	DS DP U-1470	D-153	Nov 12, 2018	
	8088368	May 12, 2030	DS DP	D-158	Feb 12, 2019	

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LEDIPASVIR; SOFOSBUVIR - HARVONI</u></b>						
N 205834 001	8273341	May 12, 2030	U-1470		D-159	Feb 12, 2019
	8334270	Mar 21, 2028	DS DP U-1470		D-160	Feb 12, 2019
	8580765	Mar 21, 2028	DS DP U-1470		I-718	Nov 12, 2018
	8618076	Dec 11, 2030	DS DP U-1470		I-719	Nov 12, 2018
	8633309	Mar 26, 2029	DS DP U-1470		I-720	Nov 12, 2018
	8735372	Mar 21, 2028	U-1470		NCE	Oct 10, 2019
	8822430	May 12, 2030	DS DP U-1470		NPP	Nov 12, 2018
	8841278	May 12, 2030	DP U-1470		NPP	Apr 07, 2020
	8889159	Mar 26, 2029	DP U-1470		ODE-136	Apr 07, 2024
	9085573	Mar 21, 2028	DS DP U-1470			
	9284342	Sep 13, 2030	DS DP U-1470			
	9393256	Sep 14, 2032	U-1470			
	9511056	May 12, 2030	DP U-1470			
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 001	5635517	Oct 04, 2019	DS		I-706	Feb 17, 2018
	6045501	Aug 28, 2018	U-1210		ODE-131	Feb 22, 2024
	6315720	Oct 23, 2020	U-1210		ODE-49	Jun 05, 2020
	6561976	Aug 28, 2018	U-1210		ODE-88	Feb 17, 2022
	6561977	Oct 23, 2020	U-1210			
	6755784	Oct 23, 2020	U-1210			
	6908432	Aug 28, 2018	U-1210			
	7189740	Apr 11, 2023	U-1215			
	7465800	Apr 27, 2027	DS DP			
	7468363	Oct 07, 2023	U-1414			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023	U-1216			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8404717	Apr 11, 2023	U-1215			
	8530498	May 15, 2023	U-1216			
	8589188	Aug 28, 2018	U-1210			
	8626531	Oct 23, 2020	U-1210			
	8648095	May 15, 2023	U-1216			
	8741929	Mar 08, 2028	U-1414			
	9056120	Apr 11, 2023	U-1215			
	9101621	May 15, 2023	U-1216			
	9101622	May 15, 2023	U-1216			
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 002	5635517	Oct 04, 2019	DS		I-706	Feb 17, 2018
	6045501	Aug 28, 2018	U-1210		ODE-131	Feb 22, 2024
	6315720	Oct 23, 2020	U-1210		ODE-49	Jun 05, 2020
	6561976	Aug 28, 2018	U-1210		ODE-88	Feb 17, 2022
	6561977	Oct 23, 2020	U-1210			
	6755784	Oct 23, 2020	U-1210			
	6908432	Aug 28, 2018	U-1210			
	7189740	Apr 11, 2023	U-1215			
	7465800	Apr 27, 2027	DS DP			
	7468363	Oct 07, 2023	U-1414			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023	U-1216			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8404717	Apr 11, 2023	U-1215			
	8530498	May 15, 2023	U-1216			
	8589188	Aug 28, 2018	U-1210			
	8626531	Oct 23, 2020	U-1210			
	8648095	May 15, 2023	U-1216			
	8741929	Mar 08, 2028	U-1414			
	9056120	Apr 11, 2023	U-1215			
	9101621	May 15, 2023	U-1216			
	9101622	May 15, 2023	U-1216			
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 003	5635517	Oct 04, 2019	DS		I-706	Feb 17, 2018
	6045501	Aug 28, 2018	U-1210		ODE-131	Feb 22, 2024
	6315720	Oct 23, 2020	U-1210		ODE-49	Jun 05, 2020
	6561976	Aug 28, 2018	U-1210		ODE-88	Feb 17, 2022

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 003	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7468363	Oct 07, 2023		U-1414		
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		
	8315886	Oct 23, 2020		U-1249		
	8404717	Apr 11, 2023		U-1215		
	8530498	May 15, 2023		U-1216		
	8589188	Aug 28, 2018		U-1210		
	8626531	Oct 23, 2020		U-1210		
	8648095	May 15, 2023		U-1216		
	8741929	Mar 08, 2028		U-1414		
	9056120	Apr 11, 2023		U-1215		
	9101621	May 15, 2023		U-1216		
	9101622	May 15, 2023		U-1216		
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 004	5635517	Oct 04, 2019	DS		I-706	Feb 17, 2018
	6045501	Aug 28, 2018		U-1210	ODE-131	Feb 22, 2024
	6315720	Oct 23, 2020		U-1210	ODE-49	Jun 05, 2020
	6561976	Aug 28, 2018		U-1210	ODE-88	Feb 17, 2022
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7468363	Oct 07, 2023		U-1414		
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		
	8315886	Oct 23, 2020		U-1249		
	8404717	Apr 11, 2023		U-1215		
	8530498	May 15, 2023		U-1216		
	8589188	Aug 28, 2018		U-1210		
	8626531	Oct 23, 2020		U-1210		
	8648095	May 15, 2023		U-1216		
	8741929	Mar 08, 2028		U-1414		
	9056120	Apr 11, 2023		U-1215		
	9101621	May 15, 2023		U-1216		
	9101622	May 15, 2023		U-1216		
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 005	5635517	Oct 04, 2019	DS		I-706	Feb 17, 2018
	6045501	Aug 28, 2018		U-1210	ODE-131	Feb 22, 2024
	6315720	Oct 23, 2020		U-1210	ODE-49	Jun 05, 2020
	6561976	Aug 28, 2018		U-1210	ODE-88	Feb 17, 2022
	6561977	Oct 23, 2020		U-1210		
	6755784	Oct 23, 2020		U-1210		
	6908432	Aug 28, 2018		U-1210		
	7189740	Apr 11, 2023		U-1215		
	7465800	Apr 27, 2027	DS DP			
	7468363	Oct 07, 2023		U-1414		
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023		U-1216		
	8204763	Aug 28, 2018		U-1249		
	8315886	Oct 23, 2020		U-1249		
	8404717	Apr 11, 2023		U-1215		
	8530498	May 15, 2023		U-1216		
	8589188	Aug 28, 2018		U-1210		
	8626531	Oct 23, 2020		U-1210		
	8648095	May 15, 2023		U-1216		
	8741929	Mar 08, 2028		U-1414		
	9056120	Apr 11, 2023		U-1215		
	9101621	May 15, 2023		U-1216		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 005	9101622	May 15, 2023		U-1216		
<b><u>LENALIDOMIDE - REVLIMID</u></b>						
N 021880 006	5635517	Oct 04, 2019	DS		I-706	Feb 17, 2018
	6045501	Aug 28, 2018	U-1210		ODE-131	Feb 22, 2024
	6315720	Oct 23, 2020	U-1210		ODE-49	Jun 05, 2020
	6561976	Aug 28, 2018	U-1210		ODE-88	Feb 17, 2022
	6561977	Oct 23, 2020	U-1210			
	6755784	Oct 23, 2020	U-1210			
	6908432	Aug 28, 2018	U-1210			
	7189740	Apr 11, 2023	U-1215			
	7465800	Apr 27, 2027	DS DP			
	7468363	Oct 07, 2023	U-1414			
	7855217	Nov 24, 2024	DS DP			
	7968569	Oct 07, 2023	U-1216			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8404717	Apr 11, 2023	U-1215			
	8530498	May 15, 2023	U-1216			
	8589188	Aug 28, 2018	U-1210			
	8626531	Oct 23, 2020	U-1210			
	8648095	May 15, 2023	U-1216			
	8741929	Mar 08, 2028	U-1414			
	9056120	Apr 11, 2023	U-1215			
	9101621	May 15, 2023	U-1216			
	9101622	May 15, 2023	U-1216			
<b><u>LENVATINIB MESYLATE - LENVIMA</u></b>						
N 206947 001	7253286	Oct 19, 2021	DS DP		I-734	May 13, 2019
	7612208	Sep 19, 2026	DS DP		NCE	Feb 13, 2020
	9006256	Jul 27, 2027	U-1695		ODE-87	Feb 13, 2022
<b><u>LENVATINIB MESYLATE - LENVIMA</u></b>						
N 206947 002	7253286	Oct 19, 2021	DS DP		I-734	May 13, 2019
	7612208	Sep 19, 2026	DS DP		NCE	Feb 13, 2020
	9006256	Jul 27, 2027	U-1695		ODE-87	Feb 13, 2022
<b><u>LESINURAD - ZURAMPIC</u></b>						
N 207988 001	8003681	Aug 25, 2025	DS		NCE	Dec 22, 2020
	8084483	Aug 17, 2029	U-1801			
	8283369	Nov 26, 2028	U-1802			
	8283369	Nov 26, 2028	U-1804			
	8357713	Nov 26, 2028	DP U-1801			
	8357713	Nov 26, 2028	DP U-1802			
	8357713	Nov 26, 2028	DP U-1803			
	8546436	Feb 29, 2032	DS DP			
	8546437	Apr 29, 2029	U-1803			
	9216179	Aug 01, 2031	U-1806			
<b><u>LETTERMOVIR - PREVYMIS</u></b>						
N 209939 001	7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP			
<b><u>LETTERMOVIR - PREVYMIS</u></b>						
N 209939 002	7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP			
<b><u>LETTERMOVIR - PREVYMIS</u></b>						
N 209940 001	7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP			
<b><u>LETTERMOVIR - PREVYMIS</u></b>						
N 209940 002	7196086	May 22, 2024	DS DP		NCE	Nov 08, 2022
	8513255	May 22, 2024	DS DP			
<b><u>LETROZOLE; RIBOCICLIB SUCCINATE - KISQALI FEMARA CO-PACK (COPACKAGED)</u></b>						
N 209935 001	8324225	Jun 17, 2028	DS DP		NCE	Mar 13, 2022
	8415355	Feb 19, 2031	DS DP			
	8685980	May 25, 2030	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LETROZOLE; RIBOCICLIB SUCCINATE - KISQALI FEMARA CO-PACK (COPACKAGED)</u></b>						
N 209935 001	8962630	Dec 09, 2029		U-1981		
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029		U-1981		
<b><u>LEUPROLIDE ACETATE - LUPRON DEPOT</u></b>						
N 020517 003	7429559	Jan 13, 2019		DP		
	8815801	Jun 28, 2022		DP		
	8921326	Feb 05, 2031		DP U-1666		
<b><u>LEUPROLIDE ACETATE - VIADUR</u></b>						
N 021088 001	6113938	Jul 24, 2018				
	6375978	Dec 17, 2018				
<b><u>LEUPROLIDE ACETATE - ELIGARD</u></b>						
N 021343 001	6565874	Oct 28, 2018		DP U-801		
	6626870	Mar 27, 2020		DP		
	6773714	Oct 28, 2018		DP U-801		
	8470359	Oct 15, 2023	DS	DP U-621		
	8486455	Oct 28, 2018	DS	DP		
	8840916	Nov 13, 2020		DP		
	9283282	Oct 28, 2018	DS	DP		
	9539333	Nov 13, 2020	DS	DP U-621		
<b><u>LEUPROLIDE ACETATE - ELIGARD</u></b>						
N 021488 001	6565874	Oct 28, 2018		DP U-801		
	6626870	Mar 27, 2020		DP		
	6773714	Oct 28, 2018		DP U-801		
	8470359	Oct 15, 2023	DS	DP U-621		
	8486455	Oct 28, 2018	DS	DP		
	8840916	Nov 13, 2020		DP		
	9283282	Oct 28, 2018	DS	DP		
	9539333	Nov 13, 2020	DS	DP U-621		
<b><u>LEUPROLIDE ACETATE - ELIGARD</u></b>						
N 021731 001	6565874	Oct 28, 2018		DP U-621		
	6626870	Mar 27, 2020		DP		
	6773714	Oct 28, 2018		U-621		
	8470359	Oct 15, 2023	DS	DP U-621		
	8486455	Oct 28, 2018	DS	DP		
	8840916	Nov 13, 2020		DP		
	9283282	Oct 28, 2018	DS	DP		
	9539333	Nov 13, 2020	DS	DP U-621		
<b><u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u></b>						
N 020837 001	6451289	Mar 21, 2021			M-151	Jan 22, 2018
<b><u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u></b>						
N 020837 002	6451289	Mar 21, 2021			M-151	Jan 22, 2018
<b><u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u></b>						
N 020837 003	6451289	Mar 21, 2021			M-151	Jan 22, 2018
<b><u>LEVALBUTEROL HYDROCHLORIDE - XOPENEX</u></b>						
N 020837 004	6451289	Mar 21, 2021		DP	M-151	Jan 22, 2018
<b><u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u></b>						
N 021730 001	7256310	Oct 08, 2024	DS	DP U-636	M-156	Mar 12, 2018
	8765153	Dec 08, 2023	DP			
<b><u>LEVETIRACETAM - ROWEEPRA</u></b>						
A 090906 003					PC	Sep 13, 2017

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LEVETIRACETAM - KEPTRA</u></b>						
N 021035 001	8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<b><u>LEVETIRACETAM - KEPTRA</u></b>						
N 021035 002	8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<b><u>LEVETIRACETAM - KEPTRA</u></b>						
N 021035 003	8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<b><u>LEVETIRACETAM - KEPTRA</u></b>						
N 021035 004	8802142	Jun 07, 2031	DP			
	8802142*PED	Dec 07, 2031				
<b><u>LEVETIRACETAM - KEPTRA XR</u></b>						
N 022285 001	7858122	Sep 17, 2028	DP			
<b><u>LEVETIRACETAM - KEPTRA XR</u></b>						
N 022285 002	7858122	Sep 17, 2028	DP			
<b><u>LEVETIRACETAM - SPRITAM</u></b>						
N 207958 001	6471992	Feb 20, 2018	DP			
	9339489	Mar 14, 2034	DP U-1850			
	9463160	Feb 20, 2018	DP			
	9669009	Mar 14, 2034	U-1850			
	9669009	Mar 14, 2034	U-2021			
	9669009	Mar 14, 2034	U-2022			
<b><u>LEVETIRACETAM - SPRITAM</u></b>						
N 207958 002	6471992	Feb 20, 2018	DP			
	9339489	Mar 14, 2034	DP U-1850			
	9463160	Feb 20, 2018	DP			
	9669009	Mar 14, 2034	U-1850			
	9669009	Mar 14, 2034	U-2021			
	9669009	Mar 14, 2034	U-2022			
<b><u>LEVETIRACETAM - SPRITAM</u></b>						
N 207958 003	6471992	Feb 20, 2018	DP			
	9339489	Mar 14, 2034	DP U-1850			
	9463160	Feb 20, 2018	DP			
	9669009	Mar 14, 2034	U-1850			
	9669009	Mar 14, 2034	U-2021			
	9669009	Mar 14, 2034	U-2022			
<b><u>LEVETIRACETAM - SPRITAM</u></b>						
N 207958 004	6471992	Feb 20, 2018	DP			
	9339489	Mar 14, 2034	DP U-1850			
	9463160	Feb 20, 2018	DP			
	9669009	Mar 14, 2034	U-1850			
	9669009	Mar 14, 2034	U-2021			
	9669009	Mar 14, 2034	U-2022			
<b><u>LEVOCARNITINE - CARNITOR</u></b>						
N 020182 001	6335369	Jan 18, 2021	U-433			
	6429230	Jan 18, 2021	U-433			
	6696493	Jan 18, 2021	U-433			
<b><u>LEVOSETIRIZINE DIHYDROCHLORIDE - Xyzal Allergy 24HR</u></b>						
N 209090 001	8633194	Oct 16, 2027	DP			
<b><u>LEVOFLOXACIN - LEVAQUIN</u></b>						
N 021721 001	6806256	Feb 26, 2022	DP			
<b><u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u></b>						
N 020140 001	6500829	Mar 07, 2022	DS DP		ODE-10	Apr 29, 2018

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u></b>						
N 020140 002	6500829	Mar 07, 2022	DS DP		ODE-10	Apr 29, 2018
<b><u>LEVOLEUCOVORIN CALCIUM - FUSILEV</u></b>						
N 020140 003	6500829	Mar 07, 2022	DS DP		ODE-10	Apr 29, 2018
<b><u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u></b>						
N 204168 001	8481598	Mar 02, 2031		U-839		
	8865937	May 23, 2032	DS DP		NCE*	Jul 25, 2018
	RE43879	Jun 03, 2023		U-839		
<b><u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u></b>						
N 204168 002	8481598	Mar 02, 2031		U-839		
	8865937	May 23, 2032	DS DP		NCE*	Jul 25, 2018
	RE43879	Jun 03, 2023		U-839		
<b><u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u></b>						
N 204168 003	8481598	Mar 02, 2031		U-839		
	8865937	May 23, 2032	DS DP		NCE*	Jul 25, 2018
	RE43879	Jun 03, 2023		U-839		
<b><u>LEVOMILNACIPRAN HYDROCHLORIDE - FETZIMA</u></b>						
N 204168 004	8481598	Mar 02, 2031		U-839		
	8865937	May 23, 2032	DS DP		NCE*	Jul 25, 2018
	RE43879	Jun 03, 2023		U-839		
<b><u>LEVONORGESTREL - MIRENA</u></b>						
N 021225 001	9615965	Sep 16, 2029	DP	U-2003		
	9668912	Apr 01, 2031	DP			
<b><u>LEVONORGESTREL - SKYLA</u></b>						
N 203159 001	7252839	Nov 13, 2023	DP			
	9615965	Sep 16, 2029	DP	U-2003		
	9668912	Apr 01, 2031	DP			
<b><u>LEVONORGESTREL - LILETTA</u></b>						
N 206229 001					NP	Feb 26, 2018
<b><u>LEVONORGESTREL - KYLEENA</u></b>						
N 208224 001	7252839	Nov 13, 2023	DP		NP	
	9615965	Sep 16, 2029	DP	U-2003		
	9668912	Apr 01, 2031	DP			
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 001	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023		U-759		
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 002	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023		U-759		
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 003	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023		U-759		
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 004	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023		U-759		
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 005	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022	DP			
	7101569	Oct 02, 2023		U-759		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 006	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Oct 02, 2023	DP		U-759	
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 007	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Oct 02, 2023	DP		U-759	
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 008	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Oct 02, 2023	DP		U-759	
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 009	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Oct 02, 2023	DP		U-759	
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 010	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Oct 02, 2023	DP		U-759	
<b><u>LEVOTHYROXINE SODIUM - LEVOXYL</u></b>						
N 021301 011	6555581	Feb 15, 2022				
	7067148	Feb 15, 2022				
	7101569	Oct 02, 2023	DP		U-759	
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 001	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 002	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 003	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 004	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 005	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 006	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 007	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 008	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 009	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 010	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - LEVO-T</u></b>						
N 021342 011	6399101	Mar 30, 2020				
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 002	7691411	Mar 14, 2024				
	7723390	Mar 14, 2024	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 002	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 003	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 004	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 005	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 006	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 007	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 008	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 009	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - TIROSINT</u></b>						
N 021924 010	7691411 7723390	Mar 14, 2024 Mar 14, 2024	DP DP			
<b><u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u></b>						
N 202231 001	9006289 9168238 9168239	Oct 03, 2032 Aug 29, 2032 Aug 29, 2032	DP DP DP			
<b><u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u></b>						
N 202231 002	9006289 9168238 9168239	Oct 03, 2032 Aug 29, 2032 Aug 29, 2032	DP DP DP			
<b><u>LEVOTHYROXINE SODIUM - LEVOTHYROXINE SODIUM</u></b>						
N 202231 003	9006289 9168238 9168239	Oct 03, 2032 Aug 29, 2032 Aug 29, 2032	DP DP DP			
<b><u>LIDOCAINE HYDROCHLORIDE - ZINGO</u></b>						
N 022114 001	8540665 9358338 9370622	Oct 22, 2029 Apr 27, 2035 Sep 28, 2035	U-1438 U-1870 U-1870			
<b><u>LIDOCAINE HYDROCHLORIDE - AKTEN</u></b>						
N 022221 001	8759401	Jul 24, 2026	DP U-1523			
<b><u>LIDOCAINE; TETRACAINE - SYNERA</u></b>						
N 021623 001	6465709	Jul 07, 2020	DP			
<b><u>LIDOCAINE; TETRACAINE - PLIAGLIS</u></b>						
N 021717 001	6528086	Sep 28, 2019	DP			
<b><u>LIFITEGRAST - XIIDRA</u></b>						
N 208073 001	7314938 7745460 7790743 7928122 8084047	Mar 10, 2025 Nov 05, 2024 Nov 05, 2024 Nov 05, 2024 May 17, 2026	DS DP DS DP U-1880 U-1880 DS DP DS DP	NCE		Jul 11, 2021

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LIFITEGRAST - XITDRA</b>						
N 208073 001	8168655	May 09, 2029	U-1880			
	8367701	Apr 15, 2029	DP U-1880			
	8592450	May 17, 2026	U-1880			
	8927574	Nov 12, 2030	DP			
	9085553	Jul 25, 2033	DP			
	9216174	Nov 05, 2024	DP			
	9353088	Oct 21, 2030	DP			
	9447077	Apr 15, 2029	U-1900			
<b>LINACLOTIDE - LINZESS</b>						
N 202811 001	7304036	Aug 30, 2026	DS DP U-1278		NCE	Aug 30, 2017
	7304036	Aug 30, 2026	DS DP U-1516			
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024	U-1278			
	8748573	Oct 30, 2031	U-1515			
	8748573	Oct 30, 2031	U-1516			
	8802628	Nov 17, 2031	DP			
	8933030	Feb 17, 2031	DP			
	9708371	Aug 16, 2033	DP U-1515			
	9708371	Aug 16, 2033	DP U-1516			
<b>LINACLOTIDE - LINZESS</b>						
N 202811 002	7304036	Aug 30, 2026	DS DP U-1278		NCE	Aug 30, 2017
	7304036	Aug 30, 2026	DS DP U-1516			
	7371727	Jan 28, 2024	DS			
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024	U-1278			
	8748573	Oct 30, 2031	U-1515			
	8748573	Oct 30, 2031	U-1516			
	8802628	Nov 17, 2031	DP			
	8933030	Feb 17, 2031	DP			
	9708371	Aug 16, 2033	DP U-1515			
<b>LINACLOTIDE - LINZESS</b>						
N 202811 003	7304036	Aug 30, 2026	DS DP U-1516		NCE	Aug 30, 2017
	7371727	Jan 28, 2024	DS		NS	Jan 25, 2020
	7704947	Jan 28, 2024	DS DP			
	7745409	Jan 28, 2024	DS DP			
	8080526	Jan 28, 2024	DS DP			
	8110553	Jan 28, 2024	U-1516			
	8933030	Feb 17, 2031	DP U-1516			
	9708371	Aug 16, 2033	DP U-1516			
<b>LINAGLIPTIN - TRADJENTA</b>						
N 201280 001	6890898	Feb 02, 2019	U-1270			
	6890898	Feb 02, 2019	U-493			
	7078381	Feb 02, 2019	U-1270			
	7078381	Feb 02, 2019	U-493			
	7407955	May 02, 2025	DS DP			
	7459428	Feb 02, 2019	U-1270			
	7459428	Feb 02, 2019	U-493			
	8119648	Aug 12, 2023	U-1270			
	8119648	Aug 12, 2023	U-774			
	8178541	Aug 12, 2023	U-1244			
	8178541	Aug 12, 2023	U-1245			
	8178541	Aug 12, 2023	U-1270			
	8178541	Aug 12, 2023	U-775			
	8673927	May 04, 2027	U-1503			
	8846695	Jun 04, 2030	U-1503		Y	
	8853156	Mar 05, 2031	U-1642			
	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP U-1503			
	9173859	May 04, 2027	DP U-1768			
	9486526	Aug 05, 2029	U-1915			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LINAGLIPITIN - TRADJENTA</b>						
N 201280 001	6890898	Feb 02, 2019	U-1270			
	6890898	Feb 02, 2019	U-493			
	7078381	Feb 02, 2019	U-1270			
	7078381	Feb 02, 2019	U-493			
	7407955	May 02, 2025	DS DP			
	7459428	Feb 02, 2019	U-1270			
	7459428	Feb 02, 2019	U-493			
	8119648	Aug 12, 2023	U-1270			
	8119648	Aug 12, 2023	U-774			
	8178541	Aug 12, 2023	U-1244			
	8178541	Aug 12, 2023	U-1245			
	8178541	Aug 12, 2023	U-1270			
	8178541	Aug 12, 2023	U-775			
	8673927	May 04, 2027	U-1503			
	8846695	Jun 04, 2030	U-1503		Y	
	8853156	Mar 05, 2031	U-1642			
	8883805	Nov 26, 2025	DP			
	9173859	May 04, 2027	DP U-1503			
	9173859	May 04, 2027	DP U-1768			
	9486526	Aug 05, 2029	U-1915			
<b>LINAGLIPITIN; METFORMIN HYDROCHLORIDE - JENTADUETO</b>						
N 201281 001	6890898	Feb 02, 2019	U-1039		M-146	Jul 30, 2017
	7078381	Feb 02, 2019	U-1039			
	7407955	May 02, 2025	DS DP			
	7459428	Feb 02, 2019	U-1039			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027	U-1503			
	8846695	Jun 04, 2030	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
<b>LINAGLIPITIN; METFORMIN HYDROCHLORIDE - JENTADUETO</b>						
N 201281 002	6890898	Feb 02, 2019	U-1039		M-146	Jul 30, 2017
	7078381	Feb 02, 2019	U-1039			
	7407955	May 02, 2025	DS DP			
	7459428	Feb 02, 2019	U-1039			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-775			
	8673927	May 04, 2027	U-1503			
	8846695	Jun 04, 2030	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
<b>LINAGLIPITIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</b>						
N 208026 001	6488962	Jun 20, 2020	DP			
	6890898	Feb 02, 2019	U-803			
	7078381	Feb 02, 2019	U-803			
	7407955	May 02, 2025	DS DP			
	7459428	Feb 02, 2019	U-803			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LINAGLITPIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</b>						
N 208026 001	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-1853			
	8673927	May 04, 2027	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
<b>LINAGLITPIN; METFORMIN HYDROCHLORIDE - JENTADUETO XR</b>						
N 208026 002	6488962	Jun 20, 2020	DP			
	6890898	Feb 02, 2019	U-803			
	7078381	Feb 02, 2019	U-803			
	7407955	May 02, 2025	DS DP			
	7459428	Feb 02, 2019	U-803			
	8119648	Aug 12, 2023	U-802			
	8178541	Aug 12, 2023	DP U-1853			
	8673927	May 04, 2027	U-1503			
	8883805	Nov 26, 2025	DP			
	9155705	May 21, 2030	DP			
	9173859	May 04, 2027	DP U-1503			
	9415016	Apr 02, 2029	DP			
	9555001	Mar 06, 2033	DP U-1967			
	9555001	Mar 06, 2033	DP U-1968			
<b>LINEZOLID - ZYVOX</b>						
N 021130 001	6514529	Mar 15, 2021	DP			
	6559305	Jan 29, 2021	DS			
<b>LINEZOLID - ZYVOX</b>						
N 021130 002	6514529	Mar 15, 2021	DP			
	6559305	Jan 29, 2021	DS			
<b>LINEZOLID - ZYVOX</b>						
N 021131 001	6559305	Jan 29, 2021	DS			
<b>LINEZOLID - ZYVOX</b>						
N 021131 002	6559305	Jan 29, 2021	DS			
	6559305*PED	Jul 29, 2021				
<b>LINEZOLID - ZYVOX</b>						
N 021131 003	6559305	Jan 29, 2021	DS			
	6559305*PED	Jul 29, 2021				
<b>LINEZOLID - ZYVOX</b>						
N 021132 001	6559305	Jan 29, 2021	DS			
<b>LIRAGLUTIDE RECOMBINANT - VICTOZA</b>						
N 022341 001	6004297	Jan 28, 2019	DP		I-750	Aug 25, 2020
	6268343	Aug 22, 2022	DS DP U-968		M-176	Apr 22, 2019
	6458924	Aug 22, 2017	DS DP U-968			
	7235627	Aug 22, 2017	DS DP			
	8114833	Aug 13, 2025	DP			
	8846618	Jun 27, 2022	DP			
	9265893	Sep 23, 2032	DP			
	RE41956	Jan 21, 2021	DP			
	RE43834	Jan 28, 2019	DP			
<b>LIRAGLUTIDE RECOMBINANT - SAXENDA</b>						
N 206321 001	6268343	Aug 22, 2022	DS DP U-1255			
	6458924	Aug 22, 2017	DS DP U-1255			
	6899699	Jan 01, 2022	DP			
	7235627	Aug 22, 2017	DS DP			
	8114833	Aug 13, 2025	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8846618	Jun 27, 2022	DP			
	8920383	Jul 17, 2026	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>LIRAGLUTIDE RECOMBINANT - SAXENDA</b>						
N 206321 001	9108002	Jan 26, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b>LISDEXAMFETAMINE DIMESYLAATE - VYVANSE</b>						
N 021977 001	7105486	Feb 24, 2023	U-727		I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		
<b>LISDEXAMFETAMINE DIMESYLAATE - VYVANSE</b>						
N 021977 002	7105486	Feb 24, 2023	U-727		I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		
<b>LISDEXAMFETAMINE DIMESYLAATE - VYVANSE</b>						
N 021977 003	7105486	Feb 24, 2023	U-727		I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 021977 004	7105486	Feb 24, 2023		U-727	I-703	Jan 30, 2018
	7105486	Feb 24, 2023		U-842	M-188	Oct 14, 2019
	7223735	Feb 24, 2023		DP		
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 021977 005	7105486	Feb 24, 2023		U-842	I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 021977 006	7105486	Feb 24, 2023		U-727	I-703	Jan 30, 2018
	7105486	Feb 24, 2023		U-842	M-188	Oct 14, 2019
	7223735	Feb 24, 2023	DP			
	7655630	Feb 24, 2023	DS			
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023	DP	U-842		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023	DP	U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-842		
	7723305	Feb 24, 2023	DP	U-842		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 021977 007	7223735	Feb 24, 2023	DP		I-703	Jan 30, 2018
	7655630	Feb 24, 2023	DS		M-188	Oct 14, 2019
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-1034		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 021977 007	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023		DP U-842		
	7678770	Feb 24, 2023		U-842		
	7678771	Feb 24, 2023		DP		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023		DP U-842		
	7700561	Feb 24, 2023		DP		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023		DP U-842		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 208510 001	7105486	Feb 24, 2023		U-727	I-703	Jan 30, 2018
	7223735	Feb 24, 2023		DP	M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS	DP		
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-727		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023		DP U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023		DP U-727		
	7678770	Feb 24, 2023		U-727		
	7678771	Feb 24, 2023		DP U-727		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023		DP U-727		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023		DP U-727		
	7723305	Feb 24, 2023		DP U-727		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 208510 002	7105486	Feb 24, 2023		U-727	I-703	Jan 30, 2018
	7223735	Feb 24, 2023		DP	M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS	DP		
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-727		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023		DP U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023		DP U-727		
	7678770	Feb 24, 2023		U-727		
	7678771	Feb 24, 2023		DP U-727		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023		DP U-727		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023		DP U-727		
	7723305	Feb 24, 2023		DP U-727		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 208510 003	7105486	Feb 24, 2023		U-727	I-703	Jan 30, 2018
	7223735	Feb 24, 2023		DP	M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS	DP		
	7659253	Feb 24, 2023	DS	DP U-727		
	7659254	Feb 24, 2023		U-727		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023		DP U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023		DP U-727		
	7678770	Feb 24, 2023		U-727		
	7678771	Feb 24, 2023		DP U-727		
	7687466	Feb 24, 2023		DP		
	7687467	Feb 24, 2023		DP U-727		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023		DP U-727		
	7723305	Feb 24, 2023		DP U-727		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 208510 004	7105486	Feb 24, 2023	U-727		I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023		U-727		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-727		
	7678770	Feb 24, 2023		U-727		
	7678771	Feb 24, 2023	DP	U-727		
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP	U-727		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-727		
	7723305	Feb 24, 2023	DP	U-727		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 208510 005	7105486	Feb 24, 2023	U-727		I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023		U-727		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-727		
	7678770	Feb 24, 2023		U-727		
	7678771	Feb 24, 2023	DP	U-727		
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP	U-727		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-727		
	7723305	Feb 24, 2023	DP	U-727		
<b>LISDEXAMFETAMINE Dimesylate - Vyvanse</b>						
N 208510 006	7105486	Feb 24, 2023	U-727		I-703	Jan 30, 2018
	7223735	Feb 24, 2023	DP		M-188	Oct 14, 2019
	7655630	Feb 24, 2023	DS DP			
	7659253	Feb 24, 2023	DS DP U-727			
	7659254	Feb 24, 2023		U-727		
	7662787	Feb 24, 2023	DS			
	7662788	Feb 24, 2023		U-727		
	7671030	Feb 24, 2023	DP	U-727		
	7671031	Feb 24, 2023		U-727		
	7674774	Feb 24, 2023	DP	U-727		
	7678770	Feb 24, 2023		U-727		
	7678771	Feb 24, 2023	DP	U-727		
	7687466	Feb 24, 2023	DP			
	7687467	Feb 24, 2023	DP	U-727		
	7713936	Feb 24, 2023		U-727		
	7718619	Feb 24, 2023	DP	U-727		
	7723305	Feb 24, 2023	DP	U-727		
<b>Lisinopril - Obrelis</b>						
N 208401 001	9463183	Nov 06, 2035	DP			
	9616096	Nov 06, 2035		U-1723		
	9616096	Nov 06, 2035		U-185		
	9616096	Nov 06, 2035		U-1864		
	9616096	Nov 06, 2035		U-1991		
	9616096	Nov 06, 2035		U-3		
	9616096	Nov 06, 2035		U-71		
	9616096	Nov 06, 2035		U-8		
	9814751	Nov 06, 2035	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LIXISENATIDE - ADLYXIN</u></b>						
N 208471 001	8475414	Dec 28, 2030	DP U-1881		NCE	Jul 27, 2021
	8882721	Jun 28, 2031	DP			
	8915888	Jun 08, 2030	DP U-1881			
	9072836	Mar 15, 2032	DP			
	9084853	Oct 05, 2031	DP			
	9308329	Dec 28, 2030	DP U-1881			
	9408893	Aug 27, 2032	U-1894			
	9511193	Jan 19, 2032	DP			
	9707176	Nov 11, 2030	DP			
	9821032	May 09, 2032	U-2200			
	RE45313	Jul 12, 2020	DS DP			
<b><u>LIXISENATIDE - ADLYXIN</u></b>						
N 208471 002	8475414	Dec 28, 2030	DP U-1881		NCE	Jul 27, 2021
	8882721	Jun 28, 2031	DP			
	8915888	Jun 08, 2030	DP U-1881			
	9072836	Mar 15, 2032	DP			
	9084853	Oct 05, 2031	DP			
	9308329	Dec 28, 2030	DP U-1881			
	9408893	Aug 27, 2032	U-1894			
	9511193	Jan 19, 2032	DP			
	9707176	Nov 11, 2030	DP			
	9821032	May 09, 2032	U-2200			
	RE45313	Jul 12, 2020	DS DP			
<b><u>LOMITAPIDE MESYLATE - JUXTAPID</u></b>						
N 203858 001	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			
<b><u>LOMITAPIDE MESYLATE - JUXTAPID</u></b>						
N 203858 002	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			
<b><u>LOMITAPIDE MESYLATE - JUXTAPID</u></b>						
N 203858 003	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			
<b><u>LOMITAPIDE MESYLATE - JUXTAPID</u></b>						
N 203858 004	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			
<b><u>LOMITAPIDE MESYLATE - JUXTAPID</u></b>						
N 203858 005	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>LOMITAPIDE MESYLATE - JUXTAPIID</u></b>						
N 203858 005	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			
<b><u>LOMITAPIDE MESYLATE - JUXTAPIID</u></b>						
N 203858 006	5712279	Feb 21, 2020	DS DP U-1317		NCE	Dec 21, 2017
	6492365	Dec 10, 2019	U-1318		ODE-36	Dec 21, 2019
	7932268	Aug 19, 2027	U-1316			
	8618135	Mar 07, 2025	U-1316			
	9265758	Mar 07, 2025	U-1316			
	9364470	Mar 07, 2025	U-1851			
	9433617	Mar 07, 2025	U-1316			
<b><u>LOPERAMIDE HYDROCHLORIDE - IMODIUM A-D EZ CHEWS</u></b>						
N 020448 001	6814978	Aug 26, 2021	DP			
<b><u>LOPERAMIDE HYDROCHLORIDE; SIMETHICONE - IMODIUM MULTI-SYMPTOM RELIEF</u></b>						
N 021140 001	6103260	Jul 17, 2017	DP			
<b><u>LOPINAVIR; RITONAVIR - KALETRA</u></b>						
N 021226 001	6232333	Nov 07, 2017				
	6458818	Nov 07, 2017				
	6521651	Nov 07, 2017	DP			
	7141593	May 22, 2020	DP			
	7432294	May 22, 2020	DP			
<b><u>LOPINAVIR; RITONAVIR - KALETRA</u></b>						
N 021251 001	6911214	Nov 28, 2021	DP U-895			
	8501219	Nov 28, 2021	DP			
<b><u>LOPINAVIR; RITONAVIR - KALETRA</u></b>						
N 021906 001	7148359	Jul 19, 2019	DP			
	7364752	Nov 10, 2020	DP U-688			
	8025899	Dec 14, 2027	DP			
	8025899*PED	Jun 14, 2028				
	8268349	Aug 25, 2024	DP			
	8309613	Dec 24, 2024	U-688			
	8377952	Oct 22, 2027	U-1372			
	8377952*PED	Apr 22, 2028				
	8399015	Aug 25, 2024	DP			
	8399015*PED	Feb 25, 2025				
	8470347	Sep 17, 2026	DP			
	8470347*PED	Mar 17, 2027				
	8691878	Aug 25, 2024	U-1513			
	8691878*PED	Feb 25, 2025				
<b><u>LOPINAVIR; RITONAVIR - KALETRA</u></b>						
N 021906 002	7148359	Jul 19, 2019	DP			
	7364752	Nov 10, 2020	DP U-688			
	8025899	Dec 14, 2027	DP			
	8025899*PED	Jun 14, 2028				
	8268349	Aug 25, 2024	DP			
	8309613	Dec 24, 2024	U-688			
	8377952	Oct 22, 2027	U-1372			
	8377952*PED	Apr 22, 2028				
	8399015	Aug 25, 2024	DP			
	8399015*PED	Feb 25, 2025				
	8470347	Sep 17, 2026	DP			
	8470347*PED	Mar 17, 2027				
	8691878	Aug 25, 2024	U-1513			
	8691878*PED	Feb 25, 2025				
<b><u>LORATADINE - CLARITIN</u></b>						
N 020641 002	6132758	Jun 01, 2018	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LORCASERIN HYDROCHLORIDE - BELVIO</b>						
N 022529 001	6953787	Apr 10, 2023	DS DP U-1252			
	6953787	Apr 10, 2023	DS DP U-1253			
	6953787	Apr 10, 2023	DS DP U-1254			
	6953787	Apr 10, 2023	DS DP U-1255			
	7514422	Apr 10, 2023	U-1252			
	7514422	Apr 10, 2023	U-1253			
	7514422	Apr 10, 2023	U-1254			
	7514422	Apr 10, 2023	U-1255			
	7977329	Apr 10, 2023	DS DP U-1252			
	7977329	Apr 10, 2023	DS DP U-1253			
	7977329	Apr 10, 2023	DS DP U-1254			
	7977329	Apr 10, 2023	DS DP U-1255			
	8168624	Apr 18, 2029	DS DP			
	8207158	Apr 10, 2023	U-1252			
	8207158	Apr 10, 2023	U-1253			
	8207158	Apr 10, 2023	U-1254			
	8207158	Apr 10, 2023	U-1255			
	8273734	Apr 10, 2023	U-1254			
	8273734	Apr 10, 2023	U-1255			
	8367657	Apr 10, 2023	DS DP U-1252			
	8367657	Apr 10, 2023	DS DP U-1253			
	8367657	Apr 10, 2023	DS DP U-1254			
	8367657	Apr 10, 2023	DS DP U-1255			
	8546379	Apr 10, 2023	DS DP U-1252			
	8546379	Apr 10, 2023	DS DP U-1253			
	8546379	Apr 10, 2023	DS DP U-1254			
	8546379	Apr 10, 2023	DS DP U-1255			
	8575149	Apr 10, 2023	U-1452			
	8697686	Dec 20, 2025	DS DP			
	8946207	Jun 16, 2024	DP			
	8980881	Dec 20, 2025	U-1252			
	8980881	Dec 20, 2025	U-1253			
	8980881	Dec 20, 2025	U-1254			
	8980881	Dec 20, 2025	U-1255			
	8999970	Feb 07, 2033	U-1688			
	8999970	Feb 07, 2033	U-1689			
	8999970	Feb 07, 2033	U-1692			
	9169213	Dec 06, 2032	U-1762			
	9169213	Dec 06, 2032	U-1763			
	9169213	Dec 06, 2032	U-1764			
	9169213	Dec 06, 2032	U-1765			
	9770455	Aug 31, 2031	U-2110			
<b>LORCASERIN HYDROCHLORIDE - BELVIO XR</b>						
N 208524 001	6953787	Apr 10, 2023	DS DP U-1252			
	6953787	Apr 10, 2023	DS DP U-1253			
	6953787	Apr 10, 2023	DS DP U-1254			
	6953787	Apr 10, 2023	DS DP U-1255			
	7514422	Apr 10, 2023	U-1252			
	7514422	Apr 10, 2023	U-1253			
	7514422	Apr 10, 2023	U-1254			
	7514422	Apr 10, 2023	U-1255			
	7977329	Apr 10, 2023	DS DP U-1252			
	7977329	Apr 10, 2023	DS DP U-1253			
	7977329	Apr 10, 2023	DS DP U-1254			
	7977329	Apr 10, 2023	DS DP U-1255			
	8168624	Apr 18, 2029	DS DP			
	8207158	Apr 10, 2023	U-1252			
	8207158	Apr 10, 2023	U-1253			
	8207158	Apr 10, 2023	U-1254			
	8207158	Apr 10, 2023	U-1255			
	8273734	Apr 10, 2023	U-1254			
	8273734	Apr 10, 2023	U-1255			
	8367657	Apr 10, 2023	DS DP U-1252			
	8367657	Apr 10, 2023	DS DP U-1253			
	8367657	Apr 10, 2023	DS DP U-1254			
	8367657	Apr 10, 2023	DS DP U-1255			
	8546379	Apr 10, 2023	DS DP U-1252			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>LORCASERIN HYDROCHLORIDE - BELVIQ XR</b>						
N 208524 001	8546379	Apr 10, 2023	DS DP U-1253			
	8546379	Apr 10, 2023	DS DP U-1254			
	8546379	Apr 10, 2023	DS DP U-1255			
	8575149	Apr 10, 2023		U-1452		
	8697686	Dec 20, 2025	DS DP			
	8946207	Jun 16, 2024	DP			
	8980881	Dec 20, 2025		U-1252		
	8980881	Dec 20, 2025		U-1253		
	8980881	Dec 20, 2025		U-1254		
	8980881	Dec 20, 2025		U-1255		
	8999970	Feb 07, 2033		U-1688		
	8999970	Feb 07, 2033		U-1689		
	8999970	Feb 07, 2033		U-1692		
	9169213	Dec 06, 2032		U-1884		
	9169213	Dec 06, 2032		U-1885		
	9169213	Dec 06, 2032		U-1886		
	9169213	Dec 06, 2032		U-1887		
	9770455	Aug 31, 2031		U-2110		
<b>LOVASTATIN - ALTOPREV</b>						
N 021316 001	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018		U-456		
	6485748	Dec 12, 2017		DP		
<b>LOVASTATIN - ALTOPREV</b>						
N 021316 002	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018		U-456		
	6485748	Dec 12, 2017		DP		
<b>LOVASTATIN - ALTOPREV</b>						
N 021316 003	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018		U-456		
	6485748	Dec 12, 2017		DP		
<b>LOVASTATIN - ALTOPREV</b>						
N 021316 004	5916595	Dec 12, 2017				
	6080778	Mar 23, 2018		U-456		
	6485748	Dec 12, 2017		DP		
<b>LOXAPINE - ADASUVE</b>						
N 022549 001	6716416	May 20, 2022		DP		
	7052679	Oct 26, 2021		DP		
	7078020	Oct 26, 2021		DP U-1375		
	7090830	Oct 26, 2021		DP		
	7458374	Aug 18, 2024		DP		
	7537009	Oct 28, 2024		DP		
	7585493	Oct 26, 2021		DP		
	7601337	Oct 26, 2021		DP		
	8074644	Jul 25, 2022		DP		
	8173107	Oct 26, 2021		DP		
	8235037	Oct 26, 2021		DP		
	8387612	Oct 23, 2026		DP		
	8955512	Oct 26, 2021		DP		
	8991387	May 21, 2024		DP		
	9370629	May 20, 2024		DP		
	9439907	Oct 26, 2021		DP		
	9440034	Oct 26, 2021		DP		
	9687487	Oct 26, 2021	DS DP			
<b>LUBIPROSTONE - AMITIZA</b>						
N 021908 001	6414016	Sep 05, 2020		U-1392		
	6414016	Sep 05, 2020		U-717		
	6583174	Oct 16, 2020		DP		
	6982283	Dec 04, 2022		U-1391		
	7064148	Aug 30, 2022		U-1404		
	7064148	Aug 30, 2022		U-739		
	7417067	Oct 16, 2020		DP		
	8026393	Oct 25, 2027		DP		
	8071613	Sep 05, 2020		U-1203		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LUBIPROSTONE - AMITIZA</b>						
N 021908 001	8071613	Sep 05, 2020		U-1393		
	8088934	May 18, 2021	DS			
	8097649	Oct 16, 2020		DP		
	8097653	Nov 14, 2022		U-1214		
	8097653	Nov 14, 2022		U-1394		
	8114890	Sep 05, 2020		DP		
	8338639	Jan 23, 2027		DP		
	8389542	Nov 14, 2022		DP U-1345		
	8389542	Nov 14, 2022		DP U-1395		
	8748481	Sep 01, 2025		U-1520		
	8779187	Jul 23, 2027		DP		
<b>LUBIPROSTONE - AMITIZA</b>						
N 021908 002	6414016	Sep 05, 2020		U-874		
	6583174	Oct 16, 2020	DP			
	7064148	Aug 30, 2022		U-739		
	7064148	Aug 30, 2022		U-873		
	7417067	Oct 16, 2020		DP		
	7795312	Sep 17, 2024		U-1085		
	8026393	Oct 25, 2027		DP		
	8071613	Sep 05, 2020		U-1202		
	8088934	May 18, 2021	DS			
	8097649	Oct 16, 2020		DP		
	8114890	Sep 05, 2020		DP		
	8338639	Jan 23, 2027		DP		
	8748481	Sep 01, 2025		U-1519		
	8779187	Jan 23, 2027		DP		
<b>LULICONAZOLE - LUZU</b>						
N 204153 001	5900488	Jan 18, 2020	DS DP		NCE	Nov 14, 2018
	8980931	Apr 28, 2034	DP			
	9012484	Sep 06, 2033	DS DP	U-540		
	9199977	Sep 06, 2033	DS DP			
	9453006	Sep 06, 2033	DS			
<b>LURASIDONE HYDROCHLORIDE - LATUDA</b>						
N 200603 001	5532372	Jul 02, 2018	DS		M-195	Jan 27, 2020
	5532372*PED	Jan 02, 2019			NPP	Jan 27, 2020
	8729085	May 26, 2026		DP	PED	Jul 27, 2020
	8729085*PED	Nov 26, 2026			PED	Jul 27, 2020
	8883794	May 26, 2026		DP		
	8883794*PED	Nov 26, 2026				
	9174975	Jun 25, 2026		U-1770		
	9174975*PED	Dec 25, 2026				
	9259423	May 23, 2031		U-1822		
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP	U-543		
	9815827	Feb 20, 2024		U-2166		
	9815827	Feb 20, 2024		U-543		
	9827242	May 23, 2031		U-2199		
	9827242	May 23, 2031		U-2201		
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<b>LURASIDONE HYDROCHLORIDE - LATUDA</b>						
N 200603 002	5532372	Jul 02, 2018	DS		NPP	Jan 27, 2020
	5532372*PED	Jan 02, 2019			PED	Jul 27, 2020
	8729085	May 26, 2026		DP		
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026		DP		
	8883794*PED	Nov 26, 2026				
	9174975	Jun 25, 2026		U-1770		
	9174975*PED	Dec 25, 2026				
	9259423	May 23, 2031		U-1822		
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP	U-543		
	9815827	Feb 20, 2024		U-2166		
	9815827	Feb 20, 2024		U-543		
	9827242	May 23, 2031		U-2199		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>LURASIDONE HYDROCHLORIDE - LATUDA</b>						
N 200603 002	9827242	May 23, 2031		U-2201		
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<b>LURASIDONE HYDROCHLORIDE - LATUDA</b>						
N 200603 003	5532372	Jul 02, 2018	DS		M-195	Jan 27, 2020
	5532372*PED	Jan 02, 2019			NPP	Jan 27, 2020
	8729085	May 26, 2026	DP		PED	Jul 27, 2020
	8729085*PED	Nov 26, 2026			PED	Jul 27, 2020
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9174975	Jun 25, 2026	U-1770			
	9174975*PED	Dec 25, 2026				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<b>LURASIDONE HYDROCHLORIDE - LATUDA</b>						
N 200603 004	5532372	Jul 02, 2018	DS			
	5532372*PED	Jan 02, 2019				
	8729085	May 26, 2026	DP			
	8729085*PED	Nov 26, 2026				
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9174975	Jun 25, 2026	U-1770			
	9174975*PED	Dec 25, 2026				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<b>LURASIDONE HYDROCHLORIDE - LATUDA</b>						
N 200603 005	5532372	Jul 02, 2018	DS		M-195	Jan 27, 2020
	5532372*PED	Jan 02, 2019			NPP	Jan 27, 2020
	8729085	May 26, 2026	DP		PED	Jul 27, 2020
	8729085*PED	Nov 26, 2026			PED	Jul 27, 2020
	8883794	May 26, 2026	DP			
	8883794*PED	Nov 26, 2026				
	9174975	Jun 25, 2026	U-1770			
	9174975*PED	Dec 25, 2026				
	9259423	May 23, 2031	U-1822			
	9259423*PED	Nov 23, 2031				
	9555027	May 26, 2026	DP U-543			
	9815827	Feb 20, 2024	U-2166			
	9815827	Feb 20, 2024	U-543			
	9827242	May 23, 2031	U-2199			
	9827242	May 23, 2031	U-2201			
	RE45573	Jun 23, 2025	DS			
	RE45573*PED	Dec 23, 2025				
<b>MACIMORELIN ACETATE - MACRILEN</b>						
N 205598 001					NCE	Dec 20, 2022
<b>MACITENTAN - OPSUMIT</b>						
N 204410 001	7094781	Oct 12, 2022	DS DP		NCE	Oct 18, 2018
	8268847	Apr 18, 2029		U-1446	ODE-54	Oct 18, 2020
	8367685	Oct 04, 2028	DP	U-1445		
	9265762	May 29, 2027	DP	U-1820		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MACITENTAN - OPSUMIT</u></b>						
N 204410 001	7094781	Oct 12, 2022	DS DP		NCE	Oct 18, 2018
	8268847	Apr 18, 2029		U-1446	ODE-54	Oct 18, 2020
	8367685	Oct 04, 2028	DP	U-1445		
	9265762	May 29, 2027	DP	U-1820		
<b><u>MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 25</u></b>						
N 021910 001	5945449	Oct 31, 2017	DP	U-785		
	7300674	Mar 04, 2023	DP	U-785		
<b><u>MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 35</u></b>						
N 021910 002	5945449	Oct 31, 2017	DP	U-785		
<b><u>MAGNESIUM SULFATE; POTASSIUM SULFATE; SODIUM SULFATE - SUPREP BOWEL PREP KIT</u></b>						
N 022372 001	6946149	Mar 07, 2023	DP	U-837		
<b><u>MALATHION - OVIDE</u></b>						
N 018613 001	7560445	Feb 01, 2027	DS	DP U-986		
	7977324	Aug 14, 2026		DP		
<b><u>MARAVIROC - SELZENTRY</u></b>						
N 022128 001	6586430	Dec 01, 2019	DS	DP U-824	NPP	Nov 04, 2019
	6667314	Aug 06, 2021	DS	DP U-824	NS	Nov 04, 2019
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<b><u>MARAVIROC - SELZENTRY</u></b>						
N 022128 002	6586430	Dec 01, 2019	DS	DP U-824	NPP	Nov 04, 2019
	6667314	Aug 06, 2021	DS	DP U-824	NS	Nov 04, 2019
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<b><u>MARAVIROC - SELZENTRY</u></b>						
N 022128 003	6586430	Dec 01, 2019	DS	DP U-824	NPP	Nov 04, 2019
	6667314	Aug 06, 2021	DS	DP U-824	NS	Nov 04, 2019
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<b><u>MARAVIROC - SELZENTRY</u></b>						
N 022128 004	6586430	Dec 01, 2019	DS	DP U-824	NPP	Nov 04, 2019
	6667314	Aug 06, 2021	DS	DP U-824	NS	Nov 04, 2019
	7368460	Nov 25, 2022		U-824		
	7576097	May 25, 2021	DS			
<b><u>MEBENDAZOLE - VERMOX</u></b>						
N 208398 001					NP	Nov 04, 2019
					NS	Oct 19, 2019
<b><u>MECASERMIN RECOMBINANT - INCRELEX</u></b>						
N 021839 001	5681814	Sep 18, 2017	DP			
<b><u>MECHLORETHAMINE HYDROCHLORIDE - VALCHLOR</u></b>						
N 202317 001	7838564	Mar 07, 2026	DP		ODE-51	Aug 23, 2020
	7872050	Jul 08, 2029		U-1427		
	8450375	Mar 07, 2026	DP			
	8501818	Mar 07, 2026	DP			
	8501819	Mar 07, 2026		U-1427		
	9382191	Mar 07, 2026	DP			
<b><u>MEDROXYPROGESTERONE ACETATE - DEPO-SUBQ PROVERA 104</u></b>						
N 021583 001	6495534	May 15, 2020	DP			
<b><u>MEGESTROL ACETATE - MEGACE ES</u></b>						
N 021778 001	6592903	Sep 21, 2020	DP			
	7101576	Apr 22, 2024		U-755		
	9040088	Apr 22, 2024	U-755			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MEGESTROL ACETATE - MEGACE ES</u></b>						
N 021778 001	9101540	Apr 22, 2024	DP	U-755		
	9101549	Apr 22, 2024		U-755		
	9107827	Apr 22, 2024		U-755		
<b><u>MELOXICAM - MOBIC</u></b>						
N 021530 001	6184220	Mar 25, 2019	DP			
<b><u>MELOXICAM - VIVLODEX</u></b>						
N 207233 001	9526734	Mar 31, 2033	DP		NP	Oct 22, 2018
	9649318	Mar 31, 2035	DP			
	9808468	Mar 31, 2035		U-2160		
	9808468	Mar 31, 2035		U-2165		
<b><u>MELOXICAM - VIVLODEX</u></b>						
N 207233 002	9526734	Mar 31, 2033	DP		NP	Oct 22, 2018
	9649318	Mar 31, 2035	DP			
	9808468	Mar 31, 2035		U-2160		
	9808468	Mar 31, 2035		U-2165		
<b><u>MELPHALAN HYDROCHLORIDE - EVOMELA</u></b>						
N 207155 001	8410077	Mar 13, 2029	DP		ODE-110	Mar 10, 2023
	9200088	Mar 13, 2029	DP			
	9493582	Feb 27, 2033	DP			
<b><u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u></b>						
N 022525 001	8039009	Mar 24, 2029	U-539		M-138	Jul 03, 2017
	8039009*PED	Sep 24, 2029			PED	Jan 03, 2018
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-539			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-539			
	8283379*PED	May 22, 2026				
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8362085	Nov 22, 2025	U-539			
	8362085*PED	May 22, 2026				
<b><u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u></b>						
N 022525 002	8039009	Mar 24, 2029	U-539		M-138	Jul 03, 2017
	8039009*PED	Sep 24, 2029			PED	Jan 03, 2018
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-539			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-539			
	8283379*PED	May 22, 2026				
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8362085	Nov 22, 2025	U-539			
	8362085*PED	May 22, 2026				
<b><u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u></b>						
N 022525 003	8039009	Mar 24, 2029	U-539		M-138	Jul 03, 2017
	8039009*PED	Sep 24, 2029			PED	Jan 03, 2018
	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-539			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-539			
	8283379*PED	May 22, 2026				
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8362085	Nov 22, 2025	U-539			
	8362085*PED	May 22, 2026				
<b><u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u></b>						
N 022525 004	8039009	Mar 24, 2029	U-539		M-138	Jul 03, 2017
	8039009*PED	Sep 24, 2029			PED	Jan 03, 2018

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MEMANTINE HYDROCHLORIDE - NAMENDA XR</u></b>						
N 022525 004	8168209	Nov 22, 2025	DP			
	8168209*PED	May 22, 2026				
	8173708	Nov 22, 2025	U-539			
	8173708*PED	May 22, 2026				
	8283379	Nov 22, 2025	U-539			
	8283379*PED	May 22, 2026				
	8329752	Nov 22, 2025	DP			
	8329752*PED	May 22, 2026				
	8362085	Nov 22, 2025	U-539			
	8362085*PED	May 22, 2026				
	8598233	Nov 22, 2025	DP			
	8598233*PED	May 22, 2026				
<b><u>MENTHOL; METHYL SALICYLATE - SALONPAS</u></b>						
N 022029 001	8809615	Jan 03, 2030	DP			
	9233184	Aug 01, 2027	DP			
<b><u>MENTHOL; METHYL SALICYLATE - SALONPAS</u></b>						
N 022029 002	8809615	Jan 03, 2030	DP			
	9233184	Aug 01, 2027	DP			
<b><u>MEQUINOL; TRETINOIN - SOLAGE</u></b>						
N 020922 001	6353029	Aug 24, 2020				
<b><u>MERCAPTOPURINE - PURIXAN</u></b>						
N 205919 001					ODE-65	Apr 28, 2021
<b><u>MEROPENEM; VABORBACTAM - VABOMERE</u></b>						
N 209776 001	8680136	Aug 17, 2031	DS DP			
	9694025	Aug 08, 2031	U-2120			
<b><u>MESALAMINE - MESALAMINE</u></b>						
A 091640 001					PC	Jan 14, 2018
<b><u>MESALAMINE - SFROWASA</u></b>						
N 019618 002	7645801	Jul 24, 2027	DS DP			
<b><u>MESALAMINE - CANASA</u></b>						
N 021252 001					M-187	Sep 02, 2019
<b><u>MESALAMINE - CANASA</u></b>						
N 021252 002	8217083	Jun 06, 2028	DP			
	8436051	Jun 06, 2028	DP			
<b><u>MESALAMINE - ASACOL HD</u></b>						
N 021830 001	6893662	Nov 15, 2021	DP U-141			
	8580302	Nov 15, 2021	DP			
	9089492	Nov 15, 2021	DP			
<b><u>MESALAMINE - LIALDA</u></b>						
N 022000 001	6773720	Jun 08, 2020	DP			
<b><u>MESALAMINE - APRISO</u></b>						
N 022301 001	6551620	Apr 20, 2018	DS DP U-907			
	8337886	Apr 20, 2018	DP U-1310			
	8496965	Apr 20, 2018	DP			
	8865688	May 01, 2030	U-1310			
	8911778	Apr 20, 2018	DP U-1310			
	8940328	Apr 20, 2018	DP			
	8956647	Apr 20, 2018	DP			
<b><u>MESALAMINE - DELZICOL</u></b>						
N 204412 001	6649180	Apr 13, 2020	DP			
<b><u>METAXALONE - SKELAXIN</u></b>						
N 013217 003	7122566	Feb 06, 2026	U-915			
	7714006	Dec 03, 2021	U-1050			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b>METFORMIN HYDROCHLORIDE - GLUCOPHAGE XR</b>						
N 021202 001	6475521	Mar 19, 2018				
	6660300	Mar 19, 2018		U-542		
<b>METFORMIN HYDROCHLORIDE - GLUCOPHAGE XR</b>						
N 021202 004	6475521	Mar 19, 2018				
	6660300	Mar 19, 2018		U-542		
<b>METFORMIN HYDROCHLORIDE - FORTAMET</b>						
N 021574 001	6099859	Mar 20, 2018		DP		
	6495162	Mar 20, 2018		DP		
	6790459	Mar 17, 2021			U-604	
	6866866	Mar 17, 2021		DP		
	7919116	Mar 20, 2018		DP		
	8475841	Mar 20, 2018			U-604	
<b>METFORMIN HYDROCHLORIDE - FORTAMET</b>						
N 021574 002	6099859	Mar 20, 2018		DP		
	6495162	Mar 20, 2018		DP		
	6790459	Mar 17, 2021			U-604	
	6866866	Mar 17, 2021		DP		
	7919116	Mar 20, 2018		DP		
	8475841	Mar 20, 2018			U-604	
<b>METFORMIN HYDROCHLORIDE - RIOMET</b>						
N 021591 001	6890957	Sep 14, 2023		DP		
<b>METFORMIN HYDROCHLORIDE - GLUMETZA</b>						
N 021748 001	6488962	Jun 20, 2020	DS DP			
	6723340	Oct 25, 2021	DS DP			
<b>METFORMIN HYDROCHLORIDE - GLUMETZA</b>						
N 021748 002	6488962	Jun 20, 2020	DS DP			
	7780987	Mar 23, 2025	DS DP			
	8323692	Mar 23, 2025	DP			
<b>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</b>						
N 021842 001	9101660	Jan 22, 2027	DP			
	9320714	Feb 03, 2029	DP			
<b>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET</b>						
N 021842 002	9101660	Jan 22, 2027	DP			
	9320714	Feb 03, 2029	DP			
<b>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</b>						
N 022024 001	6099859	Mar 20, 2018	DP			
	6495162	Mar 20, 2018	DP			
	6790459	Mar 17, 2021		U-974		
	6866866	Mar 17, 2021		DP		
	7785627	Jul 31, 2026		DP		
	7919116	Mar 20, 2018			U-1120	
	7919116	Mar 20, 2018			U-973	
	7959946	Jul 31, 2026		DP		
	8470368	Sep 19, 2023		DP		
	8475841	Mar 20, 2018			U-973	
	8668931	Sep 19, 2023		DP		
	9060941	Sep 19, 2023		DP		
<b>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</b>						
N 022024 002	6099859	Mar 20, 2018	DP			
	6495162	Mar 20, 2018	DP			
	6790459	Mar 17, 2021		U-974		
	6866866	Mar 17, 2021		DP		
	7785627	Jul 31, 2026		DP		
	7919116	Mar 20, 2018			U-1120	
	7919116	Mar 20, 2018			U-973	
	7959946	Jul 31, 2026		DP		
	8470368	Sep 19, 2023		DP		
	8475841	Mar 20, 2018			U-973	
	8668931	Sep 19, 2023		DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE - ACTOPLUS MET XR</b>						
N 022024 002	9060941	Sep 19, 2023		DP		
<b>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</b>						
N 021410 001	7358366	Apr 19, 2020	DS			
	8236345	Oct 07, 2022	DP			
<b>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</b>						
N 021410 002	7358366	Apr 19, 2020	DS			
	8236345	Oct 07, 2022	DP			
<b>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</b>						
N 021410 003	7358366	Apr 19, 2020	DS			
	8236345	Oct 07, 2022	DP			
<b>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</b>						
N 021410 005	7358366	Apr 19, 2020	DS			
	8236345	Oct 07, 2022	DP			
<b>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</b>						
N 200678 001	8628799	Jul 13, 2025	DP		M-175	Apr 05, 2019
	9339472	Jul 13, 2025	DP		M-198	Feb 27, 2020
RE44186		Jul 31, 2023	DS DP U-1097			
RE44186		Jul 31, 2023	DS DP U-1838			
<b>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</b>						
N 200678 002	9339472	Jul 13, 2025	DP		M-175	Apr 05, 2019
RE44186		Jul 31, 2023	DS DP U-1097		M-198	Feb 27, 2020
RE44186		Jul 31, 2023	DS DP U-1838			
<b>METFORMIN HYDROCHLORIDE; SAXAGLIPTIN HYDROCHLORIDE - KOMBIGLYZE XR</b>						
N 200678 003	9339472	Jul 13, 2025	DP		M-175	Apr 05, 2019
RE44186		Jul 31, 2023	DS DP U-1097		M-198	Feb 27, 2020
RE44186		Jul 31, 2023	DS DP U-1838			
<b>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</b>						
N 022044 001	6699871	Jul 26, 2022	DS DP U-802			
	6890898	Feb 02, 2019	U-1996			
7078381		Feb 02, 2019	U-1996			
7125873		Jul 26, 2022	DP U-1036			
7125873		Jul 26, 2022	DP U-1038			
7125873		Jul 26, 2022	DP U-803			
7326708		Nov 24, 2026	DS DP U-802			
7459428		Feb 02, 2019	U-1996			
8414921		Jul 21, 2028	DP U-1036			
<b>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</b>						
N 022044 002	6699871	Jul 26, 2022	DS DP U-802			
	6890898	Feb 02, 2019	U-1996			
7078381		Feb 02, 2019	U-1996			
7125873		Jul 26, 2022	DP U-1036			
7125873		Jul 26, 2022	DP U-1038			
7125873		Jul 26, 2022	DP U-803			
7326708		Nov 24, 2026	DS DP U-802			
7459428		Feb 02, 2019	U-1996			
8414921		Jul 21, 2028	DP U-1036			
<b>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</b>						
N 202270 001	6699871	Jul 26, 2022	DS DP U-1227			
	6890898	Feb 02, 2019	U-1996			
7078381		Feb 02, 2019	U-1996			
7125873		Jul 26, 2022	DP U-1227			
7326708		Nov 24, 2026	DS DP U-1227			
7459428		Feb 02, 2019	U-1996			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u></b>						
N 202270 002	6699871	Jul 26, 2022	DS DP U-1227			
	6890898	Feb 02, 2019		U-1996		
	7078381	Feb 02, 2019		U-1996		
	7125873	Jul 26, 2022		DP U-1227		
	7326708	Nov 24, 2026	DS DP	U-1227		
	7459428	Feb 02, 2019		U-1996		
<b><u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET XR</u></b>						
N 202270 003	6699871	Jul 26, 2022	DS DP	U-1227		
	6890898	Feb 02, 2019		U-1996		
	7078381	Feb 02, 2019		U-1996		
	7125873	Jul 26, 2022		DP U-1227		
	7326708	Nov 24, 2026	DS DP	U-1227		
	7459428	Feb 02, 2019		U-1996		
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 001	6746429	Apr 12, 2020		DP		
	7744582	Aug 10, 2019		DP U-1442		
	7776015	Aug 10, 2019		DP		
	8021335	Oct 04, 2026		DP		
	8480631	Mar 19, 2030		DP U-1442		
	8562564	Jan 24, 2026		DP		
	8579865	Mar 19, 2030		DP U-1442		
	8945063	Mar 19, 2030		DP U-1442		
	9421333	Mar 19, 2030		DP U-1442		
	9533102	Jan 24, 2026		DP		
	9629959	Jan 24, 2026		DP		
	RE44846	Aug 10, 2019		DP		
	RE44847	Aug 10, 2019		DP U-1442		
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 002	6746429	Apr 12, 2020		DP		
	7744582	Aug 10, 2019		DP U-1442		
	7776015	Aug 10, 2019		DP		
	8021335	Oct 04, 2026		DP		
	8480631	Mar 19, 2030		DP U-1442		
	8562564	Jan 24, 2026		DP		
	8579865	Mar 19, 2030		DP U-1442		
	8945063	Mar 19, 2030		DP U-1442		
	9421333	Mar 19, 2030		DP U-1442		
	9533102	Jan 24, 2026		DP		
	9629959	Jan 24, 2026		DP		
	RE44846	Aug 10, 2019		DP		
	RE44847	Aug 10, 2019		DP U-1442		
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 003	6746429	Apr 12, 2020		DP		
	7744582	Aug 10, 2019		DP U-1442		
	7776015	Aug 10, 2019		DP		
	8021335	Oct 04, 2026		DP		
	8480631	Mar 19, 2030		DP U-1442		
	8562564	Jan 24, 2026		DP		
	8579865	Mar 19, 2030		DP U-1442		
	8945063	Mar 19, 2030		DP U-1442		
	9421333	Mar 19, 2030		DP U-1442		
	9533102	Jan 24, 2026		DP		
	9629959	Jan 24, 2026		DP		
	RE44846	Aug 10, 2019		DP		
	RE44847	Aug 10, 2019		DP U-1442		
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 004	6746429	Apr 12, 2020		DP		
	7744582	Aug 10, 2019		DP U-1442		
	7776015	Aug 10, 2019		DP		
	8021335	Oct 04, 2026		DP		
	8480631	Mar 19, 2030		DP U-1442		
	8562564	Jan 24, 2026		DP		
	8579865	Mar 19, 2030		DP U-1442		
	8945063	Mar 19, 2030		DP U-1442		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 004	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	RE44846	Aug 10, 2019	DP			
	RE44847	Aug 10, 2019	DP U-1442			
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 005	6746429	Apr 12, 2020	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	RE44846	Aug 10, 2019	DP			
	RE44847	Aug 10, 2019	DP U-1442			
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 006	6746429	Apr 12, 2020	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	RE44846	Aug 10, 2019	DP			
	RE44847	Aug 10, 2019	DP U-1442			
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 007	6746429	Apr 12, 2020	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	RE44846	Aug 10, 2019	DP			
	RE44847	Aug 10, 2019	DP U-1442			
<b><u>METHOTREXATE - OTREXUP</u></b>						
N 204824 008	6746429	Apr 12, 2020	DP			
	8021335	Oct 04, 2026	DP			
	8480631	Mar 19, 2030	DP U-1442			
	8562564	Jan 24, 2026	DP			
	8579865	Mar 19, 2030	DP U-1442			
	8945063	Mar 19, 2030	DP U-1442			
	9421333	Mar 19, 2030	DP U-1442			
	9533102	Jan 24, 2026	DP			
	9629959	Jan 24, 2026	DP			
	RE44846	Aug 10, 2019	DP			
	RE44847	Aug 10, 2019	DP U-1442			
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 001	8664231	Jun 01, 2029	U-1442			
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 002	8664231	Jun 01, 2029	U-1442			
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 003	8664231	Jun 01, 2029	U-1442			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 004	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 005	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 006	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 007	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 008	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 009	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE - RASUVO</u></b>						
N 205776 010	8664231	Jun 01, 2029		U-1442		
<b><u>METHOTREXATE SODIUM - XATMEP</u></b>						
N 208400 001	9259427	Jan 02, 2033	DP		ODE-137 ODE-138	Apr 25, 2024 Apr 25, 2024
<b><u>METHYLENE BLUE - PROVAYBLUE</u></b>						
N 204630 001					ODE-113	Apr 08, 2023
<b><u>METHYLNALTREXONE BROMIDE - RELISTOR</u></b>						
N 021964 001	6559158	Nov 03, 2017	U-1185			
	8247425	Dec 31, 2030	U-1185			
	8420663	Sep 30, 2029	U-1185			
	8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP U-1185			
	9180125	Sep 30, 2029	DP U-1185			
	9492445	Sep 30, 2029	DP U-1185			
	9669096	Apr 08, 2024	DP			
<b><u>METHYLNALTREXONE BROMIDE - RELISTOR</u></b>						
N 021964 002	6559158	Nov 03, 2017	U-1185			
	8247425	Dec 31, 2030	U-1185			
	8420663	Sep 30, 2029	U-1185			
	8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP U-1185			
	9180125	Sep 30, 2029	DP U-1185			
	9492445	Sep 30, 2029	DP U-1185			
	9669096	Apr 08, 2024	DP			
<b><u>METHYLNALTREXONE BROMIDE - RELISTOR</u></b>						
N 021964 003	8247425	Dec 31, 2030	U-1185			
	8420663	Sep 30, 2029	U-1185			
	8552025	Apr 08, 2024	DP			
	8822490	Sep 30, 2029	DP U-1185			
	9180125	Sep 30, 2029	DP U-1185			
	9492445	Sep 30, 2029	DP U-1185			
	9669096	Apr 08, 2024	DP			
<b><u>METHYLNALTREXONE BROMIDE - RELISTOR</u></b>						
N 208271 001	6559158	Nov 03, 2017	U-1185		NP	Jul 19, 2019
	8420663	Sep 30, 2029	U-1185			
	8524276	Mar 10, 2031	DP			
	8956651	Mar 10, 2031	DP			
	9180125	Sep 30, 2029	DP U-1185			
	9314461	Mar 10, 2031	DP			
	9492445	Sep 30, 2029	DP U-1185			
	9724343	Sep 30, 2029	DP U-1185			
<b><u>METHYLPHENIDATE - DAYTRANA</u></b>						
N 021514 001	6210705	Sep 30, 2018	DP U-727			
	6348211	Sep 30, 2018	DP U-727			
	8632802	Oct 07, 2025	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>METHYLPHENIDATE - DAYTRANA</b>						
N 021514 001	9034370	Oct 07, 2025	DP			
	9668981	Oct 07, 2025		U-2024		
<b>METHYLPHENIDATE - DAYTRANA</b>						
N 021514 002	6210705	Sep 30, 2018	DP	U-727		
	6348211	Sep 30, 2018		DP	U-727	
	8632802	Oct 07, 2025		DP		
	9034370	Oct 07, 2025		DP		
	9668981	Oct 07, 2025		U-2024		
<b>METHYLPHENIDATE - DAYTRANA</b>						
N 021514 003	6210705	Sep 30, 2018	DP	U-727		
	6348211	Sep 30, 2018		DP	U-727	
	8632802	Oct 07, 2025		DP		
	9034370	Oct 07, 2025		DP		
	9668981	Oct 07, 2025		U-2024		
<b>METHYLPHENIDATE - DAYTRANA</b>						
N 021514 004	6210705	Sep 30, 2018	DP	U-727		
	6348211	Sep 30, 2018		DP	U-727	
	8632802	Oct 07, 2025		DP		
	9034370	Oct 07, 2025		DP		
	9668981	Oct 07, 2025		U-2024		
<b>METHYLPHENIDATE - COTEMPLA XR-ODT</b>						
N 205489 001	8840924	Jun 05, 2026	DP		NP	Jun 19, 2020
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<b>METHYLPHENIDATE - COTEMPLA XR-ODT</b>						
N 205489 002	8840924	Jun 05, 2026	DP		NP	Jun 19, 2020
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<b>METHYLPHENIDATE - COTEMPLA XR-ODT</b>						
N 205489 003	8840924	Jun 05, 2026	DP		NP	Jun 19, 2020
	9072680	Jun 28, 2032	DP			
	9089496	Jun 28, 2032	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</b>						
N 021121 001	6919373	Jul 31, 2017	U-666			
	6930129	Jul 31, 2017	U-666			
	8163798	Jul 31, 2017	DP			
	8629179	Jul 31, 2017	DP			
	8629179*PED	Jan 31, 2018				
	9000038	Jul 31, 2017	U-1693			
	9000038	Jul 31, 2017	U-666			
	9000038*PED	Jan 31, 2018				
	9029416	Jul 31, 2017	DP	U-666		
	9144549	Jul 31, 2017		U-1747		
	9144549	Jul 31, 2017		U-1748		
<b>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</b>						
N 021121 002	6919373	Jul 31, 2017	U-666			
	6930129	Jul 31, 2017	U-666			
	8163798	Jul 31, 2017	DP			
	8629179	Jul 31, 2017	DP			
	8629179*PED	Jan 31, 2018				
	9000038	Jul 31, 2017	U-1693			
	9000038	Jul 31, 2017	U-666			
	9000038*PED	Jan 31, 2018				
	9029416	Jul 31, 2017	DP	U-666		
	9144549	Jul 31, 2017		U-1747		
	9144549	Jul 31, 2017		U-1748		
<b>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</b>						
N 021121 003	6919373	Jul 31, 2017	U-666			
	6930129	Jul 31, 2017	U-666			
	8163798	Jul 31, 2017	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</b>						
N 021121 003	8629179	Jul 31, 2017	DP			
	8629179*PED	Jan 31, 2018				
	9000038	Jul 31, 2017		U-1693		
	9000038	Jul 31, 2017		U-666		
	9000038*PED	Jan 31, 2018				
	9029416	Jul 31, 2017	DP	U-666		
	9144549	Jul 31, 2017		U-1747		
	9144549	Jul 31, 2017		U-1748		
<b>METHYLPHENIDATE HYDROCHLORIDE - CONCERTA</b>						
N 021121 004	6919373	Jul 31, 2017		U-666		
	6930129	Jul 31, 2017		U-666		
	8163798	Jul 31, 2017	DP			
	8629179	Jul 31, 2017	DP			
	8629179*PED	Jan 31, 2018				
	9000038	Jul 31, 2017		U-1693		
	9000038	Jul 31, 2017		U-666		
	9000038*PED	Jan 31, 2018				
	9029416	Jul 31, 2017	DP	U-666		
	9144549	Jul 31, 2017		U-1747		
	9144549	Jul 31, 2017		U-1748		
<b>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</b>						
N 021259 001	6344215	Oct 27, 2020	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</b>						
N 021259 002	6344215	Oct 27, 2020	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</b>						
N 021259 003	6344215	Oct 27, 2020	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</b>						
N 021259 004	6344215	Oct 27, 2020	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</b>						
N 021284 001	6228398	Nov 01, 2019	DP	U-472		
<b>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</b>						
N 021284 002	6228398	Nov 01, 2019	DP	U-472		
<b>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</b>						
N 021284 003	6228398	Nov 01, 2019	DP	U-472		
<b>METHYLPHENIDATE HYDROCHLORIDE - RITALIN LA</b>						
N 021284 004	6228398	Nov 01, 2019	DP	U-472		
<b>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</b>						
N 021419 001	7691880	Oct 07, 2024	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - METHYLIN</b>						
N 021419 002	7691880	Oct 07, 2024	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - QUILLIVANT XR</b>						
N 202100 001	8062667	Mar 29, 2029	DP			
	8287903	Feb 15, 2031	DP			
	8465765	Feb 15, 2031	DP	U-1415		
	8563033	Feb 15, 2031	DP	U-1415		
	8778390	Feb 15, 2031	DP	U-1543		
	8956649	Feb 15, 2031	DP	U-1665		
	9040083	Feb 15, 2031	DP			
<b>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</b>						
N 205831 001	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP		Y	
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u></b>						
N 205831 002	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP	Y		
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u></b>						
N 205831 003	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP	Y		
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u></b>						
N 205831 004	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP	Y		
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u></b>						
N 205831 005	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP	Y		
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u></b>						
N 205831 006	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP	Y		
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - APTENSIO XR</u></b>						
N 205831 007	6419960	Dec 16, 2019	DP		NP	Apr 17, 2018
	7083808	Dec 16, 2019	DP			
	7247318	Dec 16, 2019	DP			
	7438930	Dec 16, 2019	DP	Y		
	8580310	Dec 16, 2019	DP			
	9066869	Dec 16, 2019	DP			
	9801823	Dec 16, 2019	DP			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u></b>						
N 207960 001	8202537	Mar 15, 2027	DP		NP	Dec 04, 2018
	8287903	Feb 15, 2031	DP			
	8999386	Aug 14, 2033	DP			
	9295642	Aug 14, 2033	DP U-1827			
	9545399	Aug 14, 2033	DP U-1827			
	9844544	Aug 14, 2033	DP U-2203			
<b><u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u></b>						
N 207960 002	8202537	Mar 15, 2027	DP		NP	Dec 04, 2018
	8287903	Feb 15, 2031	DP			
	8999386	Aug 14, 2033	DP			
	9295642	Aug 14, 2033	DP U-1827			
	9545399	Aug 14, 2033	DP U-1827			
	9844544	Aug 14, 2033	DP U-2203			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>METHYLPHENIDATE HYDROCHLORIDE - QUILLICHEW ER</u></b>						
N 207960 003	8202537	Mar 15, 2027	DP		NP	Dec 04, 2018
	8287903	Feb 15, 2031	DP			
	8999386	Aug 14, 2033	DP			
	9295642	Aug 14, 2033	DP U-1827			
	9545399	Aug 14, 2033	DP U-1827			
	9844544	Aug 14, 2033	DP U-2203			
<b><u>METOCLOPRAMIDE HYDROCHLORIDE - REGLAN ODT</u></b>						
N 021793 001	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>METOCLOPRAMIDE HYDROCHLORIDE - REGLAN ODT</u></b>						
N 021793 002	6024981	Apr 09, 2018	DP			
	6221392	Apr 09, 2018	DP			
<b><u>METOCLOPRAMIDE HYDROCHLORIDE - METOZOLV ODT</u></b>						
N 022246 001	6413549	Jul 11, 2017	DP			
<b><u>METOCLOPRAMIDE HYDROCHLORIDE - METOZOLV ODT</u></b>						
N 022246 002	6413549	Jul 11, 2017	DP			
<b><u>METRONIDAZOLE - METROGEL</u></b>						
N 021789 001	6881726	Feb 21, 2022	DP U-743			
	7348317	Feb 21, 2022	DP U-743			
<b><u>METRONIDAZOLE - VANDAZOLE</u></b>						
N 021806 001	7456207	Sep 22, 2024	DP			
<b><u>METRONIDAZOLE - NUVESSA</u></b>						
N 205223 001	7893097	Feb 19, 2028	DP			
	8658678	Jun 27, 2028		U-1682		
	8877792	Feb 02, 2028	DP			
	8946276	Jun 28, 2032		U-1664		
	9198858	Jun 28, 2032		U-1664		
<b><u>MICAFUNGIN SODIUM - MYCAMINE</u></b>						
N 021506 002	6107458	Mar 16, 2019	DS DP U-650			
	6107458	Mar 16, 2019	DS DP U-845			
	6774104	Jan 08, 2021	DP U-650			
	6774104	Jan 08, 2021	DP U-845			
<b><u>MICAFUNGIN SODIUM - MYCAMINE</u></b>						
N 021506 003	6107458	Mar 16, 2019	DS DP U-650			
	6107458	Mar 16, 2019	DS DP U-845			
	6774104	Jan 08, 2021	DP U-650			
	6774104	Jan 08, 2021	DP U-845			
<b><u>MICONAZOLE - ORAVIG</u></b>						
N 022404 001	6916485	Sep 11, 2022	DP U-1051			
	7651698	Sep 11, 2022		U-1051		
	8518442	Sep 11, 2022	DP			
<b><u>MICONAZOLE NITRATE; MICONAZOLE NITRATE - MONISTAT 1 COMBINATION PACK</u></b>						
N 021308 001	6153635	Nov 28, 2020			Y	
<b><u>MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE - VUSION</u></b>						
N 021026 001	8147852	Mar 30, 2028	U-1426			
<b><u>MIDOSTAURIN - RYDAPT</u></b>						
N 207997 001	7973031	Oct 17, 2024	U-2007		NCE	Apr 28, 2022
	8222244	Oct 29, 2022	U-2007		ODE-140	Apr 28, 2024
	8575146	Dec 02, 2030	U-2008		ODE-141	Apr 28, 2024
<b><u>MIFEPRISTONE - KORLYM</u></b>						
N 202107 001	8921348	Aug 27, 2028	U-1643		ODE-22	Feb 17, 2019
	9829495	Aug 15, 2036	U-1643			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u></b>						
N 022256 001	6602911	Jan 14, 2023	U-882			
	6992110	Nov 05, 2021	U-882			
	7888342	Nov 05, 2021	U-882			
	7994220	Sep 19, 2029	U-819			
<b><u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u></b>						
N 022256 002	6602911	Jan 14, 2023	U-882			
	6992110	Nov 05, 2021	U-882			
	7888342	Nov 05, 2021	U-882			
	7994220	Sep 19, 2029	U-819			
<b><u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u></b>						
N 022256 003	6602911	Jan 14, 2023	U-882			
	6992110	Nov 05, 2021	U-882			
	7888342	Nov 05, 2021	U-882			
	7994220	Sep 19, 2029	U-819			
<b><u>MILNACIPRAN HYDROCHLORIDE - SAVELLA</u></b>						
N 022256 004	6602911	Jan 14, 2023	U-882			
	6992110	Nov 05, 2021	U-882			
	7888342	Nov 05, 2021	U-882			
	7994220	Sep 19, 2029	U-819			
<b><u>MILTEFOSINE - IMPAVIDO</u></b>						
N 204684 001					NCE	Mar 19, 2019
					ODE-63	Mar 19, 2021
<b><u>MINOCYCLINE HYDROCHLORIDE - MINOCIN</u></b>						
N 050444 001	9084802	May 12, 2031	U-282			
	9278105	May 12, 2031	U-282			
<b><u>MINOCYCLINE HYDROCHLORIDE - ARESTIN</u></b>						
N 050781 001	6682348	Mar 29, 2022	DP			
	7699609	Mar 29, 2022	DP			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 001	5908838	Feb 19, 2018	U-917			
	7790705	Jun 24, 2025	U-1078			
	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 002	5908838	Feb 19, 2018	U-917			
	7541347	Apr 02, 2027	U-917			
	7544373	Apr 02, 2027	DP			
	7790705	Jun 24, 2025	U-1078			
	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 003	5908838	Feb 19, 2018	U-917			
	7790705	Jun 24, 2025	U-1078			
	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 004	5908838	Feb 19, 2018	U-917			
	7790705	Jun 24, 2025	U-1078			
	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
	9192615	Nov 17, 2031	DP			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 005	5908838	Feb 19, 2018	U-917			
	7790705	Jun 24, 2025	U-1078			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 005	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
	9192615	Nov 17, 2031	DP			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 006	5908838	Feb 19, 2018	U-917			
	7790705	Jun 24, 2025	U-1078			
	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
	8722650	Jun 24, 2025	U-1078			
<b><u>MINOCYCLINE HYDROCHLORIDE - SOLODYN</u></b>						
N 050808 007	5908838	Feb 19, 2018	U-917			
	7790705	Jun 24, 2025	U-1078			
	7919483	Mar 07, 2027	U-1078			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-1078			
	8722650	Jun 24, 2025	U-1078			
<b><u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u></b>						
N 201922 001	5908838	Feb 19, 2018	U-1376			
	7541347	Apr 02, 2027	U-917			
	7544373	Apr 02, 2027	DP			
	7790705	Jun 24, 2025	U-124			
	7919483	Mar 07, 2027	U-124			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-124			
<b><u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u></b>						
N 201922 003	5908838	Feb 19, 2018	U-1376			
	7541347	Apr 02, 2027	U-917			
	7544373	Apr 02, 2027	DP			
	7790705	Jun 24, 2025	U-124			
	7919483	Mar 07, 2027	U-124			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-124			
<b><u>MINOCYCLINE HYDROCHLORIDE - XIMINO</u></b>						
N 201922 005	5908838	Feb 19, 2018	U-1376			
	7541347	Apr 02, 2027	U-917			
	7544373	Apr 02, 2027	DP			
	7790705	Jun 24, 2025	U-124			
	7919483	Mar 07, 2027	U-124			
	8252776	Jun 24, 2025	U-124			
	8268804	Jun 24, 2025	U-124			
<b><u>MINOXIDIL - MEN'S ROGAINE</u></b>						
N 021812 001	6946120	Apr 20, 2019	DP U-702			
<b><u>MINOXIDIL - WOMEN'S ROGAINE</u></b>						
N 021812 002	6946120	Apr 20, 2019	DP U-702			
<b><u>MIPOMERSEN SODIUM - KYNAMRO</u></b>						
N 203568 001	6166197	Dec 26, 2017	DS		NCE	Jan 29, 2018
	7015315	Mar 21, 2023	DS		ODE-41	Jan 29, 2020
	7101993	Sep 05, 2023	DS			
	7407943	Aug 01, 2021	U-1353			
	7511131	Jan 29, 2027	DS			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MIRABEGRON - MYRBETRIO</u></b>						
N 202611 001	6346532	Mar 27, 2022	DS DP			
	6562375	Aug 01, 2020	DP			
	7342117	Nov 04, 2023	DS			
	7982049	Nov 04, 2023	DP			
	8835474	Nov 04, 2023		U-1527		
	RE44872	Nov 04, 2023		U-1527		
<b><u>MIRABEGRON - MYRBETRIO</u></b>						
N 202611 002	6346532	Mar 27, 2022	DS DP			
	6562375	Aug 01, 2020	DP			
	7342117	Nov 04, 2023	DS			
	7982049	Nov 04, 2023	DP			
	8835474	Nov 04, 2023		U-1527		
	RE44872	Nov 04, 2023		U-1527		
<b><u>MITOMYCIN - MITOSOL</u></b>						
N 022572 001	7806265	Feb 01, 2029	DP		ODE-21	
	8186511	Jul 19, 2026	DP			Feb 07, 2019
	9205075	Jul 19, 2026	DP			
	9539241	Jan 02, 2028	DS DP	U-2095		
	9649428	May 21, 2029		U-2095		
<b><u>MODAFINIL - PROVIGIL</u></b>						
N 020717 001	7297346	Nov 29, 2023	DP			
<b><u>MODAFINIL - PROVIGIL</u></b>						
N 020717 002	7297346	Nov 29, 2023	DP			
<b><u>MOMETASONE FUROATE - NASONEX</u></b>						
N 020762 001	6127353	Oct 03, 2017	DS DP			
<b><u>MOMETASONE FUROATE - ASMANEX TWISTHALER</u></b>						
N 021067 001	6503537	Mar 17, 2018	DP			
	8173172	Mar 17, 2018	DP			
<b><u>MOMETASONE FUROATE - ASMANEX TWISTHALER</u></b>						
N 021067 002	6503537	Mar 17, 2018	DP			
	8173172	Mar 17, 2018	DP			
<b><u>MOMETASONE FUROATE - ASMANEX HFA</u></b>						
N 205641 001	6068832	Aug 27, 2017	DP U-645			
<b><u>MOMETASONE FUROATE - ASMANEX HFA</u></b>						
N 205641 002	6068832	Aug 27, 2017	DP U-645			
<b><u>MOMETASONE FUROATE - SINUVA</u></b>						
N 209310 001					NP	Dec 08, 2020
<b><u>MONTELUKAST SODIUM - SINGULAIR</u></b>						
N 021409 001	8007830	Oct 24, 2022	DP			
<b><u>MORPHINE SULFATE - AVINZA</u></b>						
N 021260 001	6066339	Nov 25, 2017	DP			
<b><u>MORPHINE SULFATE - AVINZA</u></b>						
N 021260 002	6066339	Nov 25, 2017	DP			
<b><u>MORPHINE SULFATE - AVINZA</u></b>						
N 021260 003	6066339	Nov 25, 2017	DP			
<b><u>MORPHINE SULFATE - AVINZA</u></b>						
N 021260 004	6066339	Nov 25, 2017	DP			
<b><u>MORPHINE SULFATE - AVINZA</u></b>						
N 021260 005	6066339	Nov 25, 2017	DP			
<b><u>MORPHINE SULFATE - AVINZA</u></b>						
N 021260 006	6066339	Nov 25, 2017	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>MORPHINE SULFATE - MORPHINE SULFATE</b>						
N 204223 001	9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034		U-43		
	9192608	Mar 12, 2034		U-55		
	9248229	Mar 12, 2034	DP			
<b>MORPHINE SULFATE - MORPHINE SULFATE</b>						
N 204223 002	9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034		U-43		
	9192608	Mar 12, 2034		U-55		
	9248229	Mar 12, 2034	DP			
<b>MORPHINE SULFATE - MORPHINE SULFATE</b>						
N 204223 003	9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034		U-43		
	9192608	Mar 12, 2034		U-55		
	9248229	Mar 12, 2034	DP			
<b>MORPHINE SULFATE - MORPHINE SULFATE</b>						
N 204223 004	9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034		U-43		
	9192608	Mar 12, 2034		U-55		
	9248229	Mar 12, 2034	DP			
<b>MORPHINE SULFATE - MORPHINE SULFATE</b>						
N 204223 005	9072781	Mar 12, 2034	DP			
	9192608	Mar 12, 2034		U-43		
	9192608	Mar 12, 2034		U-55		
	9248229	Mar 12, 2034	DP			
<b>MORPHINE SULFATE - MORPHABOND ER</b>						
N 206544 001	7955619	Aug 12, 2028	DP		M-189	Oct 02, 2018
<b>MORPHINE SULFATE - MORPHABOND ER</b>						
N 206544 002	7955619	Aug 12, 2028	DP		M-189	Oct 02, 2018
<b>MORPHINE SULFATE - MORPHABOND ER</b>						
N 206544 003	7955619	Aug 12, 2028	DP		M-189	Oct 02, 2018
<b>MORPHINE SULFATE - MORPHABOND ER</b>						
N 206544 004	7955619	Aug 12, 2028	DP		M-189	Oct 02, 2018
<b>MORPHINE SULFATE - ARYMO ER</b>						
N 208603 001	9044402	Jul 01, 2033	DP	U-1556		
	9549899	Jul 01, 2033	DP	U-1556		
<b>MORPHINE SULFATE - ARYMO ER</b>						
N 208603 002	9044402	Jul 01, 2033	DP	U-1556		
	9549899	Jul 01, 2033	DP	U-1556		
<b>MORPHINE SULFATE - ARYMO ER</b>						
N 208603 003	9044402	Jul 01, 2033	DP	U-1556		
	9549899	Jul 01, 2033	DP	U-1556		
<b>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</b>						
N 022321 001	7682633	Jun 19, 2027		U-1510		
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
	8158156	Jun 19, 2027		U-1510		
	8623418	Nov 07, 2029		U-1640		
	8685443	Jul 03, 2025		U-1508		
	8685444	Jul 03, 2025	DP			
	8846104	Jun 19, 2027	DP			
	8877247	Jun 19, 2027	DP			
<b>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</b>						
N 022321 002	7682633	Jun 19, 2027		U-1510		
	7682634	Jun 19, 2027	DP			
	7815934	Dec 12, 2027	DP			
	8158156	Jun 19, 2027		U-1510		
	8623418	Nov 07, 2029		U-1640		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u></b>						
N 022321 002	8685443	Jul 03, 2025		U-1508		
	8685444	Jul 03, 2025		DP		
	8846104	Jun 19, 2027		DP		
	8877247	Jun 19, 2027		DP		
<b><u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u></b>						
N 022321 003	7682633	Jun 19, 2027		U-1510		
	7682634	Jun 19, 2027		DP		
	7815934	Dec 12, 2027		DP		
	8158156	Jun 19, 2027		U-1510		
	8623418	Nov 07, 2029		U-1640		
	8685443	Jul 03, 2025		U-1508		
	8685444	Jul 03, 2025		DP		
	8846104	Jun 19, 2027		DP		
	8877247	Jun 19, 2027		DP		
<b><u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u></b>						
N 022321 004	7682633	Jun 19, 2027		U-1510		
	7682634	Jun 19, 2027		DP		
	7815934	Dec 12, 2027		DP		
	8158156	Jun 19, 2027		U-1510		
	8623418	Nov 07, 2029		U-1640		
	8685443	Jul 03, 2025		U-1508		
	8685444	Jul 03, 2025		DP		
	8846104	Jun 19, 2027		DP		
	8877247	Jun 19, 2027		DP		
<b><u>MORPHINE SULFATE; NALTREXONE HYDROCHLORIDE - EMBEDA</u></b>						
N 022321 005	7682633	Jun 19, 2027		U-1510		
	7682634	Jun 19, 2027		DP		
	7815934	Dec 12, 2027		DP		
	8158156	Jun 19, 2027		U-1510		
	8623418	Nov 07, 2029		U-1640		
	8685443	Jul 03, 2025		U-1508		
	8685444	Jul 03, 2025		DP		
	8846104	Jun 19, 2027		DP		
	8877247	Jun 19, 2027		DP		
<b><u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX</u></b>						
N 021085 001	6610327	Oct 29, 2019		DP U-298		M-185 Sep 27, 2019
<b><u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u></b>						
N 021277 001	6548079	Jul 25, 2020		DP U-298		M-185 Sep 27, 2019
<b><u>MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX</u></b>						
N 021598 001	6716830	Sep 29, 2019		DP		
	7671070	Sep 29, 2019		U-709		
<b><u>MOXIFLOXACIN HYDROCHLORIDE - MOXEZA</u></b>						
N 022428 001	6716830	Sep 29, 2019		DP		
	7671070	Sep 29, 2019		DP U-709		
	8450311	May 29, 2029		DP		
	9114168	May 29, 2029		DP		
<b><u>MUPIROCIN - CENTANY</u></b>						
N 050788 001	6013657	Jul 08, 2018		DP		

**PREScription AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MYCOPHENOLIC ACID - MYFORTIC</u>						
N 050791 001	6306900	Feb 27, 2018	DP			
<u>MYCOPHENOLIC ACID - MYFORTIC</u>						
N 050791 002	6306900	Feb 27, 2018	DP			
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 019599 002					M-191	Nov 10, 2019
<u>NAFTIFINE HYDROCHLORIDE - NAFTIN</u>						
N 204286 001	8778365	Jan 31, 2033	DP			
	9161914	Jan 31, 2033	U-540			
<u>NALDEMEDINE TOSYLATE - SYMPROTC</u>						
N 208854 001	9108975	Nov 11, 2031	DS DP		NCE	Mar 23, 2022
	RE46365	Jan 11, 2028	DS DP			
	RE46375	Oct 05, 2026	DS DP U-1185			
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760 001	7056500	Jun 29, 2024	DP U-1185		NCE	Sep 16, 2019
	7662365	Oct 18, 2022	DS DP			
	7786133	Dec 19, 2027	DS DP			
	8067431	Dec 16, 2024	U-1185			
	8617530	Oct 18, 2022	U-1185			
	9012469	Apr 02, 2032	DS DP			
<u>NALOXEGOL OXALATE - MOVANTIK</u>						
N 204760 002	7056500	Jun 29, 2024	DP U-1185		NCE	Sep 16, 2019
	7662365	Oct 18, 2022	DS DP			
	7786133	Dec 19, 2027	DS DP			
	8067431	Dec 16, 2024	U-1185			
	8617530	Oct 18, 2022	U-1185			
	9012469	Apr 02, 2032	DS DP			
<u>NALOXONE HYDROCHLORIDE - EVZIO</u>						
N 205787 001	7731686	Jun 10, 2026	DP			
	7731690	Jan 15, 2025	DP			
	7749194	Oct 30, 2028	DP			
	7918823	Nov 23, 2024	DP			
	7947017	Mar 12, 2028	DP			
	8016788	Mar 21, 2025	DP			
	8021344	Nov 02, 2029	DP			
	8206360	Feb 27, 2027	DP			
	8226610	Apr 10, 2029	DP			
	8231573	Nov 25, 2028	DP			
	8313466	Nov 23, 2024	DP			
	8361029	Nov 23, 2024	DP			
	8425462	Nov 23, 2024	DP			
	8608698	Nov 23, 2024	DP			
	8627816	Feb 04, 2032	DP			
	8926594	Mar 31, 2026	DP			
	8939943	Feb 28, 2031	DP			
	9022022	Feb 28, 2031	DP			
	9056170	Nov 23, 2024	DP			
	9238108	Feb 20, 2027	DP			
	9278182	Feb 01, 2026	DP			
	9474869	Feb 28, 2031	DP U-1907			
	9517307	Jul 18, 2034	DP U-1925			
	9724471	May 23, 2027	DP U-2092			
	9737669	Nov 23, 2024	DP			
<u>NALOXONE HYDROCHLORIDE - NARCAN</u>						
N 208411 001	9211253	Mar 16, 2035	DP			
	9468747	Mar 16, 2035	DP U-1903			
	9561177	Mar 16, 2035	DP U-1903			
	9629965	Mar 16, 2035	DP U-1903			
	9775838	Mar 16, 2035	U-1903			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>NALOXONE HYDROCHLORIDE - NARCAN</u></b>						
N 208411	002	9480644	Mar 16, 2035	DP U-1903		
		9707226	Mar 16, 2035	DP U-1903		
<b><u>NALOXONE HYDROCHLORIDE - EVZIO</u></b>						
N 209862	001	7731686	Jun 01, 2026	DP		
		7731690	Jan 15, 2025	DP		
		7749194	Oct 30, 2028	DP		
		7918823	Nov 23, 2024	DP		
		7947017	Mar 12, 2028	DP		
		8016788	Mar 21, 2025	DP		
		8021344	Nov 02, 2029	DP		
		8206360	Feb 27, 2027	DP		
		8226610	Apr 10, 2029	DP		
		8231573	Nov 25, 2028	DP		
		8313466	Nov 23, 2024	DP		
		8361029	Nov 23, 2024	DP		
		8425462	Nov 23, 2024	DP		
		8608698	Nov 23, 2024	DP		
		8627816	Feb 04, 2032	DP		
		8926594	Mar 31, 2026	DP		
		8939943	Feb 28, 2031	DP		
		9022022	Feb 28, 2031	DP		
		9056170	Nov 23, 2024	DP		
		9238108	Feb 20, 2027	DP		
		9278182	Feb 01, 2026	DP		
		9474869	Feb 28, 2031	DP U-1907		
		9517307	Jul 18, 2034	DP U-1925		
		9724471	May 23, 2027	DP U-2092		
		9737669	Nov 23, 2024	DP		
<b><u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u></b>						
N 205777	001	6277384	Dec 22, 2018	DP	NC	Jul 23, 2017
		6696066	Dec 22, 2018	DP		
		8673355	Dec 22, 2018	DP		
		8822487	Dec 22, 2018	DP		
		8846090	Apr 04, 2023	DP		
		8846091	Apr 04, 2023	DP		
		8969369	May 10, 2022	DP U-1556		
		9056051	May 10, 2022	DP U-1556		
		9073933	Mar 30, 2025	DS		
		9084729	May 10, 2022	DP U-1556		
		9161937	May 10, 2022	DP U-1556		
		9168252	May 10, 2022	DP U-1556		
		9205082	Dec 22, 2018	DP U-1556		
		9283216	May 10, 2022	DP U-1819		
		9283221	May 10, 2022	DP U-1819		
		9345701	May 10, 2022	DP U-1819		
		9474750	Dec 22, 2018	DP U-1556		
		9511066	May 10, 2022	U-1921		
		9522919	Mar 30, 2025	DS DP		
		9555000	Apr 04, 2023	DP U-1556		
<b><u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u></b>						
N 205777	002	6277384	Dec 22, 2018	DP	NC	Jul 23, 2017
		6696066	Dec 22, 2018	DP		
		8673355	Dec 22, 2018	DP		
		8822487	Dec 22, 2018	DP		
		8846090	Apr 04, 2023	DP		
		8846091	Apr 04, 2023	DP		
		8969369	May 10, 2022	DP U-1556		
		9056051	May 10, 2022	DP U-1556		
		9073933	Mar 30, 2025	DS		
		9084729	May 10, 2022	DP U-1556		
		9161937	May 10, 2022	DP U-1556		
		9168252	May 10, 2022	DP U-1556		
		9205082	Dec 22, 2018	DP U-1556		
		9283216	May 10, 2022	DP U-1819		
		9283221	May 10, 2022	DP U-1819		
		9345701	May 10, 2022	DP U-1819		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u></b>						
N 205777 002	9474750	Dec 22, 2018	DP U-1556			
	9511066	May 10, 2022	U-1921			
	9522919	Mar 30, 2025	DS DP			
	9555000	Apr 04, 2023	DP U-1556			
<b><u>NALOXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TARGINIQ</u></b>						
N 205777 003	6277384	Dec 22, 2018	DP		NC	Jul 23, 2017
	6696066	Dec 22, 2018	DP			
	8673355	Dec 22, 2018	DP			
	8822487	Dec 22, 2018	DP			
	8846090	Apr 04, 2023	DP			
	8846091	Apr 04, 2023	DP			
	8969369	May 10, 2022	DP U-1556			
	9056051	May 10, 2022	DP U-1556			
	9073933	Mar 30, 2025	DS			
	9084729	May 10, 2022	DP U-1556			
	9161937	May 10, 2022	DP U-1556			
	9168252	May 10, 2022	DP U-1556			
	9205082	Dec 22, 2018	DP U-1556			
	9283216	May 10, 2022	DP U-1819			
	9283221	May 10, 2022	DP U-1819			
	9345701	May 10, 2022	DP U-1819			
	9474750	Dec 22, 2018	DP U-1556			
	9511066	May 10, 2022	U-1921			
	9522919	Mar 30, 2025	DS DP			
<b><u>NALTREXONE - VIVITROL</u></b>						
N 021897 001	6194006	Dec 30, 2018	DP			
	6264987	May 19, 2020	DP			
	6331317	Nov 12, 2019	DP			
	6379703	Dec 30, 2018	DP			
	6379704	May 19, 2020	DP			
	6395304	Nov 12, 2019	DP			
	6495164	May 25, 2020	DP			
	6495166	Nov 12, 2019	DP			
	6534092	May 19, 2020	DP			
	6537586	Nov 12, 2019	DP			
	6596316	Dec 30, 2018	DP			
	6667061	May 25, 2020	DP			
	6713090	Nov 12, 2019	DP			
	6939033	Nov 12, 2019	DP			
	7799345	May 25, 2020	DP			
	7919499	Oct 15, 2029	U-1123			
	7919499	Oct 15, 2029	U-1124			
<b><u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u></b>						
N 207621 001	7815934	Dec 12, 2027	DP		NC	Aug 19, 2019
	8685443	Jul 03, 2025	U-1508			
<b><u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u></b>						
N 207621 002	7815934	Dec 12, 2027	DP		NC	Aug 19, 2019
	8685443	Jul 03, 2025	U-1508			
<b><u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u></b>						
N 207621 003	7815934	Dec 12, 2027	DP		NC	Aug 19, 2019
	8685443	Jul 03, 2025	U-1508			
<b><u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u></b>						
N 207621 004	7815934	Dec 12, 2027	DP		NC	Aug 19, 2019
	8685443	Jul 03, 2025	U-1508			
<b><u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u></b>						
N 207621 005	7815934	Dec 12, 2027	DP		NC	Aug 19, 2019
	8685443	Jul 03, 2025	U-1508			
<b><u>NALTREXONE HYDROCHLORIDE; OXYCODONE HYDROCHLORIDE - TROXYCA ER</u></b>						
N 207621 006	7815934	Dec 12, 2027	DP		NC	Aug 19, 2019
	8685443	Jul 03, 2025	U-1508			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>NAPROXEN SODIUM - NAPROXEN SODIUM</b>						
N 021920 001	9693978	Mar 03, 2026	DP			
	9693979	Mar 03, 2026	DP			
<b>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</b>						
N 021926 001	6060499	Aug 14, 2017	DP U-867			
	6060499*PED	Feb 14, 2018				
	6586458	Aug 14, 2017	DP U-867			
	6586458*PED	Feb 14, 2018				
	7332183	Oct 02, 2025	DP U-867			
	7332183*PED	Apr 02, 2026				
	8022095	Aug 14, 2017	DP U-867			
	8022095*PED	Feb 14, 2018				
<b>NAPROXEN SODIUM; SUMATRIPTAN SUCCINATE - TREXIMET</b>						
N 021926 002	5872145	Aug 14, 2017	DP U-1719		NP	May 14, 2018
	5872145*PED	Feb 14, 2018			PED	Nov 14, 2018
	6060499	Aug 14, 2017	DP U-1719			
	6060499*PED	Feb 14, 2018				
	6586458	Aug 14, 2017	DP U-1719			
	6586458*PED	Feb 14, 2018				
	7332183	Oct 02, 2025	DP U-1719			
	7332183*PED	Apr 02, 2026				
<b>NATEGLINIDE - STARLIX</b>						
N 021204 001	6559188	Sep 15, 2020	DP U-827			
	6641841	Nov 14, 2017	DP U-214			
	6844008	Nov 14, 2017	DP U-214			
	6878749	Sep 15, 2020	DP			
<b>NATEGLINIDE - STARLIX</b>						
N 021204 002	6559188	Sep 15, 2020	DP U-827			
	6641841	Nov 14, 2017	DP U-214			
	6844008	Nov 14, 2017	DP U-214			
	6878749	Sep 15, 2020	DP			
<b>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</b>						
N 021742 002	6545040	Dec 17, 2021	DP U-3			
<b>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</b>						
N 021742 003	6545040	Dec 17, 2021	DP U-3			
<b>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</b>						
N 021742 004	6545040	Dec 17, 2021	DP U-3			
<b>NEBIVOLOL HYDROCHLORIDE - BYSTOLIC</b>						
N 021742 005	6545040	Dec 17, 2021	DP U-3			
<b>NEBIVOLOL HYDROCHLORIDE; VALSARTAN - BYVALSON</b>						
N 206302 001	7803838	Aug 29, 2026	DP		NC	Jun 03, 2019
	7838552	Oct 04, 2027	U-185			
<b>NEPafenac - NEVANAC</b>						
N 021862 001	7834059	Jan 31, 2027	U-1095			
	8071648	Dec 02, 2025	DP			
	8324281	Dec 02, 2025	DP			
<b>NEPafenac - ILEVRO</b>						
N 203491 001	6403609	Jul 17, 2018	DP			
	7947295	Jun 08, 2024	DP			
	8921337	Mar 31, 2032	DP			
	9662398	Dec 01, 2030	DP			
<b>NERATINIB MALEATE - NERLYNX</b>						
N 208051 001	6288082	Sep 24, 2019	DS DP U-2043		NCE	Jul 17, 2022
	7399865	Dec 29, 2025	DS DP			
	7982043	Oct 08, 2025	U-2043			
	8518446	Nov 20, 2030	DP U-2043			
	8790708	Nov 05, 2030	DP U-2043			
	9139558	Oct 15, 2028	U-2043			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>NERATINIB MALEATE - NERLYNX</u></b>						
N 208051 001	9211291	Mar 24, 2030	U-2043			
	9630946	Oct 15, 2028	U-2043			
<b><u>NETARSUDIL DIMESYLATE - RHOPRESA</u></b>						
N 208254 001	8394826	Nov 10, 2030	DS DP U-1524		NCE	Dec 18, 2022
	8450344	Jul 11, 2026	DS DP U-1524			
	9096569	Jul 11, 2026	DS DP U-1524			
	9415043	Mar 14, 2034	DS			
<b><u>NETUPITANT; PALONOSETRON HYDROCHLORIDE - AKYNZEO</u></b>						
N 205718 001	6297375	Feb 22, 2020	DS		NCE	Oct 10, 2019
	8623826	Nov 18, 2030	U-528			
	8951969	Nov 18, 2030	DP			
	9186357	Nov 18, 2030	U-528			
	9271975	Sep 09, 2031	U-528			
<b><u>NEVIRAPINE - VIRAMUNE XR</u></b>						
N 201152 001	8460704	Mar 12, 2029	U-1409			
<b><u>NIACIN - NIASPAN</u></b>						
N 020381 002	6469035	Mar 15, 2018	U-1142			
	6469035	Mar 15, 2018	U-1143			
	6469035	Mar 15, 2018	U-1144			
	6469035	Mar 15, 2018	U-1145			
	6469035	Mar 15, 2018	U-768			
<b><u>NIACIN - NIASPAN</u></b>						
N 020381 003	6469035	Mar 15, 2018	U-1142			
	6469035	Mar 15, 2018	U-1143			
	6469035	Mar 15, 2018	U-1144			
	6469035	Mar 15, 2018	U-1145			
	6469035	Mar 15, 2018	U-768			
<b><u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u></b>						
N 019734 002	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027	U-1029			
	8455524	Apr 18, 2027	U-1029			
	9364564	Dec 26, 2027	DP			
<b><u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u></b>						
N 019734 003	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027	U-1029			
	8455524	Apr 18, 2027	U-1029			
	9364564	Dec 26, 2027	DP			
<b><u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u></b>						
N 019734 004	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027	U-1029			
	8455524	Apr 18, 2027	U-1029			
	9364564	Dec 26, 2027	DP			
<b><u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER</u></b>						
N 019734 005	7612102	Dec 26, 2027	DP			
	7659291	Apr 18, 2027	U-1029			
	8455524	Apr 18, 2027	U-1029			
	9364564	Dec 26, 2027	DP			
<b><u>NICOTINE - NICODERM CQ</u></b>						
N 020165 004	8075911	May 22, 2021	DP			
	8663680	Feb 13, 2020	DP			
	8999379	Feb 13, 2020	U-1686			
	9205059	Dec 15, 2019	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>NICOTINE - NICODERM CQ</u></b>						
N 020165 005	8075911	May 22, 2021	DP			
	8663680	Feb 13, 2020	DP			
	8999379	Feb 13, 2020		U-1686		
	9205059	Dec 15, 2019	DP			
<b><u>NICOTINE - NICODERM CQ</u></b>						
N 020165 006	8075911	May 22, 2021	DP			
	8663680	Feb 13, 2020	DP			
	8999379	Feb 13, 2020		U-1686		
	9205059	Dec 15, 2019	DP			
<b><u>NICOTINE POLACRILEX - NICORETTE</u></b>						
N 018612 002	8323683	Apr 30, 2028				
<b><u>NICOTINE POLACRILEX - NICORETTE</u></b>						
N 020066 002	8323683	Apr 30, 2028	DP			
<b><u>NICOTINE POLACRILEX - NICORETTE</u></b>						
N 022360 001	8501164	Jun 14, 2029	DP			
	8940772	Apr 30, 2029	DP			
<b><u>NICOTINE POLACRILEX - NICORETTE</u></b>						
N 022360 002	8501164	Jun 14, 2029	DP			
	8940772	Apr 30, 2029	DP			
<b><u>NILOTINIB HYDROCHLORIDE MONOHYDRATE - TASIGNA</u></b>						
N 022068 001	7169791	Jul 04, 2023	DS DP	U-836		
	8163904	Aug 23, 2028	DS DP			
	8293756	Sep 25, 2027	DP			
	8389537	Jul 18, 2026	DS DP	U-1374		
	8415363	Jul 18, 2026	DS DP	U-1407		
	8501760	Jul 18, 2026	DS DP			
	9061029	Apr 07, 2032	DP	U-1374		
<b><u>NILOTINIB HYDROCHLORIDE MONOHYDRATE - TASIGNA</u></b>						
N 022068 002	7169791	Jul 04, 2023	DS DP	U-836		
	8163904	Aug 23, 2028	DS DP			
	8293756	Sep 25, 2027	DP			
	8389537	Jul 18, 2026	DS DP	U-1374		
	8415363	Jul 18, 2026	DS DP	U-1407		
	8501760	Jul 18, 2026	DS DP			
	9061029	Apr 07, 2032	DP	U-1374		
<b><u>NIMODIPINE - NYMALIZE</u></b>						
N 203340 001					ODE-46	May 10, 2020
<b><u>NINTEDANIB ESYLATE - OFEV</u></b>						
N 205832 001	6762180	Dec 10, 2020	DS DP		NCE	Oct 15, 2019
	7119093	Feb 21, 2024	DS DP		ODE-77	Oct 15, 2021
	7989474	Apr 06, 2024		U-1677		
<b><u>NINTEDANIB ESYLATE - OFEV</u></b>						
N 205832 002	6762180	Dec 10, 2020	DS DP		NCE	Oct 15, 2019
	7119093	Feb 21, 2024	DS DP		ODE-77	Oct 15, 2021
	7989474	Apr 06, 2024		U-1677		
<b><u>NIRAPARIB TOSYLATE - ZEJULA</u></b>						
N 208447 001	8071623	Mar 22, 2030	DS DP		NCE	Mar 27, 2022
	8436185	Apr 24, 2029	DS		ODE-133	Mar 27, 2024
<b><u>NITAZOXANIDE - ALINIA</u></b>						
N 021498 001	5965590	Jul 03, 2017		U-523		
<b><u>NITISINONE - ORFADIN</u></b>						
N 021232 001					D-169	Sep 01, 2020
<b><u>NITISINONE - ORFADIN</u></b>						
N 021232 002					D-169	Sep 01, 2020

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NITISINONE - ORFADIN</u>					D-169	Sep 01, 2020
N 021232 003						
<u>NITISINONE - ORFADIN</u>					D-169	Sep 01, 2020
N 021232 004						
<u>NITISINONE - ORFADIN</u>					D-169	Sep 01, 2020
N 206356 001 9301932		Feb 28, 2033	DP U-1836			
<u>NITRIC OXIDE - INOMAX</u>						
N 020845 002 6125846*PED		Nov 16, 2017				
8282966		Jun 30, 2029	U-1286			
8291904		Jan 06, 2031	DP U-1226			
8293284		Jun 30, 2029	U-1286			
8431163		Jun 30, 2029	U-1286			
8431163*PED		Dec 30, 2029				
8573209		Jan 06, 2031	DP			
8573209*PED		Jul 06, 2031				
8573210		Jan 06, 2031	DP U-1453			
8573210*PED		Jul 06, 2031				
8776794		Jan 06, 2031	DP U-1226			
8776794*PED		Jul 06, 2031				
8776795		Jan 06, 2031	DP U-1226			
8776795*PED		Jul 06, 2031				
8795741		Jun 30, 2029	U-1286			
8795741*PED		Dec 30, 2029				
8846112		Jun 30, 2029	U-1286			
8846112*PED		Dec 30, 2029				
<u>NITRIC OXIDE - INOMAX</u>						
N 020845 003 6125846*PED		Nov 16, 2017			M-167	Oct 09, 2018
8282966		Jun 30, 2029	U-1286			
8291904		Jan 06, 2031	DP U-1226			
8293284		Jun 30, 2029	U-1286			
8431163		Jun 30, 2029	U-1286			
8431163*PED		Dec 30, 2029				
8573209		Jan 06, 2031	DP			
8573209*PED		Jul 06, 2031				
8573210		Jan 06, 2031	DP U-1453			
8573210*PED		Jul 06, 2031				
8776794		Jan 06, 2031	DP U-1226			
8776794*PED		Jul 06, 2031				
8776795		Jan 06, 2031	DP U-1226			
8776795*PED		Jul 06, 2031				
8795741		Jun 30, 2029	U-1286			
8795741*PED		Dec 30, 2029				
8846112		Jun 30, 2029	U-1286			
8846112*PED		Dec 30, 2029				
9265911		Jan 06, 2031	DP U-1824			
9265911*PED		Jul 06, 2031				
9279794		Feb 19, 2034	DP U-1823			
9279794*PED		Aug 19, 2034				
9295802		Jan 06, 2031	DP U-1226			
9295802*PED		Jul 06, 2031				
9408993		Jan 06, 2031	DP U-1824			
9408993*PED		Jul 06, 2031				
9770570		May 03, 2036	U-2148			
9770570*PED		Nov 03, 2036				
<u>NITROGLYCERIN - NITROLINGUAL PUMPS SPRAY</u>						
N 018705 002 7872049		Mar 14, 2028	DP U-39			
<u>NITROGLYCERIN - NITROSTAT</u>						
N 021134 001 6500456		Sep 16, 2018				
<u>NITROGLYCERIN - NITROSTAT</u>						
N 021134 002 6500456		Sep 16, 2018				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>NITROGLYCERIN - NITROSTAT</u></b>						
N 021134 003	6500456	Sep 16, 2018				
<b><u>NITROGLYCERIN - GONITRO</u></b>						
N 208424 001	9101592	Mar 11, 2032	DP			
<b><u>NIZATIDINE - AXID</u></b>						
N 021494 001	6930119	Jul 17, 2022	DP			
<b><u>NUSINERSEN SODIUM - SPINRAZA</u></b>						
N 209531 001	6166197	Dec 26, 2017	DS		NCE	Dec 23, 2021
	6210892	Oct 07, 2018		U-1942	ODE-127	Dec 23, 2023
	7101993	Sep 05, 2023	DS			
	7838657	Jul 11, 2027	DS			
	8110560	Dec 05, 2025		U-1942		
	8110560	Dec 05, 2025		U-1943		
	8110560	Dec 05, 2025		U-1944		
	8361977	May 27, 2030	DS DP			
	8980853	Nov 24, 2030		U-1941		
	9717750	Jun 17, 2030		U-1942		
	9717750	Jun 17, 2030		U-1943		
	9717750	Jun 17, 2030		U-2093		
	9717750	Jun 17, 2030		U-2094		
<b><u>OBETICHOOLIC ACID - OCALIVA</u></b>						
N 207999 001	7138390	Nov 16, 2022	DS DP		NCE	May 27, 2021
	8058267	Feb 21, 2022		U-1854	ODE-119	May 27, 2023
	8377916	Feb 21, 2022		U-1854		
	9238673	Jun 17, 2033	DP			
<b><u>OBETICHOOLIC ACID - OCALIVA</u></b>						
N 207999 002	7138390	Nov 16, 2022	DS DP		NCE	May 27, 2021
	8058267	Feb 21, 2022		U-1854	ODE-119	May 27, 2023
	8377916	Feb 21, 2022		U-1854		
	9238673	Jun 17, 2033	DP			
<b><u>OLANZAPINE - ZYPREXA</u></b>						
N 020592 001	6960577	Nov 01, 2017		U-963		
<b><u>OLANZAPINE - ZYPREXA</u></b>						
N 020592 002	6960577	Nov 01, 2017		U-963		
<b><u>OLANZAPINE - ZYPREXA</u></b>						
N 020592 003	6960577	Nov 01, 2017		U-963		
<b><u>OLANZAPINE - ZYPREXA</u></b>						
N 020592 004	6960577	Nov 01, 2017		U-963		
<b><u>OLANZAPINE - ZYPREXA</u></b>						
N 020592 005	6960577	Nov 01, 2017		U-963		
<b><u>OLANZAPINE - ZYPREXA</u></b>						
N 020592 006	6960577	Nov 01, 2017		U-963		
<b><u>OLANZAPINE - ZYPREXA ZYDIS</u></b>						
N 021086 001	6960577	Nov 01, 2017		U-964		
<b><u>OLANZAPINE - ZYPREXA ZYDIS</u></b>						
N 021086 002	6960577	Nov 01, 2017		U-964		
<b><u>OLANZAPINE - ZYPREXA ZYDIS</u></b>						
N 021086 003	6960577	Nov 01, 2017		U-964		
<b><u>OLANZAPINE - ZYPREXA ZYDIS</u></b>						
N 021086 004	6960577	Nov 01, 2017		U-964		
<b><u>OLANZAPINE PAMOATE - ZYPREXA RELPREVV</u></b>						
N 022173 001	6169084	Sep 30, 2018	DS DP	U-1026		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OLANZAPINE PAMOATE - ZYPREXA RELPREVV</u></b>						
N 022173 002	6169084	Sep 30, 2018	DS DP U-1026			
<b><u>OLANZAPINE PAMOATE - ZYPREXA RELPREVV</u></b>						
N 022173 003	6169084	Sep 30, 2018	DS DP U-1026			
<b><u>OLAPARIB - LYNPARZA</u></b>						
N 206162 001	7151102	Apr 29, 2022	DS DP		NCE	Dec 19, 2019
	7449464	Oct 11, 2024	DS DP		ODE-83	Dec 19, 2021
	7981889	Oct 11, 2024	DS DP			
	8143241	Aug 12, 2027		U-1634		
	8247416	Sep 24, 2028	DS			
	8859562	Aug 04, 2031		U-1634		
	8912187	Mar 12, 2024		U-1634		
<b><u>OLAPARIB - LYNPARZA</u></b>						
N 208558 001	7151102	Apr 29, 2022	DS DP		NCE	Dec 19, 2019
	7449464	Oct 11, 2024	DS DP		NP	Aug 17, 2020
	7981889	Oct 11, 2024	DS DP		ODE-83	Dec 19, 2021
	8143241	Aug 12, 2027		U-2101		
	8143241	Aug 12, 2027		U-2102		
	8143241	Aug 12, 2027		U-2103		
	8475842	Dec 31, 2029	DP			
	8859562	Aug 04, 2031		U-2101		
	8859562	Aug 04, 2031		U-2102		
	8912187	Mar 12, 2024		U-2101		
	8912187	Mar 12, 2024		U-2102		
<b><u>OLODATEROL HYDROCHLORIDE - STRIVERDI RESPIMAT</u></b>						
N 203108 001	6846413	Aug 28, 2018	DP		NCE	Jul 31, 2019
	6977042	Aug 28, 2018	DP			
	6988496	Feb 23, 2020	DP	U-1547		
	7056916	Dec 07, 2023	DS	DP		
	7220742	May 12, 2025	DS	DP U-1547		
	7284474	Aug 26, 2024		DP		
	7396341	Oct 10, 2026		DP	U-1547	
	7491719	Nov 10, 2023	DS	DP		
	7727984	Nov 10, 2023	DS			
	7786111	Nov 10, 2023		DP		
	7802568	Feb 26, 2019		DP		
	7837235	Mar 13, 2028		DP		
	7896264	May 26, 2025		DP		
	7988001	Aug 04, 2021		DP		
	8034809	May 12, 2025		U-1547		
	8044046	Nov 10, 2023		U-1547		
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
<b><u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u></b>						
N 206756 001	6846413	Aug 28, 2018	DP		M-173	Mar 18, 2019
	6846413*PED	Feb 28, 2019			NC	May 21, 2018
	6977042	Aug 28, 2018	DP		NCE	Jul 31, 2019
	6977042*PED	Feb 28, 2019				
	6988496	Feb 23, 2020	DP			
	6988496*PED	Aug 23, 2020				
	7056916	Dec 07, 2023	DS	DP		
	7220742	May 12, 2025	DS	DP U-1703		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OLODATEROL HYDROCHLORIDE; TIOTROPIUM BROMIDE - STIOLTO RESPIMAT</u></b>						
N 206756 001	7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7491719	Nov 10, 2023	DS DP			
	7727984	Nov 10, 2023	DS			
	7786111	Nov 10, 2023	DP			
	7802568	Feb 26, 2019	DP			
	7802568*PED	Aug 26, 2019				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7988001	Aug 04, 2021	DP			
	8034809	May 12, 2025	U-1702			
	8044046	Nov 10, 2023	U-1702			
	8733341	Oct 16, 2030	DP			
	9027967	Mar 31, 2027	DP			
	RE39820	Jan 30, 2018	DS DP U-1702			
<b><u>OLOPATADINE HYDROCHLORIDE - OLOPATADINE HYDROCHLORIDE</u></b>						
A 090848 001					PC	Dec 05, 2017
<b><u>OLOPATADINE HYDROCHLORIDE - PATADAY</u></b>						
N 021545 001	6995186	Nov 12, 2023	DP U-765			
	7402609	Jun 19, 2022	DP			
<b><u>OLOPATADINE HYDROCHLORIDE - PATANASE</u></b>						
N 021861 001	7977376	Feb 02, 2023	DP			
	8399508	Sep 17, 2022	U-726			
	8399508*PED	Mar 17, 2023				
<b><u>OLOPATADINE HYDROCHLORIDE - PAZEO</u></b>						
N 206276 001	8791154	May 19, 2032	DP U-1680		NP	Jan 30, 2018
	9533053	May 19, 2032	DP		PED	Jul 30, 2018
<b><u>OMACETAXINE MEPESUCCINATE - SYNRIBO</u></b>						
N 203585 001	6987103	Oct 26, 2026	U-1300		NCE	Oct 26, 2017
	RE45128	Mar 16, 2019	DS DP U-1576		ODE-32	Oct 26, 2019
<b><u>OMBITASVIR; PARITAPREVIR; RITONAVIR - TECHNIVIE</u></b>						
N 207931 001	7148359	Jul 19, 2019	DP		I-743	Jul 24, 2018
	7148359*PED	Jan 19, 2020			NCE	Dec 19, 2019
	7364752	Nov 10, 2020	DP		NP	Jul 24, 2018
	7364752*PED	May 10, 2021				
	8268349	Aug 25, 2024	DP			
	8268349*PED	Feb 25, 2025				
	8399015	Aug 25, 2024	DP			
	8399015*PED	Feb 25, 2025				
	8420596	Apr 10, 2031	DS DP			
	8420596*PED	Oct 10, 2031				
	8642538	Sep 10, 2029	DS DP U-1638			
	8686026	Jun 09, 2031	DP			
	8691938	Apr 13, 2032	DS DP			
	9006387	Jun 10, 2030	U-1687			
	9044480	Apr 10, 2031	U-1638			
<b><u>OMEGA-3-ACID ETHYL ESTERS TYPE A - OMTRYG</u></b>						
N 204977 001					NCE	Apr 23, 2019
<b><u>OMEGA-3-CARBOXYLIC ACIDS - EPANOVA</u></b>						
N 205060 001	5792795	May 13, 2018	DP		NCE	May 05, 2019
	5948818	May 13, 2018	DP			
	7960370	Feb 07, 2025	DP			
	8383678	Feb 07, 2025	DP U-1511			
	9012501	Feb 07, 2025	DP U-1511			
	9050308	Jan 04, 2033	U-1511			
	9050309	Jan 04, 2033	DS			
	9132112	Feb 07, 2025	DP U-1511			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OMEPRAZOLE - PRILOSEC</u></b>						
N 019810 001	6147103	Oct 09, 2018				
	6150380	Nov 10, 2018				
	6166213	Oct 09, 2018				
	6191148	Oct 09, 2018				
<b><u>OMEPRAZOLE - PRILOSEC</u></b>						
N 019810 002	6147103	Oct 09, 2018				
	6150380	Nov 10, 2018				
	6166213	Oct 09, 2018				
	6191148	Oct 09, 2018				
<b><u>OMEPRAZOLE - PRILOSEC</u></b>						
N 019810 003	6147103	Oct 09, 2018				
	6150380	Nov 10, 2018				
	6166213	Oct 09, 2018				
	6191148	Oct 09, 2018				
<b><u>OMEPRAZOLE - OMEPRAZOLE</u></b>						
N 022032 001	9023391	Aug 16, 2025	DP			
<b><u>OMEPRAZOLE MAGNESIUM - OMEPRAZOLE MAGNESIUM</u></b>						
A 204152 001				PC		May 30, 2018
<b><u>OMEPRAZOLE MAGNESIUM - PRILOSEC OTC</u></b>						
N 021229 001	6403616	Nov 15, 2019				
	6428810	Nov 03, 2019				
<b><u>OMEPRAZOLE MAGNESIUM - PRILOSEC</u></b>						
N 022056 001	6428810	Nov 03, 2019	DP U-1817			
	6428810	Nov 03, 2019	DP U-864			
<b><u>OMEPRAZOLE MAGNESIUM - PRILOSEC</u></b>						
N 022056 002	6428810	Nov 03, 2019	DP U-864			
<b><u>ONDANSETRON - ZUPLENZ</u></b>						
N 022524 001	8580830	Nov 23, 2029	DP			
	9095577	Jul 13, 2030	DP			
<b><u>ONDANSETRON - ZUPLENZ</u></b>						
N 022524 002	8580830	Nov 23, 2029	DP			
	9095577	Jul 13, 2030	DP			
<b><u>ORITAVANCIN DIPHOSPHATE - ORBACTIV</u></b>						
N 206334 001	5840684	Nov 24, 2017	DS DP U-1569			
	5998581	Nov 12, 2017	DS		NCE	Aug 06, 2019
	8420592	Aug 29, 2029		U-1570	GAIN	Aug 06, 2024
	9649352	Jul 16, 2035	DP			
	9682061	Apr 26, 2030		U-1569		
<b><u>ORLISTAT - XENICAL</u></b>						
N 020766 001	6004996	Jan 06, 2018				
<b><u>ORLISTAT - ALLI</u></b>						
N 021887 001	6004996	Jan 06, 2018	DP			
<b><u>OSIMERTINIB MESYLATE - TAGRISSO</u></b>						
N 208065 001	8946235	Aug 08, 2032	DS DP U-1777		NCE	Nov 13, 2020
	9732058	Jul 25, 2032	DS DP U-1777		ODE-102	Nov 13, 2022
<b><u>OSIMERTINIB MESYLATE - TAGRISSO</u></b>						
N 208065 002	8946235	Aug 08, 2032	DS DP U-1777		NCE	Nov 13, 2020
	9732058	Jul 25, 2032	DS DP U-1777		ODE-102	Nov 13, 2022
<b><u>OSPEMIFENE - OSPHENA</u></b>						
N 203505 001	6245819	Jul 21, 2025		U-1370	NCE	Feb 26, 2018
	8236861	Aug 11, 2026		U-1369		
	8236861	Aug 11, 2026		U-1370		
	8470890	Feb 13, 2024		U-1369		
	8470890	Feb 13, 2024		U-1370		
	8642079	Jul 09, 2028	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OSPEMIFENE - OSPHENA</u></b>						
N 203505 001	8772353	Feb 13, 2024	U-1369			
	8772353	Feb 13, 2024	U-1370			
	9241915	Feb 13, 2024	U-1369			
	9241915	Feb 13, 2024	U-1370			
	9566252	Jul 21, 2020	U-1370			
<b><u>OXANDROLONE - OXANDRIN</u></b>						
N 013718 001	5872147	Dec 05, 2017	U-585			
	6090799	Jul 18, 2017	U-585			
	6576659	Dec 05, 2017	U-585			
	6828313	Dec 05, 2017	U-585			
<b><u>OXANDROLONE - OXANDRIN</u></b>						
N 013718 002	5872147	Dec 05, 2017	U-585			
	6090799	Jul 18, 2017	U-585			
	6576659	Dec 05, 2017	U-585			
	6828313	Dec 05, 2017	U-585			
<b><u>OXCARBAZEPINE - TRILEPTAL</u></b>						
N 021014 001	7037525	Feb 12, 2018	U-724			
<b><u>OXCARBAZEPINE - TRILEPTAL</u></b>						
N 021014 002	7037525	Feb 12, 2018	U-724			
<b><u>OXCARBAZEPINE - TRILEPTAL</u></b>						
N 021014 003	7037525	Feb 12, 2018	U-724			
<b><u>OXCARBAZEPINE - TRILEPTAL</u></b>						
N 021285 001	7037525	Feb 12, 2018	U-724			
	8119148	Dec 19, 2020	DP U-724			
<b><u>OXCARBAZEPINE - OXTELLAR XR</u></b>						
N 202810 001	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027	U-1298			
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027	U-1298			
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<b><u>OXCARBAZEPINE - OXTELLAR XR</u></b>						
N 202810 002	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027	U-1298			
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027	U-1298			
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<b><u>OXCARBAZEPINE - OXTELLAR XR</u></b>						
N 202810 003	7722898	Apr 13, 2027	DP			
	7910131	Apr 13, 2027	U-1298			
	8617600	Apr 13, 2027	DP			
	8821930	Apr 13, 2027	DP			
	9119791	Apr 13, 2027	U-1298			
	9351975	Apr 13, 2027	DP			
	9370525	Apr 13, 2027	DP			
	9855278	Apr 13, 2027	DP			
<b><u>OXYBUTYNIN - OXYTROL</u></b>						
N 021351 002	6743441	Apr 26, 2020	DP U-318			
	7081249	Apr 26, 2020	DP U-318			
	7081250	Apr 26, 2020	DP U-318			
	7081251	Apr 26, 2020	DP U-318			
	7081252	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	DS DP U-318			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OXYBUTYNIN - OXYTROL FOR WOMEN</u></b>						
N 202211 001	6743441	Apr 26, 2020	DP U-1329			
	7081249	Apr 26, 2020	DP U-1329			
	7081250	Apr 26, 2020	DP U-1329			
	7081251	Apr 26, 2020	DP U-1329			
	7081252	Apr 26, 2020	DP U-1329			
	7179483	Apr 26, 2020	U-1329			
<b><u>OXYBUTYNIN - GELNIQUE 3%</u></b>						
N 202513 001	7029694	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	U-318			
	7198801	Jun 25, 2022	DP			
	8241662	Apr 26, 2020	U-318			
<b><u>OXYBUTYNIN CHLORIDE - GELNIQUE</u></b>						
N 022204 001	7029694	Apr 26, 2020	DP U-318			
	7179483	Apr 26, 2020	U-318			
	8241662	Apr 26, 2020	U-318			
	8920392	Mar 26, 2031	U-1644			
	9259388	Nov 06, 2029	U-1644			
<b><u>OXYCODONE - XTAMPZA ER</u></b>						
N 208090 001	7399488	Mar 24, 2025	DP		NP	Apr 26, 2019
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
<b><u>OXYCODONE - XTAMPZA ER</u></b>						
N 208090 002	7399488	Mar 24, 2025	DP		NP	Apr 26, 2019
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
<b><u>OXYCODONE - XTAMPZA ER</u></b>						
N 208090 003	7399488	Mar 24, 2025	DP		NP	Apr 26, 2019
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	8840928	Jul 07, 2023	DP U-1556			
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023	U-1556			
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP U-1556			
	9737530	Sep 02, 2036	DP U-1556			
	9763883	Jul 07, 2023	DP			
<b><u>OXYCODONE - XTAMPZA ER</u></b>						
N 208090 004	7399488	Mar 24, 2025	DP		NP	Apr 26, 2019
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025	U-1556			
	8840928	Jul 07, 2023	DP U-1556			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OXYCODONE - XTAMPZA ER</u></b>						
N 208090 004	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023		U-1556		
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP	U-1556		
	9737530	Sep 02, 2036	DP	U-1556		
	9763883	Jul 07, 2023	DP			
<b><u>OXYCODONE - XTAMPZA ER</u></b>						
N 208090 005	7399488	Mar 24, 2025	DP		NP	Apr 26, 2019
	7771707	Mar 24, 2025	DP			
	8449909	Mar 24, 2025	DP			
	8557291	Mar 21, 2025	DP			
	8758813	Jun 10, 2025		U-1556		
	8840928	Jul 07, 2023	DP	U-1556		
	9044398	Jul 07, 2023	DP			
	9248195	Jul 07, 2023		U-1556		
	9592200	Jul 07, 2023	DP			
	9682075	Dec 10, 2030	DP	U-1556		
	9737530	Sep 02, 2036	DP	U-1556		
	9763883	Jul 07, 2023	DP			
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 001	7674799	Mar 30, 2025	DP		Y	NPP
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	DP	U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9060976	Aug 06, 2022	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS	DP		
	9675610	Jun 16, 2023	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 002	7674799	Mar 30, 2025	DP		Y	NPP
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	DP	U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894987	Mar 29, 2030	DP			
	8894988	Aug 24, 2027	DP			
	9060976	Aug 06, 2022	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS	DP		
	9675610	Jun 16, 2023	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 003	7674799	Mar 30, 2025	DP		Y	NPP
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	DP	U-1556		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 003	8808741	Aug 24, 2027		U-1556		
	8894987	Mar 29, 2030		DP		
	8894988	Aug 24, 2027		DP		
	9060976	Aug 06, 2022		DP		
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027		DP		
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027		DP		
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9675610	Jun 16, 2023		DP		
	9763933	Aug 24, 2027		DP		
	9770416	Aug 24, 2027		DP		
	9775808	Aug 24, 2027		DP		
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 004	8309060	Nov 20, 2023	DP	U-1556	NPP	
	8808741	Aug 24, 2027		U-1556		Aug 13, 2018
	8894987	Mar 29, 2030		DP		
	8894988	Aug 24, 2027		DP		
	9060976	Aug 06, 2022		DP		
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027		DP		
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027		DP		
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9675610	Jun 16, 2023		DP		
	9763933	Aug 24, 2027		DP		
	9770416	Aug 24, 2027		DP		
	9775808	Aug 24, 2027		DP		
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 005	7674799	Mar 30, 2025	DP		Y	
	7674800	Mar 30, 2025	DS		Y	
	7683072	Mar 30, 2025	DS		Y	
	7776314	Apr 19, 2025	DP		Y	
	8309060	Nov 20, 2023	DP	U-1556		
	8808741	Aug 24, 2027		U-1556		
	8894988	Aug 24, 2027		DP		
	9060976	Aug 06, 2022		DP		
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027		DP		
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027		DP		
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9675610	Jun 16, 2023		DP		
	9763933	Aug 24, 2027		DP		
	9770416	Aug 24, 2027		DP		
	9775808	Aug 24, 2027		DP		
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 006	8309060	Nov 20, 2023	DP	U-1556	NPP	
	8808741	Aug 24, 2027		U-1556		Aug 13, 2018
	8894988	Aug 24, 2027		DP		
	9060976	Aug 06, 2022		DP		
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027		DP		
	9492391	Aug 24, 2027		U-1556		
	9492392	Aug 24, 2027		DP		
	9492393	Aug 24, 2027		U-1556		
	9522919	Mar 30, 2025	DS DP			
	9675610	Jun 16, 2023		DP		
	9763933	Aug 24, 2027		DP		
	9770416	Aug 24, 2027		DP		
	9775808	Aug 24, 2027		DP		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OXYCODONE HYDROCHLORIDE - OXYCONTIN</u></b>						
N 022272 007	8309060	Nov 20, 2023	DP U-1556		NPP	Aug 13, 2018
	8808741	Aug 24, 2027	U-1556			
	8894988	Aug 24, 2027	DP			
	9060976	Aug 06, 2022	DP			
	9073933	Mar 30, 2025	DS			
	9492389	Aug 24, 2027	DP			
	9492391	Aug 24, 2027	U-1556			
	9492392	Aug 24, 2027	DP			
	9492393	Aug 24, 2027	U-1556			
	9522919	Mar 30, 2025	DS DP			
	9675610	Jun 16, 2023	DP			
	9763933	Aug 24, 2027	DP			
	9770416	Aug 24, 2027	DP			
	9775808	Aug 24, 2027	DP			
<b><u>OXYCODONE HYDROCHLORIDE - OXAYDO</u></b>						
N 202080 001	7201920	Mar 16, 2025	DP			
	7510726	Nov 26, 2023	DP			
	7981439	Nov 26, 2023	DP			
	8409616	Nov 26, 2023	DP			
	8637540	Nov 26, 2023	DP			
	9492443	May 26, 2024	DP			
<b><u>OXYCODONE HYDROCHLORIDE - OXAYDO</u></b>						
N 202080 002	7201920	Mar 16, 2025	DP			
	7510726	Nov 26, 2023	DP			
	7981439	Nov 26, 2023	DP			
	8409616	Nov 26, 2023	DP			
	8637540	Nov 26, 2023	DP			
	9492443	May 26, 2024	DP			
<b><u>OXYCODONE HYDROCHLORIDE - ROXBOND</u></b>						
N 209777 001	7955619	Aug 12, 2028	DP			
<b><u>OXYCODONE HYDROCHLORIDE - ROXBOND</u></b>						
N 209777 002	7955619	Aug 12, 2028	DP			
<b><u>OXYCODONE HYDROCHLORIDE - ROXBOND</u></b>						
N 209777 003	7955619	Aug 12, 2028	DP			
<b><u>OXYMETAZOLINE HYDROCHLORIDE - RHOFADE</u></b>						
N 208552 001	7812049	May 02, 2028	U-1959		NP	Jan 18, 2020
	8420688	Aug 02, 2024	U-1959			
	8883838	Dec 01, 2031	DP			
<b><u>OXYMETAZOLINE HYDROCHLORIDE; TETRACAINE HYDROCHLORIDE - KOVANAZE</u></b>						
N 208032 001	6413499	Mar 20, 2020	U-1876		NC	Jun 29, 2019
	8580282	Apr 02, 2030	DP U-1876			
	9308191	Apr 02, 2030	DP U-1876			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 001	7276250	Feb 04, 2023	DP U-826			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 002	7276250	Feb 04, 2023	DP U-826			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 003	7276250	Feb 04, 2023	DP U-826			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 004	7276250	Feb 04, 2023	DP U-826			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 005	7276250	Feb 04, 2023	DP U-826			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 006	7276250	Feb 04, 2023	DP U-826			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 021610 007	7276250	Feb 04, 2023	DP	U-826		
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 001	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 002	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 003	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 004	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 005	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 006	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027		U-1598		
	8871779	Nov 22, 2029	DS			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u></b>						
N 201655 007	7851482	Jul 10, 2029	DS			
	8075872	Nov 20, 2023	DP			
	8114383	Aug 08, 2024	DP			
	8192722	Sep 15, 2025	DP			
	8309060	Nov 20, 2023	DP			
	8309122	Feb 04, 2023	DP			
	8329216	Feb 04, 2023	DP			
	8808737	Jun 21, 2027	U-1598			
	8871779	Nov 22, 2029	DS			
<b><u>OZENOXACIN - XEPI</u></b>						
N 208945 001					NCE	Dec 11, 2022
<b><u>PACLITAXEL - ABRAXANE</u></b>						
N 021660 001	7758891	Feb 21, 2026	U-1434		ODE-52	Sep 06, 2020
	7820788	Oct 27, 2024	DP U-1092			
	7820788	Oct 27, 2024	DP U-1290			
	7820788	Oct 27, 2024	DP U-1434			
	7923536	Dec 09, 2023	U-1117			
	7923536	Dec 09, 2023	U-1290			
	7923536	Dec 09, 2023	U-1434			
	8034375	Aug 13, 2026	U-1290			
	8138229	Dec 09, 2023	DP U-1092			
	8138229	Dec 09, 2023	DP U-1290			
	8138229	Dec 09, 2023	DP U-1434			
	8268348	Feb 21, 2026	U-1290			
	8314156	Dec 09, 2023	U-1290			
	8314156	Dec 09, 2023	U-1434			
	8853260	Oct 10, 2020	DP U-1092			
	8853260	Oct 10, 2020	DP U-1290			
	8853260	Oct 10, 2020	DP U-1434			
	9101543	Feb 21, 2026	U-1434			
	9393318	Mar 04, 2032	U-1290			
	9511046	Jan 12, 2034	U-1434			
	9597409	Mar 04, 2032	U-1290			
<b><u>PALBOCICLIB - IBRANCE</u></b>						
N 207103 001	6936612	Jan 22, 2023	DS DP		I-725	Feb 19, 2019
	7208489	Jan 16, 2023	DS DP		NCE	Feb 03, 2020
	7456168	Jan 16, 2023	U-1998			
<b><u>PALBOCICLIB - IBRANCE</u></b>						
N 207103 002	6936612	Jan 22, 2023	DS DP		I-725	Feb 19, 2019
	7208489	Jan 16, 2023	DS DP		NCE	Feb 03, 2020
	7456168	Jan 16, 2023	U-1998			
<b><u>PALBOCICLIB - IBRANCE</u></b>						
N 207103 003	6936612	Jan 22, 2023	DS DP		I-725	Feb 19, 2019
	7208489	Jan 16, 2023	DS DP		NCE	Feb 03, 2020
	7456168	Jan 16, 2023	U-1998			
<b><u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u></b>						
N 022264 001	6555544	Nov 10, 2018	DP U-543		I-698	Nov 12, 2017
	9439906	Jan 26, 2031	U-1901		M-215	Dec 20, 2020
	9439906	Jan 26, 2031	U-543			
<b><u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u></b>						
N 022264 002	6555544	Nov 10, 2018	DP U-543		I-698	Nov 12, 2017
	9439906	Jan 26, 2031	U-1901		M-215	Dec 20, 2020
	9439906	Jan 26, 2031	U-543			
<b><u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u></b>						
N 022264 003	6555544	Nov 10, 2018	DP U-543		I-698	Nov 12, 2017
	9439906	Jan 26, 2031	U-1901		M-215	Dec 20, 2020
	9439906	Jan 26, 2031	U-543			
<b><u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u></b>						
N 022264 004	6555544	Nov 10, 2018	DP U-543		I-698	Nov 12, 2017
	9439906	Jan 26, 2031	U-1901		M-215	Dec 20, 2020
	9439906	Jan 26, 2031	U-543			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u></b>						
N 022264 004	9439906	Jan 26, 2031		U-543		
<b><u>PALIPERIDONE PALMITATE - INVEGA SUSTENNA</u></b>						
N 022264 005	6555544	Nov 10, 2018	DP	U-543	I-698	Nov 12, 2017
	9439906	Jan 26, 2031		U-1901	M-215	Dec 20, 2020
	9439906	Jan 26, 2031		U-543		
<b><u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u></b>						
N 207946 001	6077843*PED	Nov 12, 2017			NP	May 18, 2018
<b><u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u></b>						
N 207946 002	6077843*PED	Nov 12, 2017			NP	May 18, 2018
<b><u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u></b>						
N 207946 003	6077843*PED	Nov 12, 2017			NP	May 18, 2018
<b><u>PALIPERIDONE PALMITATE - INVEGA TRINZA</u></b>						
N 207946 004	6077843*PED	Nov 12, 2017			NP	May 18, 2018
<b><u>PALONOSETRON HYDROCHLORIDE - ALOXI</u></b>						
N 021372 001	7947724	Jan 30, 2024	DP			
	7947724*PED	Jul 30, 2024				
	7947725	Jan 30, 2024	DP			
	7947725*PED	Jul 30, 2024				
	7960424	Jan 30, 2024	DP			
	7960424*PED	Jul 30, 2024				
	8518981	Jan 30, 2024	DP			
	8518981*PED	Jul 30, 2024				
	8598218	Jan 30, 2024	DP			
	8598218*PED	Jul 30, 2024				
	8598219	Jan 30, 2024	DP			
	8598219*PED	Jul 30, 2024				
	8729094	Jan 30, 2024	DP	U-528		
	8729094*PED	Jul 30, 2024				
	9066980	Jan 30, 2024	DP	U-528		
	9066980*PED	Jul 30, 2024				
	9125905	Jan 30, 2024	DP			
	9125905*PED	Jul 30, 2024				
	9173942	Jan 30, 2024	DP			
	9173942*PED	Jul 30, 2024				
	9439854	Jan 30, 2024	DP			
	9439854*PED	Jul 30, 2024				
	9457020	Jan 30, 2024	DP			
	9457020*PED	Jul 30, 2024				
	9457021	Jan 30, 2024	DP			
	9457021*PED	Jul 30, 2024				
<b><u>PALONOSETRON HYDROCHLORIDE - ALOXI</u></b>						
N 021372 002	7947724	Jan 30, 2024	DP			
	7947724*PED	Jul 30, 2024				
	7947725	Jan 30, 2024	DP			
	7947725*PED	Jul 30, 2024				
	7960424	Jan 30, 2024	DP			
	7960424*PED	Jul 30, 2024				
	8518981	Jan 30, 2024	DP			
	8518981*PED	Jul 30, 2024				
	8598218	Jan 30, 2024	DP			
	8598218*PED	Jul 30, 2024				
	9173942	Jan 30, 2024	DP			
	9173942*PED	Jul 30, 2024				
	9439854	Jan 30, 2024	DP			
	9439854*PED	Jul 30, 2024				
	9457020	Jan 30, 2024	DP			
	9457020*PED	Jul 30, 2024				
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 001	7658918	Feb 20, 2028	DP			
	8221747	Feb 20, 2028	DP			
	8246950	Feb 20, 2028		U-1274		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 001	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 002	7658918	Feb 20, 2028	DP			
	8221747	Feb 20, 2028	DP			
	8246950	Feb 20, 2028	DP	U-1274		
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 003	7658918	Feb 20, 2028	DP			
	8221747	Feb 20, 2028	DP			
	8246950	Feb 20, 2028	DP	U-1274		
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 004	7658918	Feb 20, 2028	DP			
	8221747	Feb 20, 2028	DP			
	8246950	Feb 20, 2028	DP	U-1274		
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 005	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 006	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u></b>						
N 022210 007	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - PANCREAZE</u></b>						
N 022523 005	8221747	Feb 20, 2028	DP			
	8562978	Feb 20, 2028	DP			
	8562979	Feb 20, 2028	DP	U-1274		
	8562980	Feb 20, 2028	DP	U-1274		
	8562981	Feb 20, 2028	DP	U-1274		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u></b>						
N 020725 001	9198871	Feb 07, 2030	DP	U-1787		
<b><u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</u></b>						
N 020725 002	9198871	Feb 07, 2030	DP	U-1787		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</b>						
N 020725 003	9198871	Feb 07, 2030		DP U-1787		
<b>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</b>						
N 020725 004	9198871	Feb 07, 2030		DP U-1787	M-93	Jul 29, 2019
<b>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE); PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - CREON</b>						
N 020725 005	9198871	Feb 07, 2030		DP U-1787	M-93	Jul 29, 2019
<b>PANOBINOSTAT LACTATE - FARYDAK</b>						
N 205353 001	6552065	Aug 31, 2021	DS DP		NCE	Feb 23, 2020
	6833384	Sep 30, 2021	DS DP	U-1669	ODE-89	Feb 23, 2022
	7067551	Aug 31, 2021		U-1669		
	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028		U-1669		
<b>PANOBINOSTAT LACTATE - FARYDAK</b>						
N 205353 002	6552065	Aug 31, 2021	DS DP		NCE	Feb 23, 2020
	6833384	Sep 30, 2021	DS DP	U-1669	ODE-89	Feb 23, 2022
	7067551	Aug 31, 2021		U-1669		
	7989494	Jan 17, 2028	DS DP			
	8883842	Jun 13, 2028		U-1669		
<b>PANTOPRAZOLE SODIUM - PROTONIX IV</b>						
N 020988 001	6780881	Nov 17, 2021	DP			
	7351723	Nov 17, 2021	DP			
	8754108	Nov 17, 2021	DP			
	8754108*PED	May 17, 2022				
<b>PANTOPRAZOLE SODIUM - PROTONIX</b>						
N 022020 001	7544370	Jun 07, 2026	DP			
	7550153	Sep 30, 2024		U-859		
	7553498	Sep 30, 2024		U-859		
	7838027	Sep 30, 2024	DP	U-859		
<b>PARICALCITOL - ZEMPLAR</b>						
N 020819 001	6136799	Apr 08, 2018				
	6361758	Apr 08, 2018	DP			
<b>PARICALCITOL - ZEMPLAR</b>						
N 020819 002	6136799	Apr 08, 2018				
	6361758	Apr 08, 2018	DP			
<b>PARICALCITOL - ZEMPLAR</b>						
N 020819 003	6136799	Apr 08, 2018				
	6136799*PED	Oct 08, 2018				
	6361758	Apr 08, 2018	DP			
	6361758*PED	Oct 08, 2018				
<b>PARICALCITOL - ZEMPLAR</b>						
N 021606 001					NPP	Oct 18, 2019
					NPP	Oct 18, 2019
					ODE-125	Oct 18, 2023
<b>PARICALCITOL - ZEMPLAR</b>						
N 021606 002					NPP	Oct 18, 2019
					NPP	Oct 18, 2019
					ODE-125	Oct 18, 2023

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PARICALCITOL - ZEMPLAR</u></b>						
N 021606	003				NPP	Oct 18, 2019
					NPP	Oct 18, 2019
					ODE-125	Oct 18, 2023
<b><u>PAROXETINE HYDROCHLORIDE - PAXIL</u></b>						
N 020885	001	6063927 6172233	Apr 23, 2019 Jan 15, 2018			
<b><u>PAROXETINE HYDROCHLORIDE - PAXIL</u></b>						
N 020885	002	6063927 6172233	Apr 23, 2019 Jan 15, 2018			
<b><u>PAROXETINE HYDROCHLORIDE - PAXIL</u></b>						
N 020885	003	6063927 6172233	Apr 23, 2019 Jan 15, 2018			
<b><u>PAROXETINE HYDROCHLORIDE - PAXIL</u></b>						
N 020885	004	6063927 6172233	Apr 23, 2019 Jan 15, 2018			
<b><u>PAROXETINE MESYLATE - PEXEVA</u></b>						
N 021299	001	7598271	May 04, 2025	DS		
<b><u>PAROXETINE MESYLATE - PEXEVA</u></b>						
N 021299	002	7598271	May 04, 2025	DS		
<b><u>PAROXETINE MESYLATE - PEXEVA</u></b>						
N 021299	003	7598271	May 04, 2025	DS		
<b><u>PAROXETINE MESYLATE - PEXEVA</u></b>						
N 021299	004	7598271	May 04, 2025	DS		
<b><u>PAROXETINE MESYLATE - BRISDELLE</u></b>						
N 204516	001	7598271 8658663 8946251 9393237	May 04, 2025 Apr 06, 2029 Aug 04, 2026 Aug 04, 2026	DS DS DP U-904 DS DP U-904 U-904		
<b><u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u></b>						
N 200677	001	7473761 8299209	Dec 14, 2026 Dec 27, 2025	DS DP DS DP	NCE ODE-34	Dec 14, 2017 Dec 14, 2019
<b><u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u></b>						
N 200677	002	7473761 8299209	Dec 14, 2026 Dec 27, 2025	DS DP DS DP	NCE ODE-34	Dec 14, 2017 Dec 14, 2019
<b><u>PASIREOTIDE DIASPARTATE - SIGNIFOR</u></b>						
N 200677	003	7473761 8299209	Dec 14, 2026 Dec 27, 2025	DS DP DS DP	NCE ODE-34	Dec 14, 2017 Dec 14, 2019
<b><u>PASIREOTIDE PAMOATE - SIGNIFOR LAR</u></b>						
N 203255	001	7473761 7759308 8822637 9351923	Dec 14, 2026 Oct 25, 2026 Aug 06, 2023 May 23, 2028	DS DP DP U-1629 DP	NCE NP ODE-81	Dec 14, 2017 Dec 15, 2017 Dec 15, 2021
<b><u>PASIREOTIDE PAMOATE - SIGNIFOR LAR</u></b>						
N 203255	002	7473761 7759308 8822637 9351923	Dec 14, 2026 Oct 25, 2026 Aug 06, 2023 May 23, 2028	DS DP DP U-1629 DP	NCE NP ODE-81	Dec 14, 2017 Dec 15, 2017 Dec 15, 2021
<b><u>PASIREOTIDE PAMOATE - SIGNIFOR LAR</u></b>						
N 203255	003	7473761 7759308 8822637 9351923	Dec 14, 2026 Oct 25, 2026 Aug 06, 2023 May 23, 2028	DS DP DP U-1629 DP	NCE NP ODE-81	Dec 14, 2017 Dec 15, 2017 Dec 15, 2021

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PATIROMER SORBITEX CALCIUM - VELTASSA</u></b>						
N 205739 001	7556799	Feb 27, 2025		U-1766		
	8147873	Mar 11, 2026		DP		
	8216560	Mar 14, 2027		U-1766		
	8282913	Mar 11, 2026		DP		
	8287847	Mar 30, 2024		U-1766		
	8337824	May 29, 2030	DS	U-1766		
	8475780	Mar 30, 2024		U-1766		
	8778324	Mar 30, 2024		U-1766		
	8889115	Mar 30, 2024		U-1766		
	9492476	Oct 08, 2033		U-1766		
<b><u>PATIROMER SORBITEX CALCIUM - VELTASSA</u></b>						
N 205739 002	7556799	Feb 27, 2025		U-1766		
	8147873	Mar 11, 2026		DP		
	8216560	Mar 14, 2027		U-1766		
	8282913	Mar 11, 2026		DP		
	8287847	Mar 30, 2024		U-1766		
	8337824	May 29, 2030	DS	U-1766		
	8475780	Mar 30, 2024		U-1766		
	8778324	Mar 30, 2024		U-1766		
	8889115	Mar 30, 2024		U-1766		
	9492476	Oct 08, 2033		U-1766		
<b><u>PATIROMER SORBITEX CALCIUM - VELTASSA</u></b>						
N 205739 003	7556799	Feb 27, 2025		U-1766		
	8147873	Mar 11, 2026		DP		
	8216560	Mar 14, 2027		U-1766		
	8282913	Mar 11, 2026		DP		
	8287847	Mar 30, 2024		U-1766		
	8337824	May 29, 2030	DS	U-1766		
	8475780	Mar 30, 2024		U-1766		
	8778324	Mar 30, 2024		U-1766		
	8889115	Mar 30, 2024		U-1766		
	9492476	Oct 08, 2033		U-1766		
<b><u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u></b>						
N 022465 001	7105530	Oct 19, 2023	DS DP		ODE-23	
	7262203	Dec 19, 2021	DS DP			
	8114885	Dec 19, 2021	DS DP			
<b><u>PAZOPANIB HYDROCHLORIDE - VOTRIENT</u></b>						
N 022465 002	7105530	Oct 19, 2023	DS DP		ODE-23	
	7262203	Dec 19, 2021	DS DP			
	8114885	Dec 19, 2021	DS DP			
<b><u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u></b>						
N 202799 001	7084245	May 12, 2024	DS DP	U-1238		
	7414105	May 12, 2024	DS DP	U-1238		
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<b><u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u></b>						
N 202799 002	7084245	May 12, 2024	DS DP	U-1238		
	7414105	May 12, 2024	DS DP	U-1238		
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<b><u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u></b>						
N 202799 003	7084245	May 12, 2024	DS DP	U-1238		
	7414105	May 12, 2024	DS DP	U-1238		
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u></b>						
N 202799 004	7084245	May 12, 2024	DS DP U-1238			
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<b><u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u></b>						
N 202799 005	7084245	May 12, 2024	DS DP U-1238			
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<b><u>PEGINESATIDE ACETATE - OMONTYS PRESERVATIVE FREE</u></b>						
N 202799 006	7084245	May 12, 2024	DS DP U-1238			
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<b><u>PEGINESATIDE ACETATE - OMONTYS</u></b>						
N 202799 007	7084245	May 12, 2024	DS DP U-1238			
	7414105	May 12, 2024	DS DP U-1238			
	7528104	May 12, 2024	DS DP			
	7550433	Jun 02, 2026		U-1238		
	7919118	May 12, 2024	DS DP			
	7919461	Jun 02, 2026		U-1238		
<b><u>PEMETREXED DISODIUM - ALIMTA</u></b>						
N 021462 001	7772209	Nov 24, 2021		U-1077		
	7772209	Nov 24, 2021		U-1296		
<b><u>PEMETREXED DISODIUM - ALIMTA</u></b>						
N 021462 002	7772209	Nov 24, 2021		U-1296		
<b><u>PENCICLOVIR - DENAVIR</u></b>						
N 020629 001	6469015	Oct 22, 2019		U-501		
	6579981	Jun 17, 2020		U-501		
<b><u>PERAMIVIR - RAPIVAB</u></b>						
N 206426 001	6503745	Nov 05, 2019	DS		NCE	Dec 19, 2019
	6562861	Dec 17, 2018	DS		NPP	Sep 20, 2020
	8778997	May 07, 2027		U-1627		
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 202834 001	6949571	Jun 08, 2021	DS DP U-106		I-710	Jun 19, 2018
	6949571	Jun 08, 2021	DS DP U-2088		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 202834 002	6949571	Jun 08, 2021	DS DP U-106		I-710	Jun 19, 2018
	6949571	Jun 08, 2021	DS DP U-2088		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 202834 003	6949571	Jun 08, 2021	DS DP U-106		I-710	Jun 19, 2018
	6949571	Jun 08, 2021	DS DP U-2088		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 202834 004	6949571	Jun 08, 2021	DS DP U-106		I-710	Jun 19, 2018
	6949571	Jun 08, 2021	DS DP U-2088		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 202834 005	6949571	Jun 08, 2021	DS DP U-106		I-710	Jun 19, 2018
	6949571	Jun 08, 2021	DS DP U-2088		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 202834 006	6949571	Jun 08, 2021	DS DP U-106		I-710	Jun 19, 2018
	6949571	Jun 08, 2021	DS DP U-2088		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			
<b><u>PERAMPANEL - FYCOMPA</u></b>						
N 208277 001	6949571	Jun 08, 2021	DS DP U-106		NCE	Oct 22, 2017
	6949571	Jun 08, 2021	DS DP U-2088			
	6949571	Jun 08, 2021	DS DP U-2089			
	8772497	May 16, 2026	DS			
<b><u>PERFLUTREN - DEFINTITY</u></b>						
N 021064 001	8658205	Jun 18, 2019	DP			
	8685441	Jan 13, 2019	U-665			
	9545457	Jan 13, 2019	U-665			
	9789210	Mar 16, 2037	U-665			
<b><u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u></b>						
N 202088 001	6149938	Jul 23, 2018	DP			
	8440170	Mar 14, 2029	DP			
<b><u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u></b>						
N 202088 002	6149938	Jul 23, 2018	DP			
	8440170	Mar 14, 2029	DP			
<b><u>PHENTERMINE HYDROCHLORIDE - SUPRENZA</u></b>						
N 202088 003	6149938	Jul 23, 2018	DP U-1243			
	8440170	Mar 14, 2029	DP			
<b><u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u></b>						
N 022580 001	7056890	Jun 14, 2020	DP U-1262			
	7553818	Jun 14, 2020	U-1262			
	7659256	Jun 14, 2020	DP U-1262			
	7674776	Jun 14, 2020	DP U-1262			
	8580298	May 15, 2029	DP			
	8580299	Jun 14, 2029	U-1262			
	8895057	Jun 09, 2028	U-1262			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028	U-1262			
<b><u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u></b>						
N 022580 002	7056890	Jun 14, 2020	DP U-1262			
	7553818	Jun 14, 2020	U-1262			
	7659256	Jun 14, 2020	DP U-1262			
	7674776	Jun 14, 2020	DP U-1262			
	8580298	May 15, 2029	DP			
	8580299	Jun 14, 2029	U-1262			
	8895057	Jun 09, 2028	U-1262			
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u></b>						
N 022580 002	9011906	Jun 09, 2028		U-1262		
<b><u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u></b>						
N 022580 003	7056890	Jun 14, 2020	DP	U-1262		
	7553818	Jun 14, 2020		U-1262		
	7659256	Jun 14, 2020	DP	U-1262		
	7674776	Jun 14, 2020	DP	U-1262		
	8580298	May 15, 2029	DP			
	8580299	Jun 14, 2029		U-1262		
	8895057	Jun 09, 2028		U-1262		
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028		U-1262		
<b><u>PHENTERMINE HYDROCHLORIDE; TOPIRAMATE - QSYMIA</u></b>						
N 022580 004	7056890	Jun 14, 2020	DP	U-1262		
	7553818	Jun 14, 2020		U-1262		
	7659256	Jun 14, 2020	DP	U-1262		
	7674776	Jun 14, 2020	DP	U-1262		
	8580298	May 15, 2029	DP			
	8580299	Jun 14, 2029		U-1262		
	8895057	Jun 09, 2028		U-1262		
	8895058	Jun 09, 2028	DP			
	9011905	Jun 09, 2028	DP			
	9011906	Jun 09, 2028		U-1262		
<b><u>PHENTOLAMINE MESYLATE - ORAVERSE</u></b>						
N 022159 001	6764678	May 11, 2021		U-967		
	6872390	May 11, 2021	DP			NPP
	7229630	Jun 20, 2023	DP			Mar 18, 2019
	7569230	Oct 17, 2023		U-967		
	7575757	Apr 21, 2025	DP			
<b><u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u></b>						
N 203510 001	8859623	Nov 14, 2033		U-1594		
<b><u>PHENYLEPHRINE HYDROCHLORIDE - PHENYLEPHRINE HYDROCHLORIDE</u></b>						
N 203510 002	8859623	Nov 14, 2033		U-1594		
<b><u>PIMAVANSERIN TARTRATE - NUPLAZID</u></b>						
N 207318 001	6756393	Mar 06, 2021	DS DP			
	6815458	Mar 06, 2021	DS DP	U-1843		NCE
	7115634	Oct 06, 2021	DS DP			Apr 29, 2021
	7601740	Jun 17, 2027	DS DP			
	7659285	Aug 24, 2026		U-1844		
	7732615	Jun 03, 2028	DS DP			
	7858789	Dec 13, 2020	DS DP			
	7923564	Sep 26, 2025	DS DP			
	8110574	Dec 13, 2020	DS DP			
	8618130	Jan 15, 2024		U-1845		
	8921393	Jan 15, 2024		U-1846		
	9296694	Mar 06, 2021	DS DP			
	9566271	Jan 15, 2024		U-1974		
	9765053	Jul 27, 2022		U-1974		
<b><u>PIMECROLIMUS - ELIDEL</u></b>						
N 021302 001	6423722	Jun 26, 2018				
<b><u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u></b>						
N 050684 001	6900184	Apr 14, 2023	DP	U-282		
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP	U-282		
<b><u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u></b>						
N 050684 002	6900184	Apr 14, 2023	DP	U-282		
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP	U-282		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u></b>						
N 050684 003	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<b><u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN</u></b>						
N 050684 004	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<b><u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u></b>						
N 050750 001	6207661	Feb 22, 2019	DP			
	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<b><u>PIPERACILLIN SODIUM; TAZOBACTAM SODIUM - ZOSYN IN PLASTIC CONTAINER</u></b>						
N 050750 002	6207661	Feb 22, 2019	DP			
	6900184	Apr 14, 2023	DP U-282			
	7915229	Apr 14, 2023	DP			
	8133883	Apr 14, 2023	DP U-282			
<b><u>PIRFENIDONE - ESBRIET</u></b>						
N 022535 001	7566729	Apr 22, 2029	U-1600			
	7635707	Apr 22, 2029	U-1609			
	7696236	Dec 18, 2027	U-1601			
	7767225	Sep 22, 2026	DP U-1602			
	7767700	Dec 18, 2027	U-1601			
	7816383	Jan 08, 2030	U-1603			
	7910610	Jan 08, 2030	U-1604			
	7988994	Sep 22, 2026	DP U-1602			
	8013002	Jan 08, 2030	U-1603			
	8084475	Jan 08, 2030	U-1605			
	8318780	Jan 08, 2030	U-1606			
	8383150	Sep 22, 2026	DP U-1607			
	8420674	Dec 18, 2027	DP U-1608			
	8592462	Apr 22, 2029	U-1609			
	8609701	Apr 22, 2029	U-1610			
	8648098	Jan 08, 2030	U-1611			
	8753679	Sep 22, 2026	DP U-1602			
	8754109	Jan 08, 2030	U-1612			
	8778947	Aug 30, 2033	U-1613			
<b><u>PIRFENIDONE - ESBRIET</u></b>						
N 208780 001	7566729	Apr 22, 2029	U-2077			
	7566729	Apr 22, 2029	U-2078			
	7635707	Apr 22, 2029	U-2072			
	7635707	Apr 22, 2029	U-2073			
	7635707	Apr 22, 2029	U-2074			
	7635707	Apr 22, 2029	U-2075			
	7635707	Apr 22, 2029	U-2076			
	7635707	Apr 22, 2029	U-2083			
	7767700	Dec 18, 2027	U-2080			
	7816383	Jan 08, 2030	U-2042			
	7816383	Jan 08, 2030	U-2050			
	7910610	Jan 08, 2030	U-2048			
	7910610	Jan 08, 2030	U-2049			
	8013002	Jan 08, 2030	U-2047			
	8013002	Jan 08, 2030	U-2082			
	8084475	Jan 08, 2030	U-2052			
	8084475	Jan 08, 2030	U-2054			
	8318780	Jan 08, 2030	U-2046			
	8318780	Jan 08, 2030	U-2081			
	8383150	Sep 22, 2026	DP U-1607			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PIRFENIDONE - ESBRIET</u></b>						
N 208780 001	8420674	Dec 18, 2017	U-2079			
	8592462	Apr 22, 2029	U-2055			
	8592462	Apr 22, 2029	U-2056			
	8592462	Apr 22, 2029	U-2057			
	8592462	Apr 22, 2029	U-2058			
	8592462	Apr 22, 2029	U-2059			
	8592462	Apr 22, 2029	U-2060			
	8592462	Apr 22, 2029	U-2061			
	8592462	Apr 22, 2029	U-2062			
	8592462	Apr 22, 2029	U-2063			
	8609701	Apr 22, 2029	U-2064			
	8609701	Apr 22, 2029	U-2065			
	8609701	Apr 22, 2029	U-2066			
	8609701	Apr 22, 2029	U-2067			
	8609701	Apr 22, 2029	U-2068			
	8609701	Apr 22, 2029	U-2069			
	8609701	Apr 22, 2029	U-2070			
	8648098	Jan 08, 2030	U-2051			
	8648098	Jan 08, 2030	U-2052			
	8754109	Jan 08, 2030	U-2053			
	8778947	Aug 30, 2033	U-2044			
	8778947	Aug 30, 2033	U-2045			
	9561217	Jan 25, 2022	DP			
<b><u>PIRFENIDONE - ESBRIET</u></b>						
N 208780 002					NCE	Oct 15, 2019
					ODE-77	Oct 15, 2021
<b><u>PIRFENIDONE - ESBRIET</u></b>						
N 208780 003	7566729	Apr 22, 2029	U-2077		NCE	Oct 15, 2019
	7566729	Apr 22, 2029	U-2078		ODE-77	Oct 15, 2021
	7635707	Apr 22, 2029	U-2072			
	7635707	Apr 22, 2029	U-2073			
	7635707	Apr 22, 2029	U-2074			
	7635707	Apr 22, 2029	U-2075			
	7635707	Apr 22, 2029	U-2076			
	7635707	Apr 22, 2029	U-2083			
	7767700	Dec 18, 2027	U-2080			
	7816383	Jan 08, 2030	U-2042			
	7816383	Jan 08, 2030	U-2050			
	7910610	Jan 08, 2030	U-2048			
	7910610	Jan 08, 2030	U-2049			
	8013002	Jan 08, 2030	U-2047			
	8013002	Jan 08, 2030	U-2082			
	8084475	Jan 08, 2030	U-2052			
	8084475	Jan 08, 2030	U-2054			
	8318780	Jan 08, 2030	U-2046			
	8318780	Jan 08, 2030	U-2081			
	8383150	Sep 22, 2026	DP U-1607			
	8420674	Dec 18, 2017	U-2079			
	8592462	Apr 22, 2029	U-2055			
	8592462	Apr 22, 2029	U-2056			
	8592462	Apr 22, 2029	U-2057			
	8592462	Apr 22, 2029	U-2058			
	8592462	Apr 22, 2029	U-2059			
	8592462	Apr 22, 2029	U-2060			
	8592462	Apr 22, 2029	U-2061			
	8592462	Apr 22, 2029	U-2062			
	8592462	Apr 22, 2029	U-2063			
	8609701	Apr 22, 2029	U-2064			
	8609701	Apr 22, 2029	U-2065			
	8609701	Apr 22, 2029	U-2066			
	8609701	Apr 22, 2029	U-2067			
	8609701	Apr 22, 2029	U-2068			
	8609701	Apr 22, 2029	U-2069			
	8609701	Apr 22, 2029	U-2070			
	8648098	Jan 08, 2030	U-2051			
	8648098	Jan 08, 2030	U-2052			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PIRFENIDONE - ESBRIET</u></b>						
N 208780 003	8754109	Jan 08, 2030	U-2053			
	8778947	Aug 30, 2033	U-2044			
	8778947	Aug 30, 2033	U-2045			
	9561217	Jan 25, 2022	DP			
<b><u>PITAVASTATIN CALCIUM - LIVALO</u></b>						
N 022363 001	5856336	Dec 25, 2020	DS	U-998		
	7022713	Feb 19, 2024		U-998		
	8557993	Feb 02, 2024		DP		
<b><u>PITAVASTATIN CALCIUM - LIVALO</u></b>						
N 022363 002	5856336	Dec 25, 2020	DS	U-998		
	7022713	Feb 19, 2024		U-998		
	8557993	Feb 02, 2024		DP		
<b><u>PITAVASTATIN CALCIUM - LIVALO</u></b>						
N 022363 003	5856336	Dec 25, 2020	DS	U-998		
	7022713	Feb 19, 2024		U-998		
	8557993	Feb 02, 2024	DS	DP		
<b><u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u></b>						
N 208379 001	8829186	Jan 19, 2031	DS	DP		
<b><u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u></b>						
N 208379 002	8829186	Jan 19, 2031	DS	DP		
<b><u>PITAVASTATIN MAGNESIUM - ZYPITAMAG</u></b>						
N 208379 003	8829186	Jan 19, 2031	DS	DP		
<b><u>PLECANATIDE - TRULANCE</u></b>						
N 208745 001	7041786	Mar 25, 2023	DS		NCE	
	7799897	Jun 09, 2022	DS			Jan 19, 2022
	8637451	Mar 28, 2022		U-1964		
	9610321	Sep 11, 2031		U-1999		
	9616097	Jul 02, 2032		DP		
<b><u>PLERIXAFOR - MOZOBIL</u></b>						
N 022311 001	6987102	Jul 22, 2023		U-936		
	7897590	Jul 22, 2023		U-936		
	RE42152	Dec 10, 2018	DP			
<b><u>POLIDOCANOL - VARITHENA</u></b>						
N 205098 001	6572873	May 26, 2020		U-1461		
	6846412	Jul 19, 2022	DP			
	6942165	May 26, 2020	DP			
	7025290	May 26, 2020	DP	U-1461		
	7357336	May 26, 2020		U-1461		
	7604185	May 26, 2020	DS	DP U-1462		
	7731986	Nov 17, 2024	DS	DP U-1463		
	7814943	Nov 19, 2027	DP	U-1461		
	7842282	May 26, 2020		U-1461		
	7842283	May 26, 2020	DP			
	8122917	Sep 09, 2024	DP			
	8323677	May 26, 2020	DS			
	8734833	May 26, 2020	DS	DP		
	9480652	May 12, 2032	DP			
<b><u>POMALIDOMIDE - POMALYST</u></b>						
N 204026 001	6045501	Aug 28, 2018		U-1361	I-707	Apr 23, 2018
	6315720	Oct 23, 2020		U-1361	NCE	Feb 08, 2018
	6561976	Aug 28, 2018		U-1361	ODE-43	Feb 08, 2020
	6561977	Oct 23, 2020		U-1361		
	6755784	Oct 23, 2020		U-1361		
	6908432	Aug 28, 2018		U-1361		
	8198262	Oct 19, 2024		U-1360		
	8204763	Aug 28, 2018		U-1361		
	8315886	Oct 23, 2020		U-1361		
	8589188	Aug 28, 2018		U-1361		
	8626531	Oct 23, 2020		U-1361		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>POMALIDOMIDE - POMALYST</u></b>						
N 204026 001	8673939	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-1360			
	8828427	Jun 21, 2031	DS DP			
<b><u>POMALIDOMIDE - POMALYST</u></b>						
N 204026 002	6045501	Aug 28, 2018	U-1361	I-707	Apr 23, 2018	
	6315720	Oct 23, 2020	U-1361	NCE	Feb 08, 2018	
	6561976	Aug 28, 2018	U-1361	ODE-43	Feb 08, 2020	
	6561977	Oct 23, 2020	U-1361			
	6755784	Oct 23, 2020	U-1361			
	6908432	Aug 28, 2018	U-1361			
	8198262	Oct 19, 2024	U-1360			
	8204763	Aug 28, 2018	U-1361			
	8315886	Oct 23, 2020	U-1361			
	8589188	Aug 28, 2018	U-1361			
	8626531	Oct 23, 2020	U-1361			
	8673939	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-1360			
	8828427	Jun 21, 2031	DS DP			
<b><u>POMALIDOMIDE - POMALYST</u></b>						
N 204026 003	6045501	Aug 28, 2018	U-1361	I-707	Apr 23, 2018	
	6315720	Oct 23, 2020	U-1361	NCE	Feb 08, 2018	
	6561976	Aug 28, 2018	U-1361	ODE-43	Feb 08, 2020	
	6561977	Oct 23, 2020	U-1361			
	6755784	Oct 23, 2020	U-1361			
	6908432	Aug 28, 2018	U-1361			
	8198262	Oct 19, 2024	U-1360			
	8204763	Aug 28, 2018	U-1361			
	8315886	Oct 23, 2020	U-1361			
	8589188	Aug 28, 2018	U-1361			
	8626531	Oct 23, 2020	U-1361			
	8673939	May 15, 2023	U-1360			
	8735428	May 15, 2023	U-1360			
	8828427	Jun 21, 2031	DS DP			
<b><u>PONATINIB HYDROCHLORIDE - ICLUSIG</u></b>						
N 203469 001	8114874	Dec 22, 2026	DS DP			
	9029533	Dec 22, 2026	U-1283	NCE	Dec 14, 2017	
	9029533	Dec 22, 2026	U-1699	ODE-35	Dec 14, 2019	
	9029533	Dec 22, 2026	U-1700			
	9029533	Dec 22, 2026	U-1701			
	9029533	Dec 22, 2026	U-836			
	9493470	Dec 12, 2033	DS DP U-1700			
	9493470	Dec 12, 2033	DS DP U-1948			
<b><u>PONATINIB HYDROCHLORIDE - ICLUSIG</u></b>						
N 203469 002	8114874	Dec 22, 2026	DS DP			
	9029533	Dec 22, 2026	U-1283	NCE	Dec 14, 2017	
	9029533	Dec 22, 2026	U-1699	ODE-35	Dec 14, 2019	
	9029533	Dec 22, 2026	U-1700			
	9029533	Dec 22, 2026	U-1701			
	9029533	Dec 22, 2026	U-836			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PONATINIB HYDROCHLORIDE - ICLUSIG</u></b>						
N 203469 002	9493470	Dec 12, 2033	DS DP U-1700			
	9493470	Dec 12, 2033	DS DP U-1948			
<b><u>PONATINIB HYDROCHLORIDE - ICLUSIG</u></b>						
N 203469 003	8114874	Dec 22, 2026	DS DP		NCE	Dec 14, 2017
	9029533	Dec 22, 2026		U-1283	ODE-35	Dec 14, 2019
	9029533	Dec 22, 2026		U-1699		
	9029533	Dec 22, 2026		U-1700		
	9029533	Dec 22, 2026		U-1701		
	9029533	Dec 22, 2026		U-836		
	9493470	Dec 12, 2033	DS DP U-1700			
	9493470	Dec 12, 2033	DS DP U-1948			
<b><u>POSACONAZOLE - NOXAFIL</u></b>						
N 022003 001	5661151	Jul 19, 2019	DS DP U-760			
	6958337	Oct 05, 2018	DS DP U-760			
	8263600	Apr 01, 2022	DP			
<b><u>POSACONAZOLE - NOXAFIL</u></b>						
N 205053 001	5661151	Jul 19, 2019	DS DP U-1454			
<b><u>POSACONAZOLE - NOXAFIL</u></b>						
N 205596 001	5661151	Jul 19, 2019	DS DP U-1454			
	8410077	Mar 13, 2029	DP			
	9023790	Jul 04, 2031	DP U-1698			
	9358297	Jun 24, 2031	DP U-1454			
	9493582	Feb 27, 2033	DP			
	9750822	Mar 13, 2029	DP			
<b><u>PRALATREXATE - FOLOTYN</u></b>						
N 022468 001	6028071	Jul 16, 2022	DS DP U-1004			
	7622470	May 31, 2025		U-1015		
	8299078	May 31, 2025		U-1004		
<b><u>PRALATREXATE - FOLOTYN</u></b>						
N 022468 002	6028071	Jul 16, 2022	DS DP U-1004			
	7622470	May 31, 2025		U-1015		
	8299078	May 31, 2025		U-1004		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u></b>						
N 020667 001	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u></b>						
N 020667 002	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u></b>						
N 020667 003	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u></b>						
N 020667 005	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u></b>						
N 020667 006	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u></b>						
N 020667 007	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 001	7695734	Apr 26, 2028	DP			
	8679533	Sep 08, 2029	DP U-219			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 002	7695734 8679533	Apr 26, 2028 Sep 08, 2029	DP DP U-219			
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 003	7695734 8679533	Apr 26, 2028 Sep 08, 2029	DP DP U-219			
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 004	7695734 8679533	Apr 26, 2028 Sep 08, 2029	DP DP U-219			
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 005	7695734 8679533	Apr 26, 2028 Sep 08, 2029	DP DP U-219			
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 006	7695734 8679533	Apr 26, 2028 Sep 08, 2029	DP DP U-219			
<b><u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u></b>						
N 022421 007	7695734 8679533	Apr 26, 2028 Sep 08, 2029	DP DP U-219			
<b><u>PRAMINTIDE ACETATE - SYMLIN</u></b>						
N 021332 001	5686411 6114304	Mar 16, 2019 Sep 05, 2017	DS DP U-638 U-640			
<b><u>PRAMINTIDE ACETATE - SYMLIN</u></b>						
N 021332 002	5686411 6114304 6114304	Mar 16, 2019 Sep 05, 2017 Sep 05, 2017	DS DP U-638 U-637 U-640			
<b><u>PRAMINTIDE ACETATE - SYMLIN</u></b>						
N 021332 003	5686411 6114304 6114304	Mar 16, 2019 Sep 05, 2017 Sep 05, 2017	DS DP U-638 U-637 U-640			
<b><u>PRASTERONE - INTRAROSA</u></b>						
N 208470 001	8268806 8629129 8957054	Mar 19, 2031 Aug 07, 2028 Aug 07, 2028	DP DP U-1922		NCE	Nov 16, 2021
<b><u>PRASUGREL HYDROCHLORIDE - PRASUGREL</u></b>						
A 205927 001					PC	Feb 11, 2018
<b><u>PRASUGREL HYDROCHLORIDE - PRASUGREL</u></b>						
A 205927 002					PC	Feb 11, 2018
<b><u>PRASUGREL HYDROCHLORIDE - EFFIENT</u></b>						
N 022307 001	5288726*PED 8404703 8404703*PED 8569325 8569325*PED	Oct 14, 2017 Jan 02, 2023 Jul 02, 2023 Jan 02, 2023 Jul 02, 2023	U-1381     		M-182 PED	Jul 12, 2019 Jan 12, 2020
<b><u>PRASUGREL HYDROCHLORIDE - EFFIENT</u></b>						
N 022307 002	5288726*PED 8404703 8404703*PED 8569325 8569325*PED	Oct 14, 2017 Jan 02, 2023 Jul 02, 2023 Jan 02, 2023 Jul 02, 2023	U-1381     		M-182 PED	Jul 12, 2019 Jan 12, 2020
<b><u>PREDNISOLONE ACETATE - FLO-PRED</u></b>						
N 022067 001	6071523 6399079 6656482 7799331 7799331	Jun 03, 2018 Jun 03, 2018 Jun 03, 2018 Oct 11, 2028 Oct 11, 2028	DP DP DP DP U-1068 DP U-139			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PREDNISOLONE ACETATE - FLO-PRED</u></b>						
N 022067 002	6071523	Jun 03, 2018	DP			
	6399079	Jun 03, 2018	DP			
	6656482	Jun 03, 2018	DP			
	7799331	Oct 11, 2028	DP U-1068			
	7799331	Oct 11, 2028	DP U-139			
<b><u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u></b>						
N 021959 001	6740341	Nov 24, 2019	DP			
<b><u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u></b>						
N 021959 002	6740341	Nov 24, 2019	DP			
<b><u>PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT</u></b>						
N 021959 003	6740341	Nov 24, 2019	DP			
<b><u>PREDNISONE - RAYOS</u></b>						
N 202020 001	6488960	Mar 14, 2020	DP U-1267			
	6677326	Mar 14, 2020	DP U-1268			
	8309124	Apr 23, 2024	U-1292			
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<b><u>PREDNISONE - RAYOS</u></b>						
N 202020 002	6488960	Mar 14, 2020	DP U-1267			
	6677326	Mar 14, 2020	DP U-1268			
	8309124	Apr 23, 2024	U-1292			
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<b><u>PREDNISONE - RAYOS</u></b>						
N 202020 003	8168218	Jan 07, 2028	DP U-1269			
	8309124	Apr 23, 2024	U-1292			
	8394407	Apr 23, 2024	DP U-1362			
	9040085	Apr 23, 2024	U-1362			
	9186332	Apr 23, 2024	U-1362			
	9504699	Aug 03, 2027	U-1362			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 001	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 002	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 003	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 004	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 005	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 005	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 006	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 007	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA</u></b>						
N 021446 008	6001876	Dec 30, 2018	U-55	Y	M-193	Dec 22, 2019
	6001876	Dec 30, 2018	U-819	Y		
	6197819	Dec 30, 2018	DS DP			
	RE41920	Dec 30, 2018	U-1250			
<b><u>PREGABALIN - LYRICA CR</u></b>						
N 209501 001	6197819	Dec 30, 2018	DS DP		NP	Oct 11, 2020
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	9144559	Nov 02, 2026	DP			
	RE41920	Dec 30, 2018	U-2136			
	RE41920	Dec 30, 2018	U-2137			
<b><u>PREGABALIN - LYRICA CR</u></b>						
N 209501 002	6197819	Dec 30, 2018	DS DP		NP	Oct 11, 2020
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	9144559	Nov 02, 2026	DP			
	RE41920	Dec 30, 2018	U-2136			
	RE41920	Dec 30, 2018	U-2137			
<b><u>PREGABALIN - LYRICA CR</u></b>						
N 209501 003	6197819	Dec 30, 2018	DS DP		NP	Oct 11, 2020
	8945620	Nov 02, 2026	DP U-2136			
	8945620	Nov 02, 2026	DP U-2137			
	9144559	Nov 02, 2026	DP			
	RE41920	Dec 30, 2018	U-2136			
	RE41920	Dec 30, 2018	U-2137			
<b><u>PROGESTERONE - ENDOMETRIN</u></b>						
N 022057 001	7300664	Nov 17, 2019	U-856			
	7320800	Nov 17, 2019	U-856			
	7393543	Nov 17, 2019	DP U-880			
<b><u>PROPOFOL - DIPRIVAN</u></b>						
N 019627 002	8476010	Dec 01, 2024	DS DP			
	8476010*PED	Jun 01, 2025				
<b><u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u></b>						
N 021438 001	6500454	Oct 04, 2021	DP			
<b><u>PROPRANOLOL HYDROCHLORIDE - INNOPRAN XL</u></b>						
N 021438 002	6500454	Oct 04, 2021	DP			
<b><u>PROPRANOLOL HYDROCHLORIDE - HEMANGEOL</u></b>						
N 205410 001	8338489	Oct 16, 2028	U-1496		ODE-62	Mar 14, 2021

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>QUAZEPAM - DORAL</u></b>						
N 018708 001	7608616	Jun 03, 2028		U-1012		
<b><u>QUAZEPAM - DORAL</u></b>						
N 018708 003	7608616	Jun 03, 2028		U-1012		
<b><u>RADIUM RA-223 DICHLORIDE - XOFIGO</u></b>						
N 203971 001	6635234	Jan 03, 2020		U-1405	NCE	May 15, 2018
<b><u>RALTEGRAVIR POTASSIUM - ISENTRESS</u></b>						
N 022145 001	7169780	Oct 03, 2023	DS DP		D-167	May 26, 2020
	7169780*PED	Apr 03, 2024			NPP	Nov 22, 2020
	7217713	Oct 21, 2022		U-257	PED	Nov 26, 2020
	7217713*PED	Apr 21, 2023			PED	May 22, 2021
	7435734	Oct 21, 2022		U-257		
	7435734	Oct 21, 2022		U-900		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
<b><u>RALTEGRAVIR POTASSIUM - ISENTRESS HD</u></b>						
N 022145 002	7169780	Oct 03, 2023	DS DP		NPP	Nov 22, 2020
	7169780*PED	Apr 03, 2024			NS	May 26, 2020
	7217713	Oct 21, 2022		U-257	PED	Nov 26, 2020
	7217713*PED	Apr 21, 2023			PED	May 22, 2021
	7435734	Oct 21, 2022		U-257		
	7435734	Oct 21, 2022		U-900		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
	9649311	Oct 21, 2030	DP			
	9649311*PED	Apr 21, 2031				
<b><u>RALTEGRAVIR POTASSIUM - ISENTRESS</u></b>						
N 203045 001	7169780	Oct 03, 2023	DS DP		NPP	Nov 22, 2020
	7169780*PED	Apr 03, 2024			PED	May 22, 2021
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	7435734	Oct 21, 2022		U-257		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
<b><u>RALTEGRAVIR POTASSIUM - ISENTRESS</u></b>						
N 203045 002	7169780	Oct 03, 2023	DS DP		NPP	Nov 22, 2020
	7169780*PED	Apr 03, 2024			PED	May 22, 2021
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	7435734	Oct 21, 2022		U-257		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
<b><u>RALTEGRAVIR POTASSIUM - ISENTRESS</u></b>						
N 205786 001	7169780	Oct 03, 2023	DS DP		NPP	Nov 22, 2020
	7169780*PED	Apr 03, 2024			PED	May 22, 2021
	7217713	Oct 21, 2022		U-257		
	7217713*PED	Apr 21, 2023				
	7435734	Oct 21, 2022		U-257		
	7435734*PED	Apr 21, 2023				
	7754731	Mar 11, 2029	DS DP	U-257		
	7754731*PED	Sep 11, 2029				
<b><u>RAMELTEON - ROZEREM</u></b>						
N 021782 001	6034239	Jul 22, 2019	DS DP	U-674		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 019901 001	7368469	Aug 30, 2020		U-871		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>RAMIPRIL - ALTACE</u></b>						
N 019901 002	7368469	Aug 30, 2020		U-871		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 019901 003	7368469	Aug 30, 2020		U-871		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 019901 004	7368469	Aug 30, 2020		U-871		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 022021 001	7368469	Aug 30, 2020		U-871		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 022021 002	7368469	Aug 30, 2020		U-871		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 022021 003	7368469	Aug 30, 2020		U-871		
<b><u>RAMIPRIL - ALTACE</u></b>						
N 022021 004	7368469	Aug 30, 2020		U-871		
<b><u>RANOLAZINE - RANEXA</u></b>						
N 021526 001	6303607	May 27, 2019		U-705		
	6369062	May 27, 2019	DP		Y	
	6479496	May 27, 2019		U-705		
	6503911	May 27, 2019	DP			
	6525057	May 27, 2019		U-705		
	6562826	May 27, 2019		U-705		
	6617328	May 27, 2019	DP			
	6620814	May 27, 2019		U-705		
	6852724	May 27, 2019		U-705		
	6864258	May 27, 2019		U-705		
<b><u>RANOLAZINE - RANEXA</u></b>						
N 021526 002	6303607	May 27, 2019		U-705		
	6369062	May 27, 2019	DP			
	6479496	May 27, 2019		U-705		
	6503911	May 27, 2019	DP			
	6525057	May 27, 2019		U-705		
	6562826	May 27, 2019		U-705		
	6617328	May 27, 2019	DP			
	6620814	May 27, 2019		U-705		
	6852724	May 27, 2019		U-705		
	6864258	May 27, 2019		U-705		
<b><u>RASAGILINE MESYLATE - RASAGILINE MESYLATE</u></b>						
A 201823 001					PC	Jul 01, 2017
<b><u>RASAGILINE MESYLATE - RASAGILINE MESYLATE</u></b>						
A 201823 002					PC	Jul 01, 2017
<b><u>RASAGILINE MESYLATE - RASAGILINE MESYLATE</u></b>						
A 201950 001					PC	Jul 01, 2017
<b><u>RASAGILINE MESYLATE - RASAGILINE MESYLATE</u></b>						
A 201950 002					PC	Jul 01, 2017
<b><u>RASAGILINE MESYLATE - AZILECT</u></b>						
N 021641 001	7572834	Dec 05, 2026	DP			
	7815942	Aug 27, 2027	DS DP U-219			
<b><u>RASAGILINE MESYLATE - AZILECT</u></b>						
N 021641 002	7572834	Dec 05, 2026	DP			
	7815942	Aug 27, 2027	DS DP U-219			
<b><u>REGADENOSON - LEXISCAN</u></b>						
N 022161 001	6403567	Apr 10, 2022	DS DP U-869		M-194	Jan 17, 2020
	6642210	Jun 22, 2019	DS DP U-869			
	7144872	Jun 22, 2019	DS DP U-116			
	7144872	Jun 22, 2019	DS DP U-869			
	7144872	Jun 22, 2019	DS DP U-870			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>REGADENOSON - LEXISCAN</u></b>						
N 022161 001	7183264	Jun 22, 2019	DP U-116			
	7183264	Jun 22, 2019	DP U-869			
	7183264	Jun 22, 2019	DP U-870			
	7582617	Jun 22, 2019	U-1003			
	7655636	Jun 22, 2019	U-869			
	7655637	Jun 22, 2019	DS DP U-869			
	7683037	Jun 22, 2019	U-1042			
	8106029	Jun 22, 2019	U-1042			
	8106183	Feb 02, 2027	DS			
	8133879	Jun 22, 2019	DP			
	8183226	Jun 22, 2019	U-116			
	8470801	Jun 22, 2019	U-116			
	8536150	Jun 22, 2019	U-116			
	9045519	Jun 22, 2019	DP			
	9085601	Feb 02, 2027	DP			
	9289446	Jun 22, 2019	DP U-116			
<b><u>REGORAFENIB - STIVARGA</u></b>						
N 203085 001	7351834	Jun 28, 2022	DS		I-744	Apr 27, 2020
	8637553	Feb 16, 2031	DS DP		NCE	Sep 27, 2017
	8680124	Jun 02, 2030	U-1506		ODE-139	Apr 27, 2024
	9458107	Apr 08, 2031	DP		ODE-44	Feb 25, 2020
<b><u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u></b>						
N 020630 001	5866591	Sep 10, 2017	DP			
<b><u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u></b>						
N 020630 002	5866591	Sep 10, 2017	DP			
<b><u>REMIFENTANIL HYDROCHLORIDE - ULTIVA</u></b>						
N 020630 003	5866591	Sep 10, 2017	DP			
<b><u>RETAPAMULIN - ALTABAX</u></b>						
N 022055 001	7875630	Feb 14, 2027	DS			
	8207191	Aug 30, 2024	U-805			
	RE43390	Apr 12, 2021	DS DP U-805			
<b><u>RIBAVIRIN - VIRAZOLE</u></b>						
N 018859 001	6150337	Nov 21, 2017	U-400			
<b><u>RIBAVIRIN - REBETOL</u></b>						
N 020903 001	6172046	Sep 21, 2017	U-1014			
	6172046	Sep 21, 2017	U-377			
	6472373	Sep 21, 2017	U-1014			
	6472373	Sep 21, 2017	U-479			
<b><u>RIBAVIRIN - REBETOL</u></b>						
N 020903 002	6172046	Sep 21, 2017	U-1014			
	6172046	Sep 21, 2017	U-377			
	6472373	Sep 21, 2017	U-1014			
	6472373	Sep 21, 2017	U-479			
<b><u>RIBAVIRIN - REBETOL</u></b>						
N 021546 001	6172046	Sep 21, 2017	U-1014			
	6172046	Sep 21, 2017	U-521			
	6472373	Sep 21, 2017	U-521			
	6790837	Apr 05, 2023	DP			
<b><u>RIBOCICLIB SUCCINATE - KISQALI</u></b>						
N 209092 001	8324225	Jun 17, 2028	DS DP		NCE	Mar 13, 2022
	8415355	Feb 19, 2031	DS DP			
	8685980	May 25, 2030	DS DP			
	8962630	Dec 09, 2029	U-1981			
	9193732	Nov 09, 2031	DS DP			
	9416136	Aug 20, 2029	U-1981			
<b><u>RIBOFLAVIN 5'-PHOSPHATE SODIUM - PHOTREXA</u></b>						
N 203324 001					NP	Apr 15, 2019
					ODE-116	Apr 15, 2023

PREScription AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>RIOCIGUAT - ADEMPAS</b>						
N 204819 001	6743798	Jul 16, 2019	DS DP		NCE	Oct 08, 2018
	7173037	Apr 25, 2023	DS DP		ODE-53	Oct 08, 2020
<b>RIOCIGUAT - ADEMPAS</b>						
N 204819 002	6743798	Jul 16, 2019	DS DP		NCE	Oct 08, 2018
	7173037	Apr 25, 2023	DS DP		ODE-53	Oct 08, 2020
<b>RIOCIGUAT - ADEMPAS</b>						
N 204819 003	6743798	Jul 16, 2019	DS DP		NCE	Oct 08, 2018
	7173037	Apr 25, 2023	DS DP		ODE-53	Oct 08, 2020
<b>RIOCIGUAT - ADEMPAS</b>						
N 204819 004	6743798	Jul 16, 2019	DS DP		NCE	Oct 08, 2018
	7173037	Apr 25, 2023	DS DP		ODE-53	Oct 08, 2020
<b>RIOCIGUAT - ADEMPAS</b>						
N 204819 005	6743798	Jul 16, 2019	DS DP		NCE	Oct 08, 2018
	7173037	Apr 25, 2023	DS DP		ODE-53	Oct 08, 2020
<b>RISEDRONATE SODIUM - ACTONEL</b>						
N 020835 001	6165513				Jun 10, 2018	
<b>RISEDRONATE SODIUM - ACTONEL</b>						
N 020835 002	6165513				Jun 10, 2018	
<b>RISEDRONATE SODIUM - ACTONEL</b>						
N 020835 003	5994329	Jul 17, 2018		U-353		
	6015801	Jul 17, 2018		U-353		
	6165513	Jun 10, 2018	DP			
	6432932	Jul 17, 2018		U-595		
	6465443	Aug 14, 2018	DP			
<b>RISEDRONATE SODIUM - ACTONEL</b>						
N 020835 004	6165513			DP		
<b>RISEDRONATE SODIUM - ACTONEL</b>						
N 020835 005	6165513	Jun 10, 2018	DP			
	7192938	May 06, 2023		U-353		
	7718634	May 06, 2023		U-662		
<b>RISEDRONATE SODIUM - ATELVIA</b>						
N 022560 001	7645459	Jan 09, 2028	DP	U-662		
	7645460	Jan 09, 2028	DP	U-662		
	8246989	Jan 16, 2026	DP			
<b>RISPERIDONE - RISPERDAL CONSTA</b>						
N 021346 001	6194006	Dec 30, 2018	DP			
	6379703	Dec 30, 2018	DP			
	6596316	Dec 30, 2018	DP			
	6667061	May 25, 2020	DP			
<b>RISPERIDONE - RISPERDAL CONSTA</b>						
N 021346 002	6194006	Dec 30, 2018	DP			
	6379703	Dec 30, 2018	DP			
	6596316	Dec 30, 2018	DP			
	6667061	May 25, 2020	DP			
<b>RISPERIDONE - RISPERDAL CONSTA</b>						
N 021346 003	6194006	Dec 30, 2018	DP			
	6379703	Dec 30, 2018	DP			
	6596316	Dec 30, 2018	DP			
	6667061	May 25, 2020	DP			
<b>RISPERIDONE - RISPERDAL CONSTA</b>						
N 021346 004	6194006	Dec 30, 2018	DP			
	6379703	Dec 30, 2018	DP			
	6596316	Dec 30, 2018	DP			
	6667061	May 25, 2020	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>RITONAVIR - NORVIR</b>						
N 020945 001	6232333	Nov 07, 2017				
	7141593	May 22, 2020	DP			
	7432294	May 22, 2020	DP			
<b>RITONAVIR - NORVIR</b>						
N 022417 001	7148359	Jul 19, 2019	DP			
	7364752	Nov 10, 2020	DP U-688			
	8268349	Aug 25, 2024	DP			
	8399015	Aug 25, 2024	DP			
	8399015*PED	Feb 25, 2025				
	8470347	Sep 17, 2026	DP			
	8470347*PED	Mar 17, 2027				
	8691878	Aug 25, 2024	U-688			
	8691878*PED	Feb 25, 2025				
<b>RIVAROXABAN - XARELTO</b>						
N 022406 001	7157456	Aug 28, 2024	DS DP U-1301		D-168	Oct 27, 2020
	7157456	Aug 28, 2024	DS DP U-1302			
	7585860	Dec 11, 2020	DS			
	7592339	Dec 11, 2020	U-1167			
	7592339	Dec 11, 2020	U-1200			
	7592339	Dec 11, 2020	U-1301			
	7592339	Dec 11, 2020	U-1302			
	7592339	Dec 11, 2020	U-1303			
	7592339	Dec 11, 2020	U-2142			
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-1303			
	9415053	Nov 13, 2024	DP U-2142			
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-1956			
	9539218	Feb 17, 2034	U-1957			
	9539218	Feb 17, 2034	U-2143			
<b>RIVAROXABAN - XARELTO</b>						
N 022406 002	7157456	Aug 28, 2024	DS DP U-1301			
	7157456	Aug 28, 2024	DS DP U-1302			
	7585860	Dec 11, 2020	DS			
	7592339	Dec 11, 2020	U-1167			
	7592339	Dec 11, 2020	U-1200			
	7592339	Dec 11, 2020	U-1301			
	7592339	Dec 11, 2020	U-1302			
	7592339	Dec 11, 2020	U-1303			
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			
	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-1303			
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-1956			
	9539218	Feb 17, 2034	U-1957			
<b>RIVAROXABAN - XARELTO</b>						
N 022406 003	7157456	Aug 28, 2024	DS DP U-1301			
	7157456	Aug 28, 2024	DS DP U-1302			
	7585860	Dec 11, 2020	DS			
	7592339	Dec 11, 2020	U-1167	Y		
	7592339	Dec 11, 2020	U-1200	Y		
	7592339	Dec 11, 2020	U-1301	Y		
	7592339	Dec 11, 2020	U-1302	Y		
	9415053	Nov 13, 2024	DP U-1167			
	9415053	Nov 13, 2024	DP U-1200			
	9415053	Nov 13, 2024	DP U-1301			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>RIVAROXABAN - XARELTO</b>						
N 022406 003	9415053	Nov 13, 2024	DP U-1302			
	9415053	Nov 13, 2024	DP U-1303			
	9539218	Feb 17, 2034	U-1953			
	9539218	Feb 17, 2034	U-1954			
	9539218	Feb 17, 2034	U-1955			
	9539218	Feb 17, 2034	U-1957			
<b>RIVASTIGMINE - EXELON</b>						
N 022083 001	6316023	Jan 08, 2019	DP			
	6335031	Jan 08, 2019	DP			
<b>RIVASTIGMINE - EXELON</b>						
N 022083 002	6316023	Jan 08, 2019	DP			
	6335031	Jan 08, 2019	DP			
<b>RIVASTIGMINE - EXELON</b>						
N 022083 005	6316023	Jan 08, 2019	DP			
	6335031	Jan 08, 2019	DP			
<b>ROFLUMILAST - DALIRESP</b>						
N 022522 001	5712298	Jan 27, 2020	DS DP U-1115		M-208	
	8431154	Feb 19, 2023	DP			
	8536206	Mar 08, 2024	U-1115			
	8604064	Mar 08, 2024	U-1115			
	8618142	Mar 08, 2024	DP			
	9468598	Feb 19, 2023	DP			
<b>ROLAPITANT HYDROCHLORIDE - VARUBI</b>						
N 206500 001	7049320	Dec 08, 2023	DS DP U-1741		NCE	
	7563801	Apr 04, 2027	DP			
	7981905	Apr 04, 2027	U-1741			
	8178550	Apr 04, 2027	DS DP			
	8361500	Oct 09, 2029	DP			
	8404702	Apr 04, 2027	U-1741			
	8470842	Jan 18, 2029	U-1741			
	8796299	Dec 17, 2022	U-1741			
<b>ROLAPITANT HYDROCHLORIDE - VARUBI</b>						
N 208399 001	7049320	Dec 08, 2023	DS DP U-1741		NCE	
	7981905	Apr 04, 2027	U-1741			
	8178550	Apr 04, 2027	DS DP			
	8404702	Apr 04, 2027	U-1741			
	8470842	Jan 18, 2029	U-1741			
	8796299	Dec 17, 2022	U-1741			
	9101615	Jul 14, 2032	U-1741			
<b>ROMIDEPSIN - ISTODAX</b>						
N 022393 001	7608280	Aug 22, 2021	DS		ODE-12	
	7611724	Aug 22, 2021	DS			
<b>ROPINIROLE HYDROCHLORIDE - REQUIP XL</b>						
N 022008 001	7927624	Dec 02, 2021	DP U-20		M-203	
	8303986	Apr 12, 2021	DP			
<b>ROPINIROLE HYDROCHLORIDE - REQUIP XL</b>						
N 022008 002	7927624	Dec 02, 2021	DP U-20		M-203	
	8303986	Apr 12, 2021	DP			
<b>ROPINIROLE HYDROCHLORIDE - REQUIP XL</b>						
N 022008 003	7927624	Dec 02, 2021	DP U-20		M-203	
	8303986	Apr 12, 2021	DP			
<b>ROPINIROLE HYDROCHLORIDE - REQUIP XL</b>						
N 022008 004	7927624	Dec 02, 2021	DP U-20		M-203	
	8303986	Apr 12, 2021	DP			
<b>ROPINIROLE HYDROCHLORIDE - REQUIP XL</b>						
N 022008 005	7927624	Dec 02, 2021	DP U-20		M-203	
	8303986	Apr 12, 2021	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ROPINIROLE HYDROCHLORIDE - REQUIP XL</u></b>						
N 022008 006	7927624	Dec 02, 2021	DP	U-20	M-203	Mar 23, 2020
	8303986	Apr 12, 2021	DP			
<b><u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u></b>						
N 020533 006	7828787	Oct 18, 2025	DP			
	7857802	Nov 28, 2026	DP			
	8118802	May 18, 2023	DP			
	8162915	May 23, 2024	DP			
<b><u>ROPIVACAINE HYDROCHLORIDE - NAROPIN</u></b>						
N 020533 007	7828787	Oct 18, 2025	DP			
	7857802	Nov 28, 2026	DP			
	8118802	May 18, 2023	DP			
	8162915	May 23, 2024	DP			
<b><u>ROSIGLITAZONE MALEATE - AVANDIA</u></b>						
N 021071 002	7358366	Apr 19, 2020	DS			
<b><u>ROSIGLITAZONE MALEATE - AVANDIA</u></b>						
N 021071 003	7358366	Apr 19, 2020	DS			
<b><u>ROSIGLITAZONE MALEATE - AVANDIA</u></b>						
N 021071 004	7358366	Apr 19, 2020	DS			
<b><u>ROSUVASTATIN CALCIUM - CRESTOR</u></b>						
N 021366 002	6316460	Aug 04, 2020	DP		NPP	Nov 20, 2018
	6858618	Dec 17, 2021	U-1032		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-1807			
	6858618	Dec 17, 2021	U-618			
	6858618*PED	Jun 17, 2022				
	7030152	Apr 02, 2018	U-1032			
	7964614	Apr 02, 2018	U-1032			
<b><u>ROSUVASTATIN CALCIUM - CRESTOR</u></b>						
N 021366 003	6316460	Aug 04, 2020	DP		NPP	Nov 20, 2018
	6858618	Dec 17, 2021	U-1032		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-1807			
	6858618	Dec 17, 2021	U-618			
	6858618*PED	Jun 17, 2022				
	7030152	Apr 02, 2018	U-1032			
	7964614	Apr 02, 2018	U-1032			
<b><u>ROSUVASTATIN CALCIUM - CRESTOR</u></b>						
N 021366 004	6316460	Aug 04, 2020	DP		I-732	May 27, 2019
	6858618	Dec 17, 2021	U-1032		NPP	Nov 20, 2018
	6858618	Dec 17, 2021	U-1807		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-618			
	6858618*PED	Jun 17, 2022				
	7030152	Apr 02, 2018	U-1032			
	7964614	Apr 02, 2018	U-1032			
<b><u>ROSUVASTATIN CALCIUM - CRESTOR</u></b>						
N 021366 005	6316460	Aug 04, 2020	DP		ODE-118	May 27, 2023
	6858618	Dec 17, 2021	U-618			
	7030152	Apr 02, 2018	U-1032			
	7964614	Apr 02, 2018	U-1032			
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 001	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	7413747	Mar 18, 2019	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 002	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 002	7413747	Mar 18, 2019	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 003	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	7413747	Mar 18, 2019	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 004	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	7413747	Mar 18, 2019	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 005	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	7413747	Mar 18, 2019	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
<b><u>ROTIGOTINE - NEUPRO</u></b>						
N 021829 006	6699498	Nov 27, 2020	DP			
	6884434	Mar 30, 2021	DP			
	7413747	Mar 18, 2019	DP			
	8246979	Sep 01, 2027	DP U-1272			
	8246979	Sep 01, 2027	DP U-1273			
	8246980	Nov 27, 2025	DP			
	8617591	Jul 22, 2023	DP U-1474			
<b><u>RUCAPARIB CAMSYLATE - RUBRACA</u></b>						
N 209115 001	6495541	Jan 10, 2020	DS DP		NCE	Dec 19, 2021
	7351701	Jul 23, 2024	U-2012		ODE-126	Dec 19, 2023
	7531530	Jul 23, 2024	U-2012			
	8071579	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2012			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	9045487	Feb 10, 2031	DS DP			
<b><u>RUCAPARIB CAMSYLATE - RUBRACA</u></b>						
N 209115 002	6495541	Jan 10, 2020	DS DP		NCE	Dec 19, 2021
	7351701	Jul 23, 2024	U-2012		ODE-126	Dec 19, 2023
	7531530	Jul 23, 2024	U-2012			
	8071579	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2012			
	8754072	Feb 10, 2031	DS DP			
	8859562	Aug 04, 2031	U-2012			
	9045487	Feb 10, 2031	DS DP			
<b><u>RUCAPARIB CAMSYLATE - RUBRACA</u></b>						
N 209115 003	6495541	Jan 10, 2020	DS DP		NCE	Dec 19, 2021
	7351701	Jul 23, 2024	U-2012		ODE-126	Dec 19, 2023
	7531530	Jul 23, 2024	U-2012			
	8071579	Aug 12, 2027	U-2012			
	8143241	Aug 12, 2027	U-2012			
	8754072	Feb 10, 2031	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>RUCAPARIB CAMSYLATE - RUBRACA</b>						
N 209115 003	8859562	Aug 04, 2031		U-2012		
	9045487	Feb 10, 2031	DS DP			
<b>RUFINAMIDE - BANZEL</b>						
N 021911 001	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
	7750028	Oct 19, 2018		U-106		
	7750028*PED	Apr 19, 2019				
	8076362	Jun 08, 2018	DP			
	8076362*PED	Dec 08, 2018				
<b>RUFINAMIDE - BANZEL</b>						
N 021911 002	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
	7750028	Oct 19, 2018		U-106		
	7750028*PED	Apr 19, 2019				
	8076362	Jun 08, 2018	DP			
	8076362*PED	Dec 08, 2018				
<b>RUFINAMIDE - BANZEL</b>						
N 021911 003	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
	7750028	Oct 19, 2018		U-106		
	7750028*PED	Apr 19, 2019				
	8076362	Jun 08, 2018	DP			
	8076362*PED	Dec 08, 2018				
<b>RUFINAMIDE - BANZEL</b>						
N 201367 001	6740669	Nov 14, 2022	DS DP			
	6740669*PED	May 14, 2023				
	7750028	Oct 19, 2018		U-106		
	7750028*PED	Apr 19, 2019				
<b>RUXOLITINIB PHOSPHATE - JAKAFI</b>						
N 202192 001	7598257	Dec 24, 2027	DS DP U-1201		I-699	Dec 04, 2017
	7598257	Dec 24, 2027	DS DP U-1622		ODE-19	Nov 16, 2018
	8415362	Dec 24, 2027	DS DP		ODE-79	Dec 04, 2021
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	9079912	Dec 12, 2026		U-1573		
	9079912	Dec 12, 2026		U-1721		
	9662335	Dec 12, 2026	DS			
<b>RUXOLITINIB PHOSPHATE - JAKAFI</b>						
N 202192 002	7598257	Dec 24, 2027	DS DP U-1201		I-699	Dec 04, 2017
	7598257	Dec 24, 2027	DS DP U-1622		ODE-19	Nov 16, 2018
	8415362	Dec 24, 2027	DS DP		ODE-79	Dec 04, 2021
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	9079912	Dec 12, 2026		U-1573		
	9079912	Dec 12, 2026		U-1721		
	9662335	Dec 12, 2026	DS			
<b>RUXOLITINIB PHOSPHATE - JAKAFI</b>						
N 202192 003	7598257	Dec 24, 2027	DS DP U-1201		I-699	Dec 04, 2017
	7598257	Dec 24, 2027	DS DP U-1622		ODE-19	Nov 16, 2018
	8415362	Dec 24, 2027	DS DP		ODE-79	Dec 04, 2021
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	9079912	Dec 12, 2026		U-1573		
	9079912	Dec 12, 2026		U-1721		
	9662335	Dec 12, 2026	DS			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>RUXOLITINIB PHOSPHATE - JAKAFI</b>						
N 202192 004	7598257	Dec 24, 2027	DS DP U-1201		I-699	Dec 04, 2017
	7598257	Dec 24, 2027	DS DP U-1622		ODE-19	Nov 16, 2018
	8415362	Dec 24, 2027	DS DP		ODE-79	Dec 04, 2021
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	9079912	Dec 12, 2026		U-1573		
	9079912	Dec 12, 2026		U-1721		
	9662335	Dec 12, 2026	DS			
<b>RUXOLITINIB PHOSPHATE - JAKAFI</b>						
N 202192 005	7598257	Dec 24, 2027	DS DP U-1201		I-699	Dec 04, 2017
	7598257	Dec 24, 2027	DS DP U-1622		ODE-19	Nov 16, 2018
	8415362	Dec 24, 2027	DS DP		ODE-79	Dec 04, 2021
	8722693	Jun 12, 2028	DS DP			
	8822481	Jun 12, 2028		U-1573		
	8829013	Jun 12, 2028		U-1201		
	8829013	Jun 12, 2028		U-1622		
	9079912	Dec 12, 2026		U-1573		
	9079912	Dec 12, 2026		U-1721		
	9662335	Dec 12, 2026	DS			
<b>SACROSIDASE - SUCRAID</b>						
N 020772 001	9469847	Feb 07, 2034	DS DP			
<b>SACUBITRIL; VALSARTAN - ENTRESTO</b>						
N 207620 001	7468390	Nov 27, 2023	DP		NCE	Jul 07, 2020
	8101659	Jan 14, 2023	DP			
	8404744	Jan 14, 2023	DP			
	8796331	Jan 14, 2023		U-1723		
	8877938	May 27, 2027	DS DP			
	9388134	Nov 08, 2026		U-1723		
<b>SACUBITRIL; VALSARTAN - ENTRESTO</b>						
N 207620 002	7468390	Nov 27, 2023	DP		NCE	Jul 07, 2020
	8101659	Jan 14, 2023	DP			
	8404744	Jan 14, 2023	DP			
	8796331	Jan 14, 2023		U-1723		
	8877938	May 27, 2027	DS DP			
	9388134	Nov 08, 2026		U-1723		
<b>SACUBITRIL; VALSARTAN - ENTRESTO</b>						
N 207620 003	7468390	Nov 27, 2023	DP		NCE	Jul 07, 2020
	8101659	Jan 14, 2023	DP			
	8404744	Jan 14, 2023	DP			
	8796331	Jan 14, 2023		U-1723		
	8877938	May 27, 2027	DS DP			
	9388134	Nov 08, 2026		U-1723		
<b>SAFINAMIDE MESYLATE - XADAGO</b>						
N 207145 001	8076515	Dec 10, 2028	DS DP U-1993		NCE	Mar 21, 2022
	8278485	Jun 08, 2027	DS U-1993			
	8283380	Sep 01, 2027		U-1993		
<b>SAFINAMIDE MESYLATE - XADAGO</b>						
N 207145 002	8076515	Dec 10, 2028	DS DP U-1993		NCE	Mar 21, 2022
	8278485	Jun 08, 2027	DS U-1993			
	8283380	Sep 01, 2027		U-1993		
<b>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</b>						
N 022181 001	7566462	Nov 16, 2025	DP			
	7566462*PED	May 16, 2026				
	7566714	Nov 17, 2024		U-989		
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024		U-1010		
	7612073*PED	May 17, 2025				
	7727987	Nov 17, 2024	DP			
	7727987*PED	May 17, 2025				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</b>						
N 022181 001	7947681	Nov 17, 2024	U-1156	Y		
	7947681*PED	May 17, 2025				
	8003126	Nov 16, 2025				
	8003126*PED	May 16, 2026				
	8067416	Nov 17, 2024	U-989			
	8067416*PED	May 17, 2025				
	8318745	Nov 17, 2024	DP			
	8318745*PED	May 17, 2025				
	9433624	Nov 17, 2024	U-1589			
	RE43797	Nov 17, 2024	U-1156			
	RE43797*PED	May 17, 2025				
<b>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</b>						
N 205065 001	7566714	Nov 17, 2024	U-1589			
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024	U-1010			
	7612073*PED	May 17, 2025				
	8067416	Nov 17, 2024	U-1589			
	8067416*PED	May 17, 2025				
	9216178	Nov 01, 2032	DP			
	9433624	Nov 17, 2024	U-1589			
	RE43797	Nov 17, 2024	U-1590			
	RE43797*PED	May 17, 2025				
<b>SAPROPTERIN DIHYDROCHLORIDE - KUVAN</b>						
N 205065 002	7566714	Nov 17, 2024	U-1589			
	7566714*PED	May 17, 2025				
	7612073	Nov 17, 2024	U-1010			
	7612073*PED	May 17, 2025				
	8067416	Nov 17, 2024	U-1589			
	8067416*PED	May 17, 2025				
	9216178	Nov 01, 2032	DP			
	9433624	Nov 17, 2024	U-1589			
	RE43797	Nov 17, 2024	U-1590			
	RE43797*PED	May 17, 2025				
<b>SAQUINAVIR - FORTOVASE</b>						
N 020828 001	6352717	Nov 16, 2019				
<b>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</b>						
N 022350 001	7951400	Nov 30, 2028	DP		M-175	Apr 05, 2019
	RE44186	Jul 31, 2023	DS DP U-1837		M-198	Feb 27, 2020
	RE44186	Jul 31, 2023	DS DP U-995			
<b>SAXAGLIPTIN HYDROCHLORIDE - ONGLYZA</b>						
N 022350 002	7951400	Nov 30, 2028	DP		M-175	Apr 05, 2019
	RE44186	Jul 31, 2023	DS DP U-1837		M-198	Feb 27, 2020
	RE44186	Jul 31, 2023	DS DP U-995			
<b>SECNIDAZOLE - SOLOSEC</b>						
N 209363 001					NCE	Sep 15, 2022
					GAIN	Sep 15, 2027
<b>SELEGILINE - EMSAM</b>						
N 021336 001	7070808	May 10, 2018	DS DP			
	7150881	Jun 12, 2018	DS DP			
	7638140	May 10, 2018	DP			
<b>SELEGILINE - EMSAM</b>						
N 021336 002	7070808	May 10, 2018	DS DP			
	7150881	Jun 12, 2018	DS DP			
	7638140	May 10, 2018	DP			
<b>SELEGILINE - EMSAM</b>						
N 021336 003	7070808	May 10, 2018	DS DP			
	7150881	Jun 12, 2018	DS DP			
	7638140	May 10, 2018	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 001	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 002	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 003	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 004	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 005	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 006	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 007	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SELEXIPAG - UPTRAVI</u></b>						
N 207947 008	7205302	Apr 04, 2023	DS DP U-1797		NCE	Dec 21, 2020
	8791122	Aug 01, 2030	DS DP		ODE-106	Dec 21, 2022
	9173881	Aug 12, 2029		U-1798		
	9284280	Jun 25, 2030		U-1831		
<b><u>SEMAGLUTIDE - OZEMPIC</u></b>						
N 209637 001	6899699	Jan 02, 2022	DP		NCE	Dec 05, 2022
	7762994	May 23, 2024	DP			
	8114833	Aug 13, 2025	DP			
	8129343	Jan 29, 2029	DS DP U-2202			
	8536122	Mar 20, 2026	DS DP U-2202			
	8579869	Jun 30, 2023	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b><u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u></b>						
N 020990 001	6727283	Oct 11, 2019	DP U-580			
	7067555	Oct 11, 2019	DP			
	7067555*PED	Apr 11, 2020				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>SERTRALINE HYDROCHLORIDE - ZOLOFT</b>						
N 020990 001	6727283	Oct 11, 2019	DP	U-580		
	7067555	Oct 11, 2019	DP			
	7067555*PED	Apr 11, 2020				
<b>SEVELAMER CARBONATE - RENVELA</b>						
N 022127 001	7985418	Oct 27, 2025	DP		NPP	Nov 25, 2019
<b>SEVELAMER CARBONATE - RENVELA</b>						
N 022318 001	9095509	Dec 06, 2030	DP		NPP	Nov 25, 2019
<b>SEVELAMER CARBONATE - RENVELA</b>						
N 022318 002	9095509	Dec 06, 2030	DP		NPP	Nov 25, 2019
<b>SEVELAMER HYDROCHLORIDE - RENAGEL</b>						
N 021179 001	6733780	Oct 18, 2020	DP			
<b>SEVELAMER HYDROCHLORIDE - RENAGEL</b>						
N 021179 002	6733780	Oct 18, 2020	DP			
<b>SEVOFLURANE - ULTANE</b>						
N 020478 001	6074668	Jan 09, 2018				
<b>SILDENAFIL CITRATE - SILDENAFIL CITRATE</b>						
A 077342 001					PC	Jun 09, 2018
<b>SILDENAFIL CITRATE - SILDENAFIL CITRATE</b>						
A 077342 002					PC	Jun 09, 2018
<b>SILDENAFIL CITRATE - SILDENAFIL CITRATE</b>						
A 077342 003					PC	Jun 09, 2018
<b>SILDENAFIL CITRATE - VIAGRA</b>						
N 020895 001	6469012	Oct 22, 2019		U-155		
<b>SILDENAFIL CITRATE - VIAGRA</b>						
N 020895 002	6469012	Oct 22, 2019		U-155		
<b>SILDENAFIL CITRATE - VIAGRA</b>						
N 020895 003	6469012	Oct 22, 2019		U-155		
<b>SILODOSIN - RAPAFLO</b>						
N 022206 001	5387603	Dec 01, 2018	DS DP			
<b>SILODOSIN - RAPAFLO</b>						
N 022206 002	5387603	Dec 01, 2018	DS DP			
<b>SIMEPREVIR SODIUM - OLYSIO</b>						
N 205123 001	7671032	May 19, 2025	DS DP		D-151	Oct 05, 2018
	8148399	Sep 05, 2029	DS DP	U-1467	I-697	Nov 05, 2017
	8349869	Jul 28, 2026	DS DP	U-1467	I-717	Oct 05, 2018
	8741926	Jul 28, 2026	DS	U-1467	M-171	Feb 26, 2019
	8754106	Jul 28, 2026	DS	U-1467	M-179	May 20, 2019
	9040562	Jul 28, 2026	DS DP	U-1467	NCE	Nov 22, 2018
	9353103	Jul 28, 2026		U-1467		
	9623022	Jul 28, 2026		U-1467		
<b>SIMVASTATIN - FLOLIPID</b>						
N 206679 001	9597289	Feb 23, 2030	DP			
<b>SIMVASTATIN - FLOLIPID</b>						
N 206679 002	9597289	Feb 23, 2030	DP			
<b>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</b>						
N 202343 001	6699871	Jul 26, 2022	DS DP	U-1188		
	6890898	Feb 02, 2019		U-1189		
	6890898	Feb 02, 2019		U-1190		
	6890898	Feb 02, 2019		U-1191		
	7078381	Feb 02, 2019		U-1188		
	7125873	Jul 26, 2022	DP	U-1189		
	7125873	Jul 26, 2022	DP	U-1190		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 001	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	7459428	Feb 02, 2019		U-1189		
	8168637	Jun 26, 2022	DP U-1188			
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 002	6699871	Jul 26, 2022	DS DP U-1188			
	6890898	Feb 02, 2019	U-1189			
	6890898	Feb 02, 2019	U-1190			
	6890898	Feb 02, 2019	U-1191			
	7078381	Feb 02, 2019	U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	7459428	Feb 02, 2019		U-1189		
	8168637	Jun 26, 2022	DP U-1188			
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 003	6699871	Jul 26, 2022	DS DP U-1188			
	6890898	Feb 02, 2019	U-1189			
	6890898	Feb 02, 2019	U-1190			
	6890898	Feb 02, 2019	U-1191			
	7078381	Feb 02, 2019	U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	7459428	Feb 02, 2019		U-1189		
	8168637	Jun 26, 2022	DP U-1188			
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 004	6699871	Jul 26, 2022	DS DP U-1188			
	6890898	Feb 02, 2019	U-1189			
	6890898	Feb 02, 2019	U-1190			
	6890898	Feb 02, 2019	U-1191			
	7078381	Feb 02, 2019	U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	7459428	Feb 02, 2019		U-1189		
	8168637	Jun 26, 2022	DP U-1188			
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 005	6699871	Jul 26, 2022	DS DP U-1188			
	6890898	Feb 02, 2019	U-1189			
	6890898	Feb 02, 2019	U-1190			
	6890898	Feb 02, 2019	U-1191			
	7078381	Feb 02, 2019	U-1188			
	7125873	Jul 26, 2022	DP U-1189			
	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	7459428	Feb 02, 2019		U-1189		
	8168637	Jun 26, 2022	DP U-1188			
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 006	6699871	Jul 26, 2022	DS DP U-1188			
	6890898	Feb 02, 2019	U-1189			
	6890898	Feb 02, 2019	U-1190			
	6890898	Feb 02, 2019	U-1191			
	7078381	Feb 02, 2019	U-1188			
	7125873	Jul 26, 2022	DP U-1189			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>SIMVASTATIN; SITAGLIPTIN PHOSPHATE - JUVISYNC</u></b>						
N 202343 006	7125873	Jul 26, 2022	DP U-1190			
	7125873	Jul 26, 2022	DP U-1192			
	7125873	Jul 26, 2022	DP U-1193			
	7326708	Apr 11, 2026	DS DP U-1188			
	7459428	Feb 02, 2019		U-1189		
	8168637	Jun 26, 2022	DP	U-1188		
<b><u>SINCALIDE - KINEVAC</u></b>						
N 017697 001	6803046	Aug 16, 2022	DP			
<b><u>SINECATECHINS - VEREGEN</u></b>						
N 021902 001	5795911	Oct 31, 2020	U-172			
	7858662	Oct 02, 2026	DP U-172			
	9770406	Jul 12, 2025	DP U-172			
<b><u>SIROLIMUS - RAPAMUNE</u></b>						
N 021083 001					ODE-92	May 28, 2022
<b><u>SIROLIMUS - RAPAMUNE</u></b>						
N 021110 001	5989591	Mar 11, 2018	DP		ODE-92	May 28, 2022
<b><u>SIROLIMUS - RAPAMUNE</u></b>						
N 021110 002	5989591	Mar 11, 2018	DP		ODE-92	May 28, 2022
<b><u>SIROLIMUS - RAPAMUNE</u></b>						
N 021110 003	5989591	Mar 11, 2018	DP		ODE-92	May 28, 2022
<b><u>SIROLIMUS - RAPAMUNE</u></b>						
N 021110 004	5989591	Mar 11, 2018	DP		ODE-92	May 28, 2022
<b><u>SITAGLIPTIN PHOSPHATE - JANUVIA</u></b>						
N 021995 001	6699871	Jul 26, 2022	DS DP U-774			
	6890898	Feb 02, 2019	U-1997			
	7078381	Feb 02, 2019	U-1997			
	7125873	Jul 26, 2022	U-1036			
	7125873	Jul 26, 2022	U-1037			
	7125873	Jul 26, 2022	U-1038			
	7125873	Jul 26, 2022	U-775			
	7326708	Nov 24, 2026	DS DP U-802			
	7459428	Feb 02, 2019	U-1945			
<b><u>SITAGLIPTIN PHOSPHATE - JANUVIA</u></b>						
N 021995 002	6699871	Jul 26, 2022	DS DP U-774			
	6890898	Feb 02, 2019	U-1997			
	7078381	Feb 02, 2019	U-1997			
	7125873	Jul 26, 2022	U-1036			
	7125873	Jul 26, 2022	U-1037			
	7125873	Jul 26, 2022	U-1038			
	7125873	Jul 26, 2022	U-775			
	7326708	Nov 24, 2026	DS DP U-802			
	7459428	Feb 02, 2019	U-1945			
<b><u>SITAGLIPTIN PHOSPHATE - JANUVIA</u></b>						
N 021995 003	6699871	Jul 26, 2022	DS DP U-774			
	6890898	Feb 02, 2019	U-1997			
	7078381	Feb 02, 2019	U-1997			
	7125873	Jul 26, 2022	U-1036			
	7125873	Jul 26, 2022	U-1037			
	7125873	Jul 26, 2022	U-1038			
	7125873	Jul 26, 2022	U-775			
	7326708	Nov 24, 2026	DS DP U-802			
	7459428	Feb 02, 2019	U-1945			
<b><u>SODIUM NITRITE - SODIUM NITRITE</u></b>						
N 203922 001	8568793	Dec 24, 2031	DS DP		ODE-5	Jan 14, 2018
<b><u>SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOTE</u></b>						
N 201444 001	8496973	Mar 29, 2031	DS DP U-1419		ODE-5	Jan 14, 2018
	8568793	Dec 24, 2031	DS DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>SODIUM NITRITE; SODIUM THIOSULFATE - NITHIODOTE</b>						
N 201444 001	9345724	Mar 29, 2031	DS DP U-2015			
	9585912	Mar 29, 2031	DS DP			
<b>SODIUM OXYBATE - XYREM</b>						
N 021196 001	6780889	Jul 04, 2020	DP			
	7262219	Jul 04, 2020	DP			
	7668730	Jun 16, 2024	U-1110			
	7765106	Jun 16, 2024	U-1069			
	7765107	Jun 16, 2024	U-1070			
	7851506	Dec 22, 2019	U-1101			
	7851506	Dec 22, 2019	U-1102			
	7895059	Dec 17, 2022	U-1110			
	8263650	Dec 22, 2019	DP U-1101			
	8263650	Dec 22, 2019	DP U-1102			
	8324275	Dec 22, 2019	U-1101			
	8324275	Dec 22, 2019	U-1102			
	8457988	Dec 17, 2022	U-1110			
	8589182	Dec 17, 2022	U-1110			
	8731963	Dec 17, 2022	U-1110			
	8772306	Mar 15, 2033	U-1532			
	8859619	Dec 22, 2019	DP			
	8952062	Dec 22, 2019	U-1101			
	8952062	Dec 22, 2019	U-1102			
	9050302	Mar 15, 2033	U-1532			
	9486426	Mar 15, 2033	U-1532			
	9539330	Dec 22, 2019	DP			
<b>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</b>						
N 021892 001	7687075	Jun 22, 2028	DS DP			
<b>SODIUM THIOSULFATE - SODIUM THIOSULFATE</b>						
N 203923 001	8496973	Mar 29, 2031	DS DP U-1419		ODE-5	
	9345724	Mar 29, 2031	DS DP U-2015			
	9585912	Mar 29, 2031	DS DP			
<b>SOFOSBUVIR - SOVALDI</b>						
N 204671 001	7964580	Mar 26, 2029	DS DP U-1470		NCE	Dec 06, 2018
	8334270	Mar 21, 2028	DS DP U-1470		NPP	Apr 07, 2020
	8580765	Mar 21, 2028	DS DP U-1470		ODE-135	Apr 07, 2024
	8618076	Dec 11, 2030	DS DP U-1470			
	8633309	Mar 26, 2029	DS DP U-1470			
	8889159	Mar 26, 2029	DP U-1470			
	9085573	Mar 21, 2028	DS DP U-1470			
	9284342	Sep 13, 2030	DS DP U-1470			
	9549941	Mar 26, 2029	U-1958			
<b>SOFOSBUVIR; VELPATASVIR - EPCLUSA</b>						
N 208341 001	7964580	Mar 26, 2029	DS DP U-1470		NCE	Dec 06, 2018
	8334270	Mar 21, 2028	DS DP U-1470		NCE	Jun 28, 2021
	8575135	Nov 16, 2032	DS DP U-1470		NPP	Aug 01, 2020
	8580765	Mar 21, 2028	DS DP U-1470			
	8618076	Dec 11, 2030	DS DP U-1470			
	8633309	Mar 26, 2029	DS DP U-1470			
	8735372	Mar 21, 2028	U-1470			
	8889159	Mar 26, 2029	DP U-1470			
	8921341	Nov 16, 2032	DS DP U-1470			
	8940718	Nov 16, 2032	DS DP U-1470			
	9085573	Mar 21, 2028	DS DP U-1470			
	9284342	Sep 13, 2030	DS DP U-1470			
	9757406	Jan 30, 2034	DP			
<b>SOFOSBUVIR; VELPATASVIR; VOXILAPREVIR - VOSEVI</b>						
N 209195 001	7964580	Mar 26, 2029	DS DP U-2039		NCE	
	7964580	Mar 26, 2029	DS DP U-2040			
	8334270	Mar 21, 2028	DS DP U-2039			
	8334270	Mar 21, 2028	DS DP U-2040			
	8575135	Nov 05, 2033	DS DP U-2039			
	8575135	Nov 05, 2033	DS DP U-2040			
	8580765	Mar 21, 2028	DS DP U-2039			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>SOFOSBUVIR; VELFATASVIR; VOXILAPREVIR - VOSEVI</b>						
N 209195 001	8580765	Mar 21, 2028	DS DP U-2040			
	8618076	Dec 11, 2030	DS DP U-2039			
	8618076	Dec 11, 2030	DS DP U-2040			
	8633309	Mar 26, 2029	DS DP U-2039			
	8633309	Mar 26, 2029	DS DP U-2040			
	8735372	Mar 21, 2028	DS DP U-2039			
	8735372	Mar 21, 2028	DS DP U-2040			
	8889159	Mar 26, 2029	DS DP U-2039			
	8889159	Mar 26, 2029	DS DP U-2040			
	8921341	Nov 16, 2032	DS DP U-2039			
	8921341	Nov 16, 2032	DS DP U-2040			
	8940718	Nov 16, 2032	DS DP U-2039			
	8940718	Nov 16, 2032	DS DP U-2040			
	9085573	Mar 21, 2028	DS DP U-2039			
	9085573	Mar 21, 2028	DS DP U-2040			
	9284342	Sep 13, 2030	DS DP U-2039			
	9284342	Sep 13, 2030	DS DP U-2040			
	9296782	Jul 17, 2034	DS DP			
	9585906	Mar 21, 2028	DS DP U-2039			
	9585906	Mar 21, 2028	DS DP U-2040			
<b>SOLIFENACIN SUCCINATE - VESICARE</b>						
N 021518 001	6017927	Nov 19, 2018	DS DP			
	6017927*PED	May 19, 2019				
<b>SOLIFENACIN SUCCINATE - VESICARE</b>						
N 021518 002	6017927	Nov 19, 2018	DS DP			
	6017927*PED	May 19, 2019				
<b>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</b>						
N 020280 001	6152897	Nov 20, 2018	DP			
<b>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</b>						
N 020280 002	6152897	Nov 20, 2018	DP			
<b>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</b>						
N 020280 003	6152897	Nov 20, 2018	DP			
<b>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</b>						
N 020280 005	6152897	Nov 20, 2018	DP			
<b>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</b>						
N 020280 008	6152897	Nov 20, 2018	DP			
<b>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</b>						
N 020280 009	6152897	Nov 20, 2018	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</b>						
N 021148 004	6004297	Jan 28, 2019	DP			
	6235004	Jan 28, 2019	DP			
	RE41956	Jan 21, 2021	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</b>						
N 021148 005	6004297	Jan 28, 2019	DP			
	6235004	Jan 28, 2019	DP			
	RE41956	Jan 21, 2021	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</b>						
N 021148 006	6004297	Jan 28, 2019	DP			
	6235004	Jan 28, 2019	DP			
	RE41956	Jan 21, 2021	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN NORDIFLEX</b>						
N 021148 007	6004297	Jan 28, 2019	DP			
	8841252	Dec 26, 2017	DP			
	RE41956	Jan 21, 2021	DP			
	RE43834	Jan 28, 2019	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</b>						
N 021148 008	6899699	Jan 02, 2022	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8841252	Dec 26, 2017	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</b>						
N 021148 009	6899699	Jan 02, 2022	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8841252	Dec 26, 2017	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</b>						
N 021148 010	6899699	Jan 02, 2022	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8841252	Dec 26, 2017	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b>SOMATROPIN RECOMBINANT - NORDITROPIN FLEXPRO</b>						
N 021148 011	6899699	Jan 02, 2022	DP			
	8672898	Jan 02, 2022	DP			
	8684969	Oct 20, 2025	DP			
	8841252	Dec 26, 2017	DP			
	8920383	Jul 17, 2026	DP			
	9108002	Jan 20, 2026	DP			
	9132239	Feb 01, 2032	DP			
	9457154	Sep 27, 2027	DP			
	9486588	Jan 02, 2022	DP			
	9687611	Feb 27, 2027	DP			
	9775953	Jul 17, 2026	DP			
	RE46363	Aug 03, 2026	DP			
<b>SONIDEGRIB PHOSPHATE - ODOMZO</b>						
N 205266 001	8063043	Sep 15, 2029	DS DP		NCE	Jul 24, 2020
	8178563	Feb 06, 2029	DS U-1722			
<b>SORAFENIB TOSYLATE - NEXAVAR</b>						
N 021923 001	7235576	Jan 12, 2020	DS DP		ODE-56	Nov 22, 2020
	7351834	Jan 12, 2020	DS			
	7897623	Jan 12, 2020	DP			
	8124630	Jan 12, 2020	U-1459			
	8618141	Feb 11, 2023	U-1480			
	8841330	Jan 12, 2020	U-1696			
	8877933	Dec 24, 2027	DS DP U-1624			
	9737488	Sep 10, 2028	DP U-1480			
	9737488	Sep 10, 2028	DP U-1696			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>SORAFENIB TOSYLATE - NEXAVAR</u></b>						
N 021923 001	9737488	Sep 10, 2028	DP U-2107			
<b><u>SOTALOL HYDROCHLORIDE - SOTYLIZE</u></b>						
N 205108 001	9724297	Aug 31, 2035	DP U-2096			
<b><u>SPINOSAD - NATROBA</u></b>						
N 022408 001	6063771	Jul 25, 2023	DP U-1670		M-152	
	6342482	Jun 22, 2019	DP U-1105			
	7030095	Jul 02, 2021	DP U-1105			
<b><u>SPIRONOLACTONE - CAROSPIR</u></b>						
N 209478 001	9757394	Oct 28, 2036	DP U-2109			
<b><u>STAVUDINE - ZERIT XR</u></b>						
N 021453 001	7135465	Feb 18, 2023	DP U-167			
<b><u>STAVUDINE - ZERIT XR</u></b>						
N 021453 002	7135465	Feb 18, 2023	DP U-167			
<b><u>STAVUDINE - ZERIT XR</u></b>						
N 021453 003	7135465	Feb 18, 2023	DP U-167			
<b><u>STAVUDINE - ZERIT XR</u></b>						
N 021453 004	7135465	Feb 18, 2023	DP U-167			
<b><u>SUCROFERRIC OXYHYDROXIDE - VELPHORO</u></b>						
N 205109 001	6174442	Dec 19, 2018	DS U-1468			
	9561251	Jan 23, 2030	DP U-1468			
<b><u>SUGAMMADEX SODIUM - BRIDION</u></b>						
N 022225 001	6949527	Jan 27, 2021	U-1795		NCE	
	7265099	Aug 07, 2020	U-1795			
	RE44733	Jan 27, 2021	DS DP U-1794			
<b><u>SUGAMMADEX SODIUM - BRIDION</u></b>						
N 022225 002	6949527	Jan 27, 2021	U-1795		NCE	
	7265099	Aug 07, 2020	U-1795			
	RE44733	Jan 27, 2021	DS DP U-1794			
<b><u>SULFUR HEXAFLUORIDE LIPID-TYPE A MICROSPHERES - LUMASON</u></b>						
N 203684 001	5686060	Nov 11, 2019	DS DP		I-728	
					NCE	
						Mar 31, 2019
						Oct 10, 2019
<b><u>SUMATRIPTAN SUCCINATE - SUMAVENT DOSEPRO</u></b>						
N 022239 001	7776007	Nov 22, 2026	DP			
	7901385	Jul 31, 2026	DP			
	8118771	Aug 10, 2023	DP			
	8241243	Aug 10, 2023	DP			
	8241244	Nov 21, 2022	DP			
	8267903	Mar 18, 2023	DP			
	8287489	Dec 06, 2024	DP			
	8343130	Oct 18, 2022	DP			
	8491524	Nov 21, 2022	DP			
<b><u>SUMATRIPTAN SUCCINATE - ALSUMA</u></b>						
N 022377 001	7811254	Aug 26, 2027	DP U-1083			
<b><u>SUMATRIPTAN SUCCINATE - ZECURITY</u></b>						
N 202278 001	6745071	Feb 21, 2023	DP			
	7973058	Apr 12, 2027	U-1328			
	8155737	Apr 12, 2027	U-1328			
	8366600	Apr 21, 2029	U-1327			
	8470853	Apr 12, 2027	U-1328			
	8597272	Apr 12, 2027	DP			
	8983594	Nov 19, 2030	DP U-1328			
	9272137	Sep 07, 2027	DP			
	9327114	Oct 08, 2032	DP U-1328			
	9427578	Apr 12, 2027	DP U-1328			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>SUMATRIPTAN SUCCINATE - ONZETRA XSAIL</b>						
N 206099 001	6715485	Mar 03, 2020	DP			
	7975690	Aug 18, 2025	DP U-1809			
	8047202	Jul 02, 2023	DP			
	8327844	Oct 03, 2023	U-1809			
	8550073	Oct 22, 2029	DP			
	8555877	Mar 03, 2020	DP			
	8590530	Sep 15, 2025	DP U-1809			
	8875704	Apr 07, 2028	DP U-1809			
	8899229	Aug 18, 2030	DP			
	8978647	Dec 06, 2030	DP			
	9108015	Sep 15, 2025	DP			
	9119932	Apr 23, 2024	DP			
	9649456	Oct 21, 2030	DP U-1719			
	9649456	Oct 21, 2030	DP U-2010			
	9649456	Oct 21, 2030	DP U-2011			
<b>SUNITINIB MALATE - SUTENT</b>						
N 021938 001	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171			
	7125905	Feb 15, 2021	DS DP			
	7211600	Dec 22, 2020	U-883			
<b>SUNITINIB MALATE - SUTENT</b>						
N 021938 002	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171			
	7125905	Feb 15, 2021	DS DP			
	7211600	Dec 22, 2020	U-883			
<b>SUNITINIB MALATE - SUTENT</b>						
N 021938 003	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171			
	7125905	Feb 15, 2021	DS DP			
	7211600	Dec 22, 2020	U-883			
<b>SUNITINIB MALATE - SUTENT</b>						
N 021938 004	6573293	Feb 15, 2021	DS DP U-1154		I-755	Nov 16, 2020
	6573293	Feb 15, 2021	DS DP U-2171			
	7125905	Feb 15, 2021	DS DP			
	7211600	Dec 22, 2020	U-883			
<b>SUVOREXANT - BELSOMRA</b>						
N 204569 001	7951797	Nov 20, 2029	DS DP U-620		NCE	Aug 13, 2019
<b>SUVOREXANT - BELSOMRA</b>						
N 204569 002	7951797	Nov 20, 2029	DS DP U-620		NCE	Aug 13, 2019
<b>SUVOREXANT - BELSOMRA</b>						
N 204569 003	7951797	Nov 20, 2029	DS DP U-620		NCE	Aug 13, 2019
<b>SUVOREXANT - BELSOMRA</b>						
N 204569 004	7951797	Nov 20, 2029	DS DP U-620		NCE	Aug 13, 2019
<b>TACROLIMUS - ASTAGRAF XL</b>						
N 204096 001	6440458	Mar 25, 2019	DP			
	6576259	Mar 25, 2019	DP U-1420			
	6884433	Mar 25, 2019	DP U-1420			
	8551522	Mar 25, 2019	DP			
<b>TACROLIMUS - ASTAGRAF XL</b>						
N 204096 002	6440458	Mar 25, 2019	DP			
	6576259	Mar 25, 2019	DP U-1420			
	6884433	Mar 25, 2019	DP U-1420			
	8551522	Mar 25, 2019	DP			
<b>TACROLIMUS - ASTAGRAF XL</b>						
N 204096 003	6440458	Mar 25, 2019	DP			
	6576259	Mar 25, 2019	DP U-1420			
	6884433	Mar 25, 2019	DP U-1420			
	8551522	Mar 25, 2019	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>TACROLIMUS - ENVARSUS XR</b>						
N 206406 001	7994214	Aug 30, 2024	DP		ODE-94	Jul 10, 2022
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	8664239	May 30, 2028	U-1752			
	8685998	May 30, 2028	DP U-1752			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<b>TACROLIMUS - ENVARSUS XR</b>						
N 206406 002	7994214	Aug 30, 2024	DP		ODE-94	Jul 10, 2022
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	8664239	May 30, 2028	U-1752			
	8685998	May 30, 2028	DP U-1752			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<b>TACROLIMUS - ENVARSUS XR</b>						
N 206406 003	7994214	Aug 30, 2024	DP		ODE-94	Jul 10, 2022
	8486993	Aug 30, 2024	DP U-1752			
	8586084	Aug 30, 2024	U-1752			
	8591946	Aug 30, 2024	DP			
	8617599	Aug 30, 2024	DP			
	8623410	Aug 30, 2024	DP			
	8623411	Aug 30, 2024	U-1752			
	8664239	May 30, 2028	U-1752			
	8685998	May 30, 2028	DP U-1752			
	8889185	Aug 30, 2024	U-1752			
	8889186	Aug 30, 2024	U-1752			
	9161907	Aug 30, 2024	DP U-1752			
	9549918	May 30, 2028	DP			
	9757362	Aug 30, 2024	DP			
	9763920	Aug 30, 2024	DP			
<b>TADALAFIL - CIALIS</b>						
N 021368 001	5859006	Nov 21, 2017	DS DP		Y	
	5859006*PED	May 21, 2018				
	6821975	Nov 19, 2020	DS DP U-1184			
	6821975	Nov 19, 2020	DS DP U-533			
	6821975	Nov 19, 2020	DS DP U-614			
	6821975*PED	May 19, 2021				
	6943166	Apr 26, 2020	U-1184			
	6943166	Apr 26, 2020	U-155			
	6943166	Apr 26, 2020	U-614			
	6943166*PED	Oct 26, 2020				
	7182958	Apr 26, 2020	DP U-1184			
	7182958	Apr 26, 2020	DP U-155			
	7182958*PED	Oct 26, 2020				
<b>TADALAFIL - CIALIS</b>						
N 021368 002	5859006	Nov 21, 2017	DS DP			
	5859006*PED	May 21, 2018				
	6821975	Nov 19, 2020	DS DP U-533			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TADALAFIL - CIALIS</u></b>						
N 021368 002	6821975	Nov 19, 2020	DS DP U-614			
	6821975*PED	May 19, 2021				
	6943166	Apr 26, 2020		U-155		
	6943166	Apr 26, 2020		U-614		
	6943166*PED	Oct 26, 2020				
	7182958	Apr 26, 2020		DP U-155		
	7182958*PED	Oct 26, 2020				
<b><u>TADALAFIL - CIALIS</u></b>						
N 021368 003	5859006	Nov 21, 2017	DS DP			
	5859006*PED	May 21, 2018				
	6821975	Nov 19, 2020	DS DP U-533			
	6821975	Nov 19, 2020	DS DP U-614			
	6821975*PED	May 19, 2021				
	6943166	Apr 26, 2020		U-614		
	6943166*PED	Oct 26, 2020				
	7182958	Apr 26, 2020		DP U-155		
	7182958*PED	Oct 26, 2020				
<b><u>TADALAFIL - CIALIS</u></b>						
N 021368 004	5859006	Nov 21, 2017	DS DP			
	5859006*PED	May 21, 2018				
	6821975	Nov 19, 2020	DS DP U-533			
	6821975	Nov 19, 2020	DS DP U-614			
	6821975*PED	May 19, 2021				
	6943166	Apr 26, 2020		U-155		
	6943166*PED	Oct 26, 2020				
	7182958	Apr 26, 2020		DP U-155		
	7182958*PED	Oct 26, 2020				
<b><u>TADALAFIL - ADCIRCA</u></b>						
N 022332 001	5859006	Nov 21, 2017	DS DP U-975			
	5859006*PED	May 21, 2018				
	6821975	Nov 19, 2020	DS DP			
	6821975*PED	May 19, 2021				
	7182958	Apr 26, 2020		DP		
	7182958*PED	Oct 26, 2020				
<b><u>TAFLUPROST - ZIOPTAN</u></b>						
N 202514 001	5886035	Dec 18, 2022	DS DP U-778			
<b><u>TALC - STERITALC</u></b>						
N 205555 001					ODE-143	May 01, 2024
<b><u>TALC - STERITALC</u></b>						
N 205555 002					ODE-143	May 01, 2024
<b><u>TALC - STERITALC</u></b>						
N 205555 003					ODE-143	May 01, 2024
<b><u>TALIGLUCERASE ALFA - ELELYSO</u></b>						
N 022458 001	8227230	Feb 24, 2024	DS DP		NPP	Aug 27, 2017
	8741620	Feb 24, 2024	DS DP			
	8790641	Oct 18, 2025		U-1564		
	8790641	Oct 18, 2025		U-1574		
<b><u>TAMOXIFEN CITRATE - SOLTAMOX</u></b>						
N 021807 001	6127425	Jun 26, 2018	DP			
<b><u>TAPENTadol HYDROCHLORIDE - NUCYNTA</u></b>						
N 022304 001	7994364	Jun 27, 2025	DS DP U-931			
	RE39593	Aug 05, 2022	DS DP U-931			
<b><u>TAPENTadol HYDROCHLORIDE - NUCYNTA</u></b>						
N 022304 002	7994364	Jun 27, 2025	DS DP U-931			
	RE39593	Aug 05, 2022	DS DP U-931			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>TAPENTADOL HYDROCHLORIDE - NUCYNTA</b>						
N 022304 003	7994364	Jun 27, 2025	DS DP U-931			
	RE39593	Aug 05, 2022	DS DP U-931			
<b>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</b>						
N 200533 001	7994364	Jun 27, 2025	DS DP U-1178			
	7994364	Jun 27, 2025	DS DP U-1276			
	8075872	Nov 20, 2023	DP			
	8114383	Oct 10, 2024	DP		Y	
	8309060	Nov 20, 2023	DP U-1178			
	8309060	Nov 20, 2023	DP U-1276			
	8420056	Nov 20, 2023	DP			
	8536130	Sep 22, 2028	U-1276			
	RE39593	Aug 05, 2022	DS DP U-1178			
	RE39593	Aug 05, 2022	DS DP U-1276			
<b>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</b>						
N 200533 002	7994364	Jun 27, 2025	DS DP U-1178			
	7994364	Jun 27, 2025	DS DP U-1276			
	8075872	Nov 20, 2023	DP			
	8114383	Oct 10, 2024	DP		Y	
	8309060	Nov 20, 2023	DP U-1178			
	8309060	Nov 20, 2023	DP U-1276			
	8420056	Nov 20, 2023	DP			
	8536130	Sep 22, 2028	U-1276			
	RE39593	Aug 05, 2022	DS DP U-1178			
	RE39593	Aug 05, 2022	DS DP U-1276			
<b>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</b>						
N 200533 003	7994364	Jun 27, 2025	DS DP U-1178			
	7994364	Jun 27, 2025	DS DP U-1276			
	8075872	Nov 20, 2023	DP			
	8114383	Oct 10, 2024	DP		Y	
	8309060	Nov 20, 2023	DP U-1178			
	8309060	Nov 20, 2023	DP U-1276			
	8420056	Nov 20, 2023	DP			
	8536130	Sep 22, 2028	U-1276			
	RE39593	Aug 05, 2022	DS DP U-1178			
	RE39593	Aug 05, 2022	DS DP U-1276			
<b>TAPENTADOL HYDROCHLORIDE - NUCYNTA ER</b>						
N 200533 004	7994364	Jun 27, 2025	DS DP U-1178			
	7994364	Jun 27, 2025	DS DP U-1276			
	8075872	Nov 20, 2023	DP			
	8114383	Oct 10, 2024	DP		Y	
	8309060	Nov 20, 2023	DP U-1178			
	8309060	Nov 20, 2023	DP U-1276			
	8420056	Nov 20, 2023	DP			
	8536130	Sep 22, 2028	U-1276			
	RE39593	Aug 05, 2022	DS DP U-1178			
	RE39593	Aug 05, 2022	DS DP U-1276			
<b>TAPENTADOL HYDROCHLORIDE - NUCYNTA</b>						
N 203794 001	7994364	Jun 27, 2025	DS DP U-1289			
	RE39593	Aug 05, 2022	DS DP U-1289			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>TASIMELTEON - HETLIOZ</b>						
N 205677 001	5856529	Dec 09, 2018	DS DP U-2149		NCE	Jan 31, 2019
	9060995	Jan 25, 2033	U-1710		ODE-59	Jan 31, 2021
	9539234	Jan 25, 2033	U-1934			
	9549913	Jan 25, 2033	U-1486			
	9730910	May 17, 2034	U-2085			
	RE46604	Jan 25, 2033	U-2147			
<b>TAVABOROLE - KERYDIN</b>						
N 204427 001	7582621	May 26, 2027	U-2016		NCE	Jul 07, 2019
	9549938	Feb 16, 2026	U-1951			
	9566289	Feb 16, 2026	DP			
	9566290	Feb 16, 2026	U-1970			
	9572823	Feb 16, 2026	U-1970			
<b>TAZAROTENE - FABIOR</b>						
N 202428 001	8808716	Feb 24, 2030	DP			
<b>TECHNETIUM TC-99M SULFUR COLLOID KIT - AN-SULFUR COLLOID</b>						
N 017858 001					ODE-29	Aug 13, 2019
<b>TECHNETIUM TC-99M TEBOROXIME KIT - CARDIOTEC</b>						
N 019928 001	6056941	Jul 28, 2019	DP			
<b>TECHNETIUM TC-99M TETROFOSMIN KIT - MYOVIEW 30ML</b>						
N 020372 002	9549999	Mar 10, 2030	DP			
<b>TECHNETIUM TC-99M TILMANOCEPT - LYMPHOSEEK KIT</b>						
N 202207 001	6409990	May 12, 2020	DS		NCE	Mar 13, 2018
	9439985	Sep 27, 2033	DS DP		ODE-67	Jun 13, 2021
<b>TEDIZOLID PHOSPHATE - SIVEXTRO</b>						
N 205435 001	7816379	Feb 23, 2028	DS DP U-282		NCE	Jun 20, 2019
	8420676	Feb 23, 2028	DS DP U-282		GAIN	Jun 20, 2024
	8426389	Dec 31, 2030	DS DP U-282			
<b>TEDIZOLID PHOSPHATE - SIVEXTRO</b>						
N 205436 001	7816379	Feb 23, 2028	DS DP U-282		NCE	Jun 20, 2019
	8420676	Feb 23, 2028	DS DP U-282		GAIN	Jun 20, 2024
	8426389	Dec 31, 2030	DS DP U-282			
<b>TEDUGLUTIDE RECOMBINANT - GATTEX KIT</b>						
N 203441 001	5789379	Apr 14, 2020	DS DP U-1320		NCE	Dec 21, 2017
	7056886	Sep 18, 2022	DP U-1320		ODE-37	Dec 21, 2019
	7847061	Nov 01, 2025	U-1320			
	9060992	Nov 01, 2025	U-1320			
	9539310	Nov 01, 2025	U-1320			
	9545434	Nov 01, 2025	U-1320			
	9545435	Nov 01, 2025	U-1320			
	9555079	Nov 01, 2025	U-1320			
	9572867	Nov 01, 2025	U-1320			
	9592273	Nov 01, 2025	U-1320			
	9592274	Nov 01, 2025	U-1320			
<b>TELAPREVIR - INCIVEK</b>						
N 201917 001	7820671	Feb 25, 2025	DS DP			
	8431615	May 30, 2028	U-1398			
	8529882	Aug 31, 2021	U-1398			
<b>TELAVANCIN HYDROCHLORIDE - VIBATIV</b>						
N 022110 001	6635618	Sep 11, 2023	DS DP U-728			
	6858584	Aug 24, 2022	DP			
	6872701	Jun 05, 2021	DP			
	7008923	May 06, 2021	U-1005			
	7208471	May 01, 2021	DS DP			
	7351691	May 01, 2021	DS DP U-728			
	7531623	Jan 01, 2027	DS			
	7544364	May 01, 2021	DP			
	7700550	May 01, 2021	U-282			
	8101575	May 01, 2021	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u></b>						
N 022110 001	8158580	May 01, 2021	DP			
<b><u>TELAVANCIN HYDROCHLORIDE - VIBATIV</u></b>						
N 022110 002	6635618	Sep 11, 2023	DS DP U-728			
	6858584	Aug 24, 2022	DP			
	6872701	Jun 05, 2021	DP			
	7008923	May 06, 2021	U-1005			
	7208471	May 01, 2021	DS DP			
	7351691	May 01, 2021	DS DP U-728			
	7531623	Jan 01, 2027	DS			
	7544364	May 01, 2021	DP			
	7700550	May 01, 2021	U-282			
	8101575	May 01, 2021	DP			
	8158580	May 01, 2021	DP			
<b><u>TELBIVUDINE - TYZEKA</u></b>						
N 022011 001	6395716	Aug 10, 2019	U-782			
	6444652	Aug 10, 2019	U-782			
	6566344	Aug 10, 2019	U-782			
	6569837	Oct 25, 2020	U-782			
	6569837	Oct 25, 2020	U-999			
	7589079	Sep 11, 2023	DS DP U-999			
	7795238	Aug 10, 2019	U-999			
	7858594	Sep 11, 2023	DS DP U-999			
<b><u>TELBIVUDINE - TYZEKA</u></b>						
N 022154 001	6395716	Aug 10, 2019	U-999			
	6444652	Aug 10, 2019	U-999			
	6566344	Aug 10, 2019	U-999			
	6569837	Oct 25, 2020	U-999			
	7795238	Aug 10, 2019	U-999			
	7858594	Sep 11, 2023	DS DP U-999			
<b><u>TELITHROMYCIN - KETEK</u></b>						
N 021144 001	5635485	Apr 01, 2018	DS DP U-578			
<b><u>TELITHROMYCIN - KETEK</u></b>						
N 021144 002	5635485	Apr 01, 2018	DS DP U-578			
<b><u>TELMISARTAN - MICARDIS</u></b>						
N 020850 001	6358986	Jan 10, 2020				
<b><u>TELMISARTAN - MICARDIS</u></b>						
N 020850 002	6358986	Jan 10, 2020				
	7998953	Jun 06, 2020	U-1177			
	8003679	Oct 06, 2022	U-1176			
<b><u>TELMISARTAN - MICARDIS</u></b>						
N 020850 003	6358986	Jan 10, 2020				
<b><u>TELOTRISTAT ETIPRATE - XERMELO</u></b>						
N 208794 001	7553840	Dec 11, 2027	DS			
	7709493	Dec 11, 2027	DS U-1979			
	7968559	Dec 11, 2027	U-1979			
	8193204	Feb 27, 2031	DS			
	8653094	Dec 19, 2028	U-1979			
<b><u>TEMOZOLOMIDE - TEMODAR</u></b>						
N 022277 001	6987108	Sep 08, 2023	DP			
	7786118	Feb 21, 2023	DP			
	8623868	Feb 21, 2023	DP			
<b><u>TEMSIROLIMUS - TORISEL</u></b>						
N 022088 001	5362718	Feb 15, 2019	DS DP			
	5362718*PED	Aug 15, 2019				
	8026276	Jan 20, 2026	DP			
	8299116	Jul 25, 2023	DP			
	8455539	Jul 25, 2023	DP			
	8455539*PED	Jan 25, 2024				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TEMSIROLIMUS - TORISEL</u></b>						
N 022088 001	8722700	Jul 25, 2023	DP			
	8722700*PED	Jan 25, 2024				
	8791097	May 10, 2032		U-1550		
	8791097	May 10, 2032		U-1551		
	8791097*PED	Nov 10, 2032				
	RE44768	Feb 15, 2019	DS DP			
	RE44768*PED	Aug 15, 2019				
<b><u>TENOFOVIR ALAFENAMIDE FUMARATE - VEMLIDY</u></b>						
N 208464 001	7390791	May 07, 2022	DS DP		NCE	Nov 05, 2020
	7803788	Feb 02, 2022		U-999	NP	Nov 11, 2019
	8754065	Aug 15, 2032	DS DP	U-999		
	9296769	Aug 15, 2032	DS DP	U-999		
<b><u>TENOFOVIR DISOPROXIL FUMARATE - VIRREAD</u></b>						
N 021356 001	5922695	Jul 25, 2017	DS	U-1275	PED	Sep 24, 2017
	5922695	Jul 25, 2017	DS	U-248		
	5922695	Jul 25, 2017	DS	U-250		
	5922695	Jul 25, 2017	DS	U-256		
	5922695	Jul 25, 2017	DS	U-999		
	5935946	Jul 25, 2017	DS DP	U-1275		
	5935946	Jul 25, 2017	DS DP	U-248		
	5935946	Jul 25, 2017	DS DP	U-250		
	5935946	Jul 25, 2017	DS DP	U-256		
	5935946	Jul 25, 2017	DS DP	U-999		
	5977089	Jul 25, 2017	DS DP	U-1275		
	5977089	Jul 25, 2017	DS DP	U-248		
	5977089	Jul 25, 2017	DS DP	U-250		
	5977089	Jul 25, 2017	DS DP	U-256		
	5977089	Jul 25, 2017	DS DP	U-999		
	6043230	Jul 25, 2017		U-1275		
	6043230	Jul 25, 2017		U-248		
	6043230	Jul 25, 2017		U-250		
	6043230	Jul 25, 2017		U-256		
	6043230	Jul 25, 2017		U-999		
<b><u>TENOFOVIR DISOPROXIL FUMARATE - VIRREAD</u></b>						
N 021356 002	5922695	Jul 25, 2017	DS	U-1275	PED	Sep 24, 2017
	5922695	Jul 25, 2017	DS	U-248		
	5922695	Jul 25, 2017	DS	U-250		
	5922695	Jul 25, 2017	DS	U-256		
	5922695	Jul 25, 2017	DS	U-999		
	5935946	Jul 25, 2017	DS DP	U-1275		
	5935946	Jul 25, 2017	DS DP	U-248		
	5935946	Jul 25, 2017	DS DP	U-250		
	5935946	Jul 25, 2017	DS DP	U-256		
	5935946	Jul 25, 2017	DS DP	U-999		
	5977089	Jul 25, 2017	DS DP	U-1275		
	5977089	Jul 25, 2017	DS DP	U-248		
	5977089	Jul 25, 2017	DS DP	U-250		
	5977089	Jul 25, 2017	DS DP	U-256		
	5977089	Jul 25, 2017	DS DP	U-999		
	6043230	Jul 25, 2017		U-1275		
	6043230	Jul 25, 2017		U-248		
	6043230	Jul 25, 2017		U-250		
	6043230	Jul 25, 2017		U-256		
	6043230	Jul 25, 2017		U-999		
<b><u>TENOFOVIR DISOPROXIL FUMARATE - VIRREAD</u></b>						
N 021356 003	5922695	Jul 25, 2017	DS	U-1275	PED	Sep 24, 2017
	5922695	Jul 25, 2017	DS	U-248		
	5922695	Jul 25, 2017	DS	U-250		
	5922695	Jul 25, 2017	DS	U-256		
	5922695	Jul 25, 2017	DS	U-999		
	5935946	Jul 25, 2017	DS DP	U-1275		
	5935946	Jul 25, 2017	DS DP	U-248		
	5935946	Jul 25, 2017	DS DP	U-250		
	5935946	Jul 25, 2017	DS DP	U-256		
	5935946	Jul 25, 2017	DS DP	U-999		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</b>						
N 021356 003	5977089	Jul 25, 2017	DS DP U-1275			
	5977089	Jul 25, 2017	DS DP U-248			
	5977089	Jul 25, 2017	DS DP U-250			
	5977089	Jul 25, 2017	DS DP U-256			
	5977089	Jul 25, 2017	DS DP U-999			
	6043230	Jul 25, 2017		U-1275		
	6043230	Jul 25, 2017		U-248		
	6043230	Jul 25, 2017		U-250		
	6043230	Jul 25, 2017		U-256		
	6043230	Jul 25, 2017		U-999		
<b>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</b>						
N 021356 004	5922695	Jul 25, 2017	DS	U-1275		PED
	5922695	Jul 25, 2017	DS	U-248		Sep 24, 2017
	5922695	Jul 25, 2017	DS	U-250		
	5922695	Jul 25, 2017	DS	U-256		
	5922695	Jul 25, 2017	DS	U-999		
	5935946	Jul 25, 2017	DS DP	U-1275		
	5935946	Jul 25, 2017	DS DP	U-248		
	5935946	Jul 25, 2017	DS DP	U-250		
	5935946	Jul 25, 2017	DS DP	U-256		
	5935946	Jul 25, 2017	DS DP	U-999		
	5977089	Jul 25, 2017	DS DP	U-1275		
	5977089	Jul 25, 2017	DS DP	U-248		
	5977089	Jul 25, 2017	DS DP	U-250		
	5977089	Jul 25, 2017	DS DP	U-256		
	5977089	Jul 25, 2017	DS DP	U-999		
	6043230	Jul 25, 2017		U-1275		
	6043230	Jul 25, 2017		U-248		
	6043230	Jul 25, 2017		U-250		
	6043230	Jul 25, 2017		U-256		
	6043230	Jul 25, 2017		U-999		
<b>TENOFOVIR DISOPROXIL FUMARATE - VIREAD</b>						
N 022577 001	5922695	Jul 25, 2017	DS	U-1275		PED
	5922695	Jul 25, 2017	DS	U-248		Sep 24, 2017
	5922695	Jul 25, 2017	DS	U-250		
	5922695	Jul 25, 2017	DS	U-256		
	5922695	Jul 25, 2017	DS	U-999		
	5935946	Jul 25, 2017	DS DP	U-1275		
	5935946	Jul 25, 2017	DS DP	U-248		
	5935946	Jul 25, 2017	DS DP	U-250		
	5935946	Jul 25, 2017	DS DP	U-256		
	5935946	Jul 25, 2017	DS DP	U-999		
	5977089	Jul 25, 2017	DS DP	U-1275		
	5977089	Jul 25, 2017	DS DP	U-248		
	5977089	Jul 25, 2017	DS DP	U-250		
	5977089	Jul 25, 2017	DS DP	U-256		
	5977089	Jul 25, 2017	DS DP	U-999		
	6043230	Jul 25, 2017		U-1275		
	6043230	Jul 25, 2017		U-248		
	6043230	Jul 25, 2017		U-250		
	6043230	Jul 25, 2017		U-256		
	6043230	Jul 25, 2017		U-999		
<b>TERIFLUNOMIDE - AUBAGIO</b>						
N 202992 001	6794410	Sep 12, 2026		U-1285		
	8802735	Sep 14, 2030	DP			
	9186346	Feb 04, 2034		U-1786		
<b>TERIFLUNOMIDE - AUBAGIO</b>						
N 202992 002	6794410	Sep 12, 2026		U-1285		
	8802735	Sep 14, 2030	DP			
	9186346	Feb 04, 2034		U-1786		
<b>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</b>						
N 021318 001	6770623	Dec 08, 2018	DP	U-597		
	6977077	Aug 19, 2019		U-597		
	7144861	Dec 08, 2018	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u></b>						
N 021318 001	7163684	Aug 19, 2019	U-790			
	7351414	Aug 19, 2019	U-865			
	7517334	Mar 25, 2025	DP			
	7550434	Dec 08, 2018	DP U-982			
<b><u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u></b>						
N 021318 002	6770623	Dec 08, 2018	DP U-982			
	6977077	Aug 19, 2019	U-982			
	6977077	Aug 19, 2019	U-994			
	7144861	Dec 08, 2018	DP			
	7163684	Aug 19, 2019	U-983			
	7163684	Aug 19, 2019	U-994			
	7351414	Aug 19, 2019	U-984			
	7351414	Aug 19, 2019	U-994			
	7517334	Mar 25, 2025	DP			
	7550434	Dec 08, 2018	DP U-982			
<b><u>TESAMORELIN ACETATE - EGRIFTA</u></b>						
N 022505 001	5861379	May 26, 2020	DS DP U-1100			
	7144577	Jul 14, 2020	U-1100			
	7316997	Aug 14, 2023	U-1100			
<b><u>TESAMORELIN ACETATE - EGRIFTA</u></b>						
N 022505 002	7144577	Jul 14, 2020	U-1100			
	7316997	Aug 14, 2023	U-1100			
<b><u>TESTOSTERONE - TESTODERM TTS</u></b>						
N 020791 001	6348210	Nov 10, 2019	U-440			
<b><u>TESTOSTERONE - ANDROGEL</u></b>						
N 021015 001	6503894	Aug 30, 2020	U-490			
	9125816	Aug 30, 2020	U-490			
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020	U-490			
	9132089*PED	Mar 02, 2021				
<b><u>TESTOSTERONE - ANDROGEL</u></b>						
N 021015 002	6503894	Aug 30, 2020	U-490			
	9125816	Aug 30, 2020	U-490			
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020	U-490			
	9132089*PED	Mar 02, 2021				
<b><u>TESTOSTERONE - ANDROGEL</u></b>						
N 021015 003	6503894	Aug 30, 2020	U-490			
	9125816	Aug 30, 2020	U-490			
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020	U-490			
	9132089*PED	Mar 02, 2021				
<b><u>TESTOSTERONE - TESTIM</u></b>						
N 021454 001	7320968	Jan 18, 2025	U-843			
	7608605	Apr 21, 2023	U-1009			
	7608606	Apr 21, 2023	U-1009			
	7608607	Apr 21, 2023	U-1009			
	7608608	Apr 21, 2023	U-1009			
	7608609	Apr 21, 2023	U-1009			
	7608610	Apr 21, 2023	U-1009			
	7935690	Apr 21, 2023	U-1009			
	8063029	Apr 21, 2023	U-843			
	8178518	Apr 21, 2023	DP			
<b><u>TESTOSTERONE - FORTESTA</u></b>						
N 021463 001	6319913	Nov 09, 2018	U-490			
	6579865	Nov 09, 2018	DP			
<b><u>TESTOSTERONE - STRIANT</u></b>						
N 021543 001	6248358	Aug 23, 2019	U-527			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TESTOSTERONE - ANDROGEL</u></b>						
N 022309 001	6503894	Aug 30, 2020	U-1103			
	6503894*PED	Mar 02, 2021				
	8466136	Oct 12, 2026	DP			
	8466137	Oct 12, 2026		U-1103		
	8466138	Oct 12, 2026		U-1103		
	8486925	Oct 12, 2026	DP			
	8729057	Oct 12, 2026	DP			
	8741881	Oct 12, 2026		U-1103		
	8754070	Oct 12, 2026	DP			
	8759329	Oct 12, 2026	DP			
	9125816	Aug 30, 2020		U-1103		
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020	U-1103			
	9132089*PED	Mar 02, 2021				
<b><u>TESTOSTERONE - ANDROGEL</u></b>						
N 022309 002	6503894	Aug 30, 2020	U-1103			
	6503894*PED	Mar 02, 2021				
	8466136	Oct 12, 2026	DP			
	8466137	Oct 12, 2026		U-1103		
	8466138	Oct 12, 2026		U-1103		
	8486925	Oct 12, 2026	DP			
	8729057	Oct 12, 2026	DP			
	8741881	Oct 12, 2026		U-1103		
	8754070	Oct 12, 2026	DP			
	8759329	Oct 12, 2026	DP			
	9125816	Aug 30, 2020		U-1103		
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020	U-1103			
	9132089*PED	Mar 02, 2021				
<b><u>TESTOSTERONE - ANDROGEL</u></b>						
N 022309 003	6503894	Aug 30, 2020	U-1103			
	6503894*PED	Mar 02, 2021				
	8466136	Oct 12, 2026	DP			
	8466137	Oct 12, 2026		U-1103		
	8466138	Oct 12, 2026		U-1103		
	8486925	Oct 12, 2026	DP			
	8729057	Oct 12, 2026	DP			
	8741881	Oct 12, 2026		U-1103		
	8754070	Oct 12, 2026	DP			
	8759329	Oct 12, 2026	DP			
	9125816	Aug 30, 2020		U-1103		
	9125816*PED	Mar 02, 2021				
	9132089	Aug 30, 2020	U-1103			
	9132089*PED	Mar 02, 2021				
<b><u>TESTOSTERONE - AXIRON</u></b>						
N 022504 001	8419307	Feb 26, 2027	U-1386			
	8435944	Sep 27, 2027	U-1390			
	8784878	Jul 13, 2023	DP U-1545			
	8807861	Feb 26, 2027	DP U-1563			
	8993520	Jun 02, 2026	U-1390			
	9180194	Jun 02, 2026	U-1390			
	9289586	Feb 26, 2027	U-1390			
<b><u>TESTOSTERONE - VOGELXO</u></b>						
N 204399 002	8785426	Feb 11, 2034	DP U-1531			
	9295675	Feb 11, 2034	DP U-1531			
	9662340	Feb 11, 2034	DP U-1531			
<b><u>TESTOSTERONE - VOGELXO</u></b>						
N 204399 003	8785426	Feb 11, 2034	DP U-1531			
	9295675	Feb 11, 2034	DP U-1531			
	9662340	Feb 11, 2034	DP U-1531			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TESTOSTERONE - NATESTO</u></b>						
N 205488 001	8574622	Feb 04, 2024	DP			
	8784869	Feb 04, 2024	DP			
	8784882	Feb 04, 2024	DP U-1557			
	8877230	Feb 04, 2024	U-1616			
<b><u>TESTOSTERONE UNDECANOATE - AVEED</u></b>						
N 022219 001	7718640	Mar 14, 2027	DP			
	8338395	Feb 27, 2026	U-1500			
<b><u>THALIDOMIDE - THALOMID</u></b>						
N 020785 001	6045501	Aug 28, 2018	U-371			
	6045501	Aug 28, 2018	U-731			
	6315720	Oct 23, 2020	U-442			
	6315720	Oct 23, 2020	U-731			
	6561976	Aug 28, 2018	U-371			
	6561976	Aug 28, 2018	U-731			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	6908432	Aug 28, 2018	U-371			
	6908432	Aug 28, 2018	U-731			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023	DP			
	7435745	Nov 03, 2017	U-899			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-442			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7959566	Oct 23, 2020	U-1155			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8589188	Aug 28, 2018	U-1465			
	8626531	Oct 23, 2020	U-1465			
<b><u>THALIDOMIDE - THALOMID</u></b>						
N 020785 002	6045501	Aug 28, 2018	U-371			
	6045501	Aug 28, 2018	U-731			
	6315720	Oct 23, 2020	U-442			
	6315720	Oct 23, 2020	U-731			
	6561976	Aug 28, 2018	U-371			
	6561976	Aug 28, 2018	U-731			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	6908432	Aug 28, 2018	U-371			
	6908432	Aug 28, 2018	U-731			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023	DP			
	7435745	Nov 03, 2017	U-899			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-442			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>THALIDOMIDE - THALOMID</u></b>						
N 020785 002	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7959566	Oct 23, 2020	U-1155			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8589188	Aug 28, 2018	U-1465			
	8626531	Oct 23, 2020	U-1465			
<b><u>THALIDOMIDE - THALOMID</u></b>						
N 020785 003	6045501	Aug 28, 2018	U-371			
	6045501	Aug 28, 2018	U-731			
	6315720	Oct 23, 2020	U-442			
	6315720	Oct 23, 2020	U-731			
	6561976	Aug 28, 2018	U-371			
	6561976	Aug 28, 2018	U-731			
	6561977	Oct 23, 2020	U-371			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-371			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-371			
	6869399	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-732			
	6869399	Oct 23, 2020	U-733			
	6908432	Aug 28, 2018	U-371			
	6908432	Aug 28, 2018	U-731			
	7141018	Oct 23, 2020	U-371			
	7141018	Oct 23, 2020	U-731			
	7141018	Oct 23, 2020	U-732			
	7141018	Oct 23, 2020	U-733			
	7230012	Dec 09, 2023	DP			
	7435745	Nov 03, 2017	U-899			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-442			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7959566	Oct 23, 2020	U-1155			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8589188	Aug 28, 2018	U-1465			
	8626531	Oct 23, 2020	U-1465			
<b><u>THIOTEPA - TEPADINA</u></b>						
N 020785 004	6045501	Aug 28, 2018	U-731			
	6315720	Oct 23, 2020	U-731			
	6561976	Aug 28, 2018	U-731			
	6561977	Oct 23, 2020	U-731			
	6755784	Oct 23, 2020	U-731			
	6869399	Oct 23, 2020	U-731			
	6908432	Aug 28, 2018	U-731			
	7141018	Oct 23, 2020	U-731			
	7435745	Nov 03, 2017	U-899			
	7874984	Aug 28, 2018	U-1109			
	7874984	Aug 28, 2018	U-371			
	7874984	Aug 28, 2018	U-442			
	7874984	Aug 28, 2018	U-732			
	7874984	Aug 28, 2018	U-733			
	7959566	Oct 23, 2020	U-1155			
	8204763	Aug 28, 2018	U-1249			
	8315886	Oct 23, 2020	U-1249			
	8589188	Aug 28, 2018	U-1465			
	8626531	Oct 23, 2020	U-1465			

**THIOTEPA - TEPADINA**

N 208264 001

I-747

Jan 26, 2020

ODE-129

Jan 26, 2024

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<b><u>THIOTEPA - TEPADINA</u></b>							
N 208264	002				I-747 ODE-129	Jan 26, 2020 Jan 26, 2024	
<b><u>TICAGRELOR - BRILINTA</u></b>							
N 022433	001	6251910 6525060 6525060 6525060 6525060 7250419 7250419 7250419 7250419 7250419 7250419 7250419 7250419 7250419 7265124 7265124 7265124 7265124 8425934 RE46276 RE46276 RE46276 RE46276	Jul 15, 2018 Dec 02, 2019 Dec 02, 2019 Jul 09, 2021 Jul 09, 2021 Jul 09, 2021 Jul 09, 2021 Apr 17, 2030 Oct 30, 2024 Oct 30, 2024 Oct 30, 2024 Oct 30, 2024	DS DS DP U-1171 DS DP U-1860 DS DP U-1862 DS DP U-1863 DS DP U-1171 DS DP U-1860 DS DP U-1864 DS DP U-1865 DS DP U-1866 DS DP U-1867 DS DP U-1171 DS DP U-1860 DS DP U-1868 DS DP U-1869 DP DS DP U-1935 DS DP U-1936 DS DP U-1937 DS DP U-1938		I-714	Sep 03, 2018
N 022433	002	6251910 6525060 6525060 6525060 6525060 7250419 7250419 7250419 7250419 7250419 7250419 7250419 7250419 7250419 7265124 7265124 7265124 7265124 8425934 RE46276 RE46276 RE46276 RE46276	Jul 15, 2018 Dec 02, 2019 Dec 02, 2019 Jul 09, 2021 Jul 09, 2021 Jul 09, 2021 Jul 09, 2021 Apr 17, 2030 Oct 30, 2024 Oct 30, 2024 Oct 30, 2024 Oct 30, 2024	DS DS DP U-1171 DS DP U-1860 DS DP U-1862 DS DP U-1863 DS DP U-1171 DS DP U-1860 DS DP U-1864 DS DP U-1865 DS DP U-1866 DS DP U-1867 DS DP U-1171 DS DP U-1860 DS DP U-1868 DS DP U-1869 DP DS DP U-1935 DS DP U-1936 DS DP U-1937 DS DP U-1938		NS	Sep 03, 2018
<b><u>TIGECYCLINE - TYGACIL</u></b>							
N 021821	001	7879828 8372995 8975242 9254328 9694078	Feb 05, 2029 Oct 08, 2030 Oct 24, 2028 Mar 13, 2026 Mar 13, 2026	DP DP DP DP DP			
<b><u>TIMOLOL MALEATE - TIMOLOL MALEATE</u></b>							
N 020963	001	6174524	Mar 26, 2019	DP			
<b><u>TIMOLOL MALEATE - TIMOLOL MALEATE</u></b>							
N 020963	002	6174524	Mar 26, 2019	DP			
<b><u>TIMOLOL MALEATE - ISTALOL</u></b>							
N 021516	001	6335335 6645963	Nov 02, 2018 Nov 16, 2018	DP DP			
<b><u>TIOTROPIUM BROMIDE - SPIRIVA</u></b>							
N 021395	001	6777423 6777423*PED 6908928 6908928	Sep 24, 2021 Mar 24, 2022 Sep 24, 2021 Sep 24, 2021	DS DP DS DP U-566 DS DP U-762			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TIOTROPIUM BROMIDE - SPIRIVA</u></b>						
N 021395 001	6908928*PED	Mar 24, 2022				
	7070800	Jan 22, 2022	DP	U-566		
	7070800*PED	Jul 22, 2022				
	7309707	Sep 24, 2021	DS	DP		
	7309707*PED	Mar 24, 2022				
	7642268	Sep 24, 2021	DS	DP		
	7642268*PED	Mar 24, 2022				
	7694676	Mar 12, 2027	DP			
	7694676*PED	Sep 12, 2027				
	8022082	Jan 19, 2026	DP	U-1186		
	8022082*PED	Jul 19, 2026				
	9010323	Apr 19, 2030	DP			
	RE38912	Oct 11, 2021	DP			
	RE38912*PED	Apr 11, 2022				
	RE39820	Jan 30, 2018	DS	DP	U-566	
	RE39820*PED	Jul 30, 2018				
<b><u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u></b>						
N 021936 001	6846413	Aug 28, 2018	DP		NP	Sep 24, 2017
	6846413*PED	Feb 28, 2019			NPP	Feb 15, 2020
	6977042	Aug 28, 2018	DP		PED	Mar 24, 2018
	6977042*PED	Feb 28, 2019			PED	Aug 15, 2020
	6988496	Feb 23, 2020	DP			
	6988496*PED	Aug 23, 2020				
	7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7802568	Feb 26, 2019	DP			
	7802568*PED	Aug 26, 2019				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	7988001	Aug 04, 2021	DP			
	7988001*PED	Feb 04, 2022				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Oct 01, 2027				
	RE39820	Jan 30, 2018	DS	DP	U-1593	
	RE39820*PED	Jul 30, 2018				
<b><u>TIOTROPIUM BROMIDE - SPIRIVA RESPIMAT</u></b>						
N 021936 002	6846413	Aug 28, 2018	DP		NP	Sep 15, 2018
	6846413*PED	Feb 28, 2019			NPP	Feb 15, 2020
	6977042	Aug 28, 2018	DP		PED	Mar 15, 2019
	6977042*PED	Feb 28, 2019			PED	Aug 15, 2020
	6988496	Feb 23, 2020	DP			
	6988496*PED	Aug 23, 2020				
	7284474	Aug 26, 2024	DP			
	7284474*PED	Feb 26, 2025				
	7396341	Oct 10, 2026	DP			
	7396341*PED	Apr 10, 2027				
	7802568	Feb 26, 2019	DP			
	7802568*PED	Aug 26, 2019				
	7837235	Mar 13, 2028	DP			
	7837235*PED	Sep 13, 2028				
	7896264	May 26, 2025	DP			
	7896264*PED	Nov 26, 2025				
	7988001	Aug 04, 2021	DP			
	7988001*PED	Feb 04, 2022				
	8733341	Oct 16, 2030	DP			
	8733341*PED	Apr 16, 2031				
	9027967	Mar 31, 2027	DP			
	9027967*PED	Oct 01, 2027				
	RE39820	Jan 30, 2018	DS	DP		
	RE39820*PED	Jul 30, 2018				

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURF</u></b>						
N 207981 001	6479500	Mar 16, 2020		U-1751		
	9527833	Jun 17, 2034	DS DP			
	RE46284	Dec 16, 2026		U-1751		
<b><u>TIPIRACIL HYDROCHLORIDE; TRIFLURIDINE - LONSURF</u></b>						
N 207981 002	6479500	Mar 16, 2020		U-1751		
	9527833	Jun 17, 2034	DS DP			
	RE46284	Dec 16, 2026		U-1751		
<b><u>TIPRANAVIR - APTIVUS</u></b>						
N 021814 001	5852195	Jun 22, 2019	DS			
	6147095	Oct 29, 2019		U-670		
	6231887	Jul 27, 2018	DP			
<b><u>TIPRANAVIR - APTIVUS</u></b>						
N 022292 001	5852195	Jun 22, 2019	DS			
	6147095	Oct 29, 2019		U-670		
<b><u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u></b>						
N 020912 001	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
	6770660	May 01, 2023		U-1444		
<b><u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u></b>						
N 020912 002	5978698	Oct 08, 2017		U-1897		
	6136794	Jan 29, 2019		U-1898		
	6770660	May 01, 2023		U-1444		
<b><u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u></b>						
N 020913 001	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
	6770660	May 01, 2023		U-1444		
<b><u>TIROFIBAN HYDROCHLORIDE - AGGRASTAT</u></b>						
N 020913 002	5978698	Oct 08, 2017				
	6136794	Jan 29, 2019				
	6770660	May 01, 2023		U-1444		
<b><u>TOBRAMYCIN - TOBI PODHALER</u></b>						
N 201688 001	7368102	Dec 19, 2022	DP	U-909		
	7442388	May 10, 2020	DP			
	7516741	Jan 11, 2024	DP			
	7559325	Oct 27, 2025	DP			
	8069851	Sep 24, 2024	DP			
	8349294	May 10, 2020	DP			
	8715623	Dec 19, 2022	DP	U-909		
<b><u>TOBRAMYCIN - BETHKIS</u></b>						
N 201820 001	6987094	Sep 22, 2022	DP			
	7696178	Mar 17, 2023	DP			
	7939502	Jun 14, 2022		U-1324		
<b><u>TOFACITINIB CITRATE - XELJANZ</u></b>						
N 203214 001	6956041	Dec 08, 2020	DP		I-761	Dec 14, 2020
	6965027	Mar 25, 2023	DS		NCE	Nov 06, 2017
	7091208	Dec 08, 2020		U-247		
	7265221	Dec 08, 2020	DS			
	7301023	May 23, 2022	DS			
	RE41783	Dec 08, 2025	DS			
<b><u>TOFACITINIB CITRATE - XELJANZ XR</u></b>						
N 208246 001	6956041	Dec 08, 2020	DP		I-761	Dec 14, 2020
	6965027	Mar 25, 2023	DS		NCE	Nov 06, 2017
	7091208	Dec 08, 2020		U-247		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TOFACITINIB CITRATE - XELJANZ XR</u></b>						
N 208246 001	7265221	Dec 08, 2020	DS			
	7301023	May 23, 2022	DS			
	RE41783	Dec 08, 2025	DS			
<b><u>TOLTERODINE TARTRATE - DETROL LA</u></b>						
N 021228 001	6630162	Nov 11, 2019	DP U-544			
	6770295	Aug 26, 2019	DP U-544			
	6911217	Nov 11, 2019	DP U-544			
	6911217*PED	May 11, 2020				
<b><u>TOLTERODINE TARTRATE - DETROL LA</u></b>						
N 021228 002	6630162	Nov 11, 2019	DP U-544			
	6770295	Aug 26, 2019	DP U-544			
	6911217	Nov 11, 2019	DP U-544			
	6911217*PED	May 11, 2020				
<b><u>TOLVAPTAN - SAMSCA</u></b>						
N 022275 001	5753677	May 19, 2020	U-978			
	8501730	Sep 01, 2026	DS			
<b><u>TOLVAPTAN - SAMSCA</u></b>						
N 022275 002	5753677	May 19, 2020	U-978			
	8501730	Sep 01, 2026	DS			
<b><u>TOLVAPTAN - SAMSCA</u></b>						
N 022275 003	5753677	May 19, 2020	U-978			
	8501730	Sep 01, 2026	DS			
<b><u>TOPIRAMATE - TOPAMAX</u></b>						
N 020844 001	7125560	Mar 01, 2019	U-766			
<b><u>TOPIRAMATE - TOPAMAX</u></b>						
N 020844 002	7125560	Mar 01, 2019	U-766			
<b><u>TOPIRAMATE - TOPAMAX SPRINKLE</u></b>						
N 020844 003	7125560	Mar 01, 2019	U-766			
<b><u>TOPIRAMATE - TROKENDI XR</u></b>						
N 201635 001	8298576	Apr 04, 2028	DP U-106			
	8298576	Apr 04, 2028	DP U-1992			
	8298580	Nov 16, 2027	DP U-106			
	8298580	Nov 16, 2027	DP U-1992			
	8663683	Nov 16, 2027	DP U-106			
	8663683	Nov 16, 2027	DP U-1992			
	8877248	Nov 16, 2027	DP U-106			
	8877248	Nov 16, 2027	DP U-1992			
	8889191	Nov 16, 2027	U-106			
	8889191	Nov 16, 2027	U-1992			
	8992989	Nov 16, 2027	DP U-1675			
	8992989	Nov 16, 2027	DP U-1992			
	9549940	Nov 16, 2027	DP U-1675			
	9549940	Nov 16, 2027	DP U-1992			
	9555004	Nov 16, 2027	DP U-1675			
	9555004	Nov 16, 2027	DP U-1992			
	9622983	Nov 16, 2027	DP U-1675			
	9622983	Nov 16, 2027	DP U-1992			
<b><u>TOPIRAMATE - TROKENDI XR</u></b>						
N 201635 002	8298576	Apr 04, 2028	DP U-106			
	8298576	Apr 04, 2028	DP U-1992			
	8298580	Nov 16, 2027	DP U-106			
	8298580	Nov 16, 2027	DP U-1992			
	8663683	Nov 16, 2027	DP U-106			
	8663683	Nov 16, 2027	DP U-1992			
	8877248	Nov 16, 2027	DP U-106			
	8877248	Nov 16, 2027	DP U-1992			
	8889191	Nov 16, 2027	U-106			
	8889191	Nov 16, 2027	U-1992			
	8992989	Nov 16, 2027	DP U-1675			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TOPIRAMATE - TROKENDI XR</u></b>						
N 201635 002	8992989	Nov 16, 2027	DP U-1992			
	9549940	Nov 16, 2027	DP U-1675			
	9549940	Nov 16, 2027	DP U-1992			
	9555004	Nov 16, 2027	DP U-1675			
	9555004	Nov 16, 2027	DP U-1992			
	9622983	Nov 16, 2027	DP U-1675			
	9622983	Nov 16, 2027	DP U-1992			
<b><u>TOPIRAMATE - TROKENDI XR</u></b>						
N 201635 003	8298576	Apr 04, 2028	DP U-106			
	8298576	Apr 04, 2028	DP U-1992			
	8298580	Nov 16, 2027	DP U-106			
	8298580	Nov 16, 2027	DP U-1992			
	8663683	Nov 16, 2027	DP U-106			
	8663683	Nov 16, 2027	DP U-1992			
	8877248	Nov 16, 2027	DP U-106			
	8877248	Nov 16, 2027	DP U-1992			
	8889191	Nov 16, 2027	U-106			
	8889191	Nov 16, 2027	U-1992			
	8992989	Nov 16, 2027	DP U-1675			
	8992989	Nov 16, 2027	DP U-1992			
	9549940	Nov 16, 2027	DP U-1675			
	9549940	Nov 16, 2027	DP U-1992			
	9555004	Nov 16, 2027	DP U-1675			
	9555004	Nov 16, 2027	DP U-1992			
	9622983	Nov 16, 2027	DP U-1675			
	9622983	Nov 16, 2027	DP U-1992			
<b><u>TOPIRAMATE - TROKENDI XR</u></b>						
N 201635 004	8298576	Apr 04, 2028	DP U-106			
	8298576	Apr 04, 2028	DP U-1992			
	8298580	Nov 16, 2027	DP U-106			
	8298580	Nov 16, 2027	DP U-1992			
	8663683	Nov 16, 2027	DP U-106			
	8663683	Nov 16, 2027	DP U-1992			
	8877248	Nov 16, 2027	DP U-106			
	8877248	Nov 16, 2027	DP U-1992			
	8889191	Nov 16, 2027	U-106			
	8889191	Nov 16, 2027	U-1992			
	8992989	Nov 16, 2027	DP U-1675			
	8992989	Nov 16, 2027	DP U-1992			
	9549940	Nov 16, 2027	DP U-1675			
	9549940	Nov 16, 2027	DP U-1992			
	9555004	Nov 16, 2027	DP U-1675			
	9555004	Nov 16, 2027	DP U-1992			
	9622983	Nov 16, 2027	DP U-1675			
	9622983	Nov 16, 2027	DP U-1992			
<b><u>TOPIRAMATE - QUDEXY XR</u></b>						
N 205122 001	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<b><u>TOPIRAMATE - QUDEXY XR</u></b>						
N 205122 002	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<b><u>TOPIRAMATE - QUDEXY XR</u></b>						
N 205122 003	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TOPIRAMATE - QUDEXY XR</u></b>						
N 205122 004	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<b><u>TOPIRAMATE - QUDEXY XR</u></b>						
N 205122 005	8652527	Mar 19, 2033	DP			
	8889190	Mar 19, 2033	DP			
	9101545	Mar 19, 2033	DP			
	9555005	Mar 19, 2033	DP			
<b><u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u></b>						
N 020981 001	8158645	Dec 10, 2024	DP			
<b><u>TOPOTECAN HYDROCHLORIDE - HYCAMTIN</u></b>						
N 020981 002	8158645	Dec 10, 2024	DP			
<b><u>TRABECTEDIN - YONDELIS</u></b>						
N 207953 001	8895557	Jan 07, 2028	DP		NCE ODE-100	Oct 23, 2020 Oct 23, 2022
<b><u>TRAMADOL HYDROCHLORIDE - ULTRAM</u></b>						
N 020281 001	6339105	Oct 12, 2019	U-435			
<b><u>TRAMADOL HYDROCHLORIDE - ULTRAM</u></b>						
N 020281 002	6339105	Oct 12, 2019	U-435			
<b><u>TRAMADOL HYDROCHLORIDE - RYBIX ODT</u></b>						
N 021693 001	6106861	Dec 05, 2017	DP			
<b><u>TRAMADOL HYDROCHLORIDE - RYZOLT</u></b>						
N 021745 001	6607748	Jun 29, 2020	DP			
	7988998	Oct 27, 2023	DP			
<b><u>TRAMADOL HYDROCHLORIDE - RYZOLT</u></b>						
N 021745 002	6607748	Jun 29, 2020	DP			
	7988998	Oct 27, 2023	DP			
<b><u>TRAMADOL HYDROCHLORIDE - RYZOLT</u></b>						
N 021745 003	6607748	Jun 29, 2020	DP			
	7988998	Oct 27, 2023	DP			
<b><u>TRAMADOL HYDROCHLORIDE - CONZIP</u></b>						
N 022370 001	7858118	Apr 11, 2022	DP U-1104			
<b><u>TRAMADOL HYDROCHLORIDE - CONZIP</u></b>						
N 022370 002	7858118	Apr 11, 2022	DP U-1104			
<b><u>TRAMADOL HYDROCHLORIDE - CONZIP</u></b>						
N 022370 003	7858118	Apr 11, 2022	DP U-1104			
<b><u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u></b>						
N 204114 001	7378423	Sep 13, 2025	DS DP		I-745	Jun 22, 2020
	8580304	Jan 28, 2032	DP		M-170	Nov 20, 2018
	8703781	Oct 15, 2030	DS DP U-1712		NCE	May 29, 2018
	8703781	Oct 15, 2030	DS DP U-2033		ODE-148	Jun 22, 2024
	8835443	Sep 13, 2025	U-1581		ODE-48	May 29, 2020
	8835443	Sep 13, 2025	U-1582		ODE-57	Jan 08, 2021
	8835443	Sep 13, 2025	U-2020			
	8835443	Sep 13, 2025	U-2037			
	8952018	Oct 15, 2030	U-2020			
	9155706	Jan 28, 2032	DP			
	9271941	Jan 28, 2032	DP			
<b><u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u></b>						
N 204114 002	7378423	Sep 13, 2025	DS DP		I-745	Jun 22, 2020
	8580304	Jan 28, 2032	DP		NCE	May 29, 2018
	8703781	Oct 15, 2030	DS DP U-1712		ODE-148	Jun 22, 2024
	8703781	Oct 15, 2030	DS DP U-2033		ODE-48	May 29, 2020
	8835443	Sep 13, 2025	U-1581		ODE-57	Jan 08, 2021

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u></b>						
N 204114 002	8835443	Sep 13, 2025		U-1582		
	8835443	Sep 13, 2025		U-2020		
	8835443	Sep 13, 2025		U-2037		
	8952018	Oct 15, 2030		U-2020		
	9155706	Jan 28, 2032	DP			
	9271941	Jan 28, 2032	DP			
<b><u>TRAMETINIB DIMETHYL SULFOXIDE - MEKINIST</u></b>						
N 204114 003	7378423	Sep 13, 2025	DS DP		I-745	Jun 22, 2020
	8580304	Jan 28, 2032	DP		M-170	Nov 20, 2018
	8703781	Oct 15, 2030	DS DP	U-1712	NCE	May 29, 2018
	8703781	Oct 15, 2030	DS DP	U-2033	ODE-148	Jun 22, 2024
	8835443	Sep 13, 2025		U-1581	ODE-48	May 29, 2020
	8835443	Sep 13, 2025		U-1582	ODE-57	Jan 08, 2021
	8835443	Sep 13, 2025		U-2020		
	8835443	Sep 13, 2025		U-2037		
	8952018	Oct 15, 2030		U-2020		
	9155706	Jan 28, 2032	DP			
	9271941	Jan 28, 2032	DP			
<b><u>TRANEXAMIC ACID - LYSTEDA</u></b>						
N 022430 001	7947739	Mar 04, 2025	DP			
	8022106	Mar 04, 2025		U-1182		
	8273795	Mar 04, 2025		U-1182		
	8487005	Mar 04, 2025	DP	U-1182		
	8791160	Mar 04, 2025	DP	U-1182		
	8809394	Mar 04, 2025	DP	U-1182		
	8957113	Mar 04, 2025	DP	U-1182		
	9060939	Mar 04, 2025	DP			
<b><u>TRAVOPROST - TRAVATAN Z</u></b>						
N 021994 001	8268299	Oct 13, 2029	DP			
	8323630	Sep 20, 2027	DP			
	8388941	Sep 20, 2027	DP			
<b><u>TRAVOPROST - IZBA</u></b>						
N 204822 001	8178582	Oct 10, 2029	DP			
	8722735	Oct 10, 2029	DP			
	8754123	May 19, 2029	DP			
	9144561	Mar 13, 2029	DP			
<b><u>TRAZODONE HYDROCHLORIDE - DESYREL</u></b>						
N 018207 001	8133893	Mar 13, 2029	DS DP			
<b><u>TRAZODONE HYDROCHLORIDE - DESYREL</u></b>						
N 018207 002	8133893	Mar 13, 2029	DS DP			
<b><u>TRAZODONE HYDROCHLORIDE - DESYREL</u></b>						
N 018207 003	8133893	Mar 13, 2029	DS DP			
<b><u>TRAZODONE HYDROCHLORIDE - DESYREL</u></b>						
N 018207 004	8133893	Mar 13, 2029	DS DP			
<b><u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u></b>						
N 022411 001	6607748	Jun 29, 2020	DP			
	7829120	Mar 27, 2027		DP U-796		
	8133893	Mar 13, 2029	DS	DP		
<b><u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u></b>						
N 022411 002	6607748	Jun 29, 2020	DP			
	7829120	Mar 27, 2027		DP U-796		
	8133893	Mar 13, 2029	DS	DP		
<b><u>TREPROSTINIL - REMODULIN</u></b>						
N 021272 001	6765117	Oct 24, 2017	DS			
	7999007	Mar 29, 2029		DP U-1437		
	8497393	Dec 15, 2028	DS			
	8653137	Sep 05, 2028		U-1437		
	8658694	Sep 05, 2028		U-1437		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TREPROSTINIL - REMODULIN</u></b>						
N 021272 001	9199908	May 24, 2024		U-1771		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024		U-2036		
<b><u>TREPROSTINIL - REMODULIN</u></b>						
N 021272 002	6765117	Oct 24, 2017	DS			
	7999007	Mar 29, 2029	DP	U-1437		
	8497393	Dec 15, 2028	DS			
	8653137	Sep 05, 2028		U-1437		
	8658694	Sep 05, 2028		U-1437		
	9199908	May 24, 2024		U-1771		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024		U-2036		
<b><u>TREPROSTINIL - REMODULIN</u></b>						
N 021272 003	6765117	Oct 24, 2017	DS			
	7999007	Mar 29, 2029	DP	U-1437		
	8497393	Dec 15, 2028	DS			
	8653137	Sep 05, 2028		U-1437		
	8658694	Sep 05, 2028		U-1437		
	9199908	May 24, 2024		U-1771		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024		U-2036		
<b><u>TREPROSTINIL - REMODULIN</u></b>						
N 021272 004	6765117	Oct 24, 2017	DS			
	7999007	Mar 29, 2029	DP	U-1437		
	8497393	Dec 15, 2028	DS			
	8653137	Sep 05, 2028		U-1437		
	8658694	Sep 05, 2028		U-1437		
	9199908	May 24, 2024		U-1771		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
	9713599	Dec 16, 2024		U-2036		
<b><u>TREPROSTINIL - TYVASO</u></b>						
N 022387 001	6521212	Nov 13, 2018		U-1018		
	6756033	Nov 13, 2018		U-1018		
	6765117	Oct 24, 2017	DS			
	8497393	Dec 15, 2028	DS			
	9339507	Mar 10, 2028	DP			
	9358240	May 05, 2028		U-1849		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
		Dec 16, 2024		U-2036		
<b><u>TREPROSTINIL DIOLAMINE - ORENITRAM</u></b>						
N 203496 001	6765117	Oct 24, 2017	DS		D-156	Jan 28, 2019
	7417070	Jul 30, 2026	DS		D-157	Jan 28, 2019
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024	DP			
	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS			
	8747897	Oct 08, 2029	DP			
	9050311	May 24, 2024	DS	DP		
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026	DP	U-1877		
	9422223	May 24, 2024	DP			
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<b><u>TREPROSTINIL DIOLAMINE - ORENITRAM</u></b>						
N 203496 002	6765117	Oct 24, 2017	DS		D-156	Jan 28, 2019
	7417070	Jul 30, 2026	DS		D-157	Jan 28, 2019
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024	DP			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>TREPROSTINIL DIOLAMINE - ORENITRAM</b>						
N 203496 002	8349892	Jan 22, 2031	DP			
	8410169	Feb 13, 2030	DP			
	8497393	Dec 15, 2028	DS			
	8747897	Oct 08, 2029	DP			
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026		DP U-1877		
	9422223	May 24, 2024		DP		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<b>TREPROSTINIL DIOLAMINE - ORENITRAM</b>						
N 203496 003	6765117	Oct 24, 2017	DS		D-156	Jan 28, 2019
	7417070	Jul 30, 2026	DS		D-157	Jan 28, 2019
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024		DP		
	8349892	Jan 22, 2031		DP		
	8410169	Feb 13, 2030		DP		
	8497393	Dec 15, 2028	DS			
	8747897	Oct 08, 2029		DP		
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026		DP U-1877		
	9422223	May 24, 2024		DP		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<b>TREPROSTINIL DIOLAMINE - ORENITRAM</b>						
N 203496 004	6765117	Oct 24, 2017	DS		D-156	Jan 28, 2019
	7417070	Jul 30, 2026	DS		D-157	Jan 28, 2019
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024		DP		
	8349892	Jan 22, 2031		DP		
	8410169	Feb 13, 2030		DP		
	8497393	Dec 15, 2028	DS			
	8747897	Oct 08, 2029		DP		
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026		DP U-1877		
	9422223	May 24, 2024		DP		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<b>TREPROSTINIL DIOLAMINE - ORENITRAM</b>						
N 203496 005	6765117	Oct 24, 2017	DS		D-156	Jan 28, 2019
	7417070	Jul 30, 2026	DS		D-157	Jan 28, 2019
	7544713	Jul 14, 2024		U-1475		
	8252839	May 24, 2024		DP		
	8349892	Jan 22, 2031		DP		
	8410169	Feb 13, 2030		DP		
	8497393	Dec 15, 2028	DS			
	8747897	Oct 08, 2029		DP		
	9050311	May 24, 2024	DS DP			
	9278901	May 24, 2024		U-1475		
	9393203	Apr 27, 2026		DP U-1877		
	9422223	May 24, 2024		DP		
	9593066	Dec 15, 2028	DS			
	9604901	Dec 15, 2028	DS			
<b>TRETINOIN - RENOVA</b>						
N 021108 001	6531141	Mar 07, 2020				
<b>TRIAMCINOLONE ACETONIDE - TRIESENCE</b>						
N 022048 001	6395294	Jan 13, 2020		DP U-846		
	8128960	Dec 17, 2029		DP		
	8211880	Mar 10, 2029		U-1257		
	8211880	Mar 10, 2029		U-1258		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<b><u>TRIAMCINOLONE ACETONIDE - ZILRETTA</u></b>						
N 208845 001	8828440	Aug 04, 2031	DP		NP	Oct 06, 2020
	9555048	Aug 04, 2031		U-2151		
<b><u>TRIMETREXATE GLUCURONATE - NEUTREXIN</u></b>						
N 020326 001	6017922	May 18, 2018				
<b><u>TRIMETREXATE GLUCURONATE - NEUTREXIN</u></b>						
N 020326 002	6017922	May 18, 2018				
<b><u>TRIPTORELIN PAMOATE - TRIPTODUR KIT</u></b>						
N 208956 001					NP	Jun 29, 2020
					ODE-149	Jun 29, 2024
<b><u>TROGLITAZONE - PRELAY</u></b>						
N 020719 001	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
<b><u>TROGLITAZONE - PRELAY</u></b>						
N 020719 002	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
<b><u>TROGLITAZONE - PRELAY</u></b>						
N 020719 003	5859037	Nov 13, 2017	U-251			
	6011049	Nov 13, 2017	U-301			
<b><u>TROSPiUM CHLORIDE - SANCTURA XR</u></b>						
N 022103 001	7410978	Feb 01, 2025	DP			
	7759359	Nov 04, 2024	U-1071			
	7763635	Nov 04, 2024	U-1071			
	7781448	Nov 04, 2024	U-1071			
	7781449	Nov 04, 2024	U-1071			
<b><u>ULIPRISTAL ACETATE - ELLA</u></b>						
N 022474 001	8426392	Jun 12, 2030	U-1389			
	8512745	Jun 02, 2030	DP			
	8735380	Feb 20, 2029	DP			
	8962603	Jun 12, 2030	U-1657			
	9283233	Apr 13, 2030	U-1821			
<b><u>UMECLIDINIUM BROMIDE - INCRUSE ELLIPTA</u></b>						
N 205382 001	7488827	Dec 18, 2027	DS DP		M-172	Feb 24, 2019
	7498440	Apr 27, 2025	DS DP		NCE	Dec 18, 2018
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8183257	Jul 27, 2025	U-1476			
	8201556	Feb 05, 2029	DP			
	8309572	Apr 27, 2025	U-1476			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
<b><u>UMECLIDINIUM BROMIDE; VILANTEROL TRIFENATATE - ANORO ELLIPTA</u></b>						
N 203975 001	7439393	May 21, 2025	DS DP U-1476		NCE	May 10, 2018
	7488827	Dec 18, 2027	DS DP		NCE	Dec 18, 2018
	7498440	Apr 27, 2025	DS DP			
	7776895	Sep 11, 2022	DP			
	8113199	Oct 23, 2027	DP			
	8161968	Feb 05, 2028	DP			
	8183257	Jul 27, 2025	U-1476			
	8309572	Apr 27, 2025	U-1476			
	8511304	Jun 14, 2027	DP U-1476			
	8534281	Aug 10, 2029	DP			
	8746242	Oct 11, 2030	DP			
	9333310	Oct 02, 2027	DP			
	9750726	Nov 29, 2030	DP			
	RE44874	Mar 23, 2023	DS DP U-1476			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>UNOPROSTONE ISOPROPYL - RESCULA</u></b>						
N 021214 001	6458836	Jul 09, 2021		U-1315		
	6458836	Jul 09, 2021		U-333		
	6770675	Nov 24, 2018	DP	U-1322		
<b><u>URIDINE TRIACETATE - VISTOGARD</u></b>						
N 208159 001	6258795	Jul 10, 2018	DP		NCE	Sep 04, 2020
	7776838	Aug 17, 2027		U-1791	NP	Dec 11, 2018
					ODE-104	Dec 11, 2022
<b><u>URIDINE TRIACETATE - XURIDEN</u></b>						
N 208169 001	6258795	Jul 10, 2018	DP		NCE	Sep 04, 2020
					ODE-98	Sep 04, 2022
<b><u>VALBENAZINE TOSYLATE - INGREZZA</u></b>						
N 209241 001	8039627	Oct 06, 2029	DS DP		NCE	Apr 11, 2022
	8357697	Nov 08, 2027		U-1995		
<b><u>VALBENAZINE TOSYLATE - INGREZZA</u></b>						
N 209241 002	8039627	Oct 06, 2029	DS DP		NCE	Apr 11, 2022
	8357697	Nov 08, 2027		U-1995		
<b><u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u></b>						
N 021304 001					D-148	Apr 23, 2018
					NPP	Apr 23, 2018
<b><u>VALGANCICLOVIR HYDROCHLORIDE - VALCYTE</u></b>						
N 022257 001	8889109	Dec 11, 2027	DP		D-148	Apr 23, 2018
	9642911	Dec 11, 2027	DP		NPP	Apr 23, 2018
<b><u>VANDETANIB - CAPRELSA</u></b>						
N 022405 001	8067427	Aug 08, 2028	DP		ODE-9	Apr 06, 2018
	RE42353	Jun 27, 2022	DS DP			
<b><u>VANDETANIB - CAPRELSA</u></b>						
N 022405 002	8067427	Aug 08, 2028	DP		ODE-9	Apr 06, 2018
	RE42353	Jun 27, 2022	DS DP			
<b><u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u></b>						
N 021400 001	6362178	Oct 31, 2018	DS DP U-533			
	7696206	Oct 31, 2018	DS DP U-533			
	8273876	Jul 23, 2027		U-1288		
	8841446	Jul 03, 2023	DP			
<b><u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u></b>						
N 021400 002	6362178	Oct 31, 2018	DS DP U-533			
	7696206	Oct 31, 2018	DS DP U-533			
	8273876	Jul 23, 2027		U-1288		
	8841446	Jul 03, 2023	DP			
<b><u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u></b>						
N 021400 003	6362178	Oct 31, 2018	DS DP U-533			
	7696206	Oct 31, 2018	DS DP U-533			
	8273876	Jul 23, 2027		U-1288		
	8841446	Jul 03, 2023	DP			
<b><u>VARDENAFIL HYDROCHLORIDE - LEVITRA</u></b>						
N 021400 004	6362178	Oct 31, 2018	DS DP U-533			
	7696206	Oct 31, 2018	DS DP U-533			
	8273876	Jul 23, 2027		U-1288		
	8841446	Jul 03, 2023	DP			
<b><u>VARDENAFIL HYDROCHLORIDE - STAXYN</u></b>						
N 200179 001	6362178	Oct 31, 2018		U-155		
	7696206	Oct 31, 2018		U-155		
	8613950	Dec 23, 2028	DP			
<b><u>VARENICLINE TARTRATE - CHANTIX</u></b>						
N 021928 001	6410550	May 10, 2020	DS DP U-56		M-143	Oct 15, 2017
	6890927	May 06, 2022	DS DP U-56		M-144	Oct 15, 2017
	7265119	Aug 03, 2022	DS DP U-56		M-183	Aug 12, 2019

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>VARENICLINE TARTRATE - CHANTIX</b>						
N 021928 001					M-192	Dec 16, 2019
<b>VARENICLINE TARTRATE - CHANTIX</b>						
N 021928 002	6410550	May 10, 2020	DS DP U-56		M-143	Oct 15, 2017
	6890927	May 06, 2022	DS DP U-56		M-144	Oct 15, 2017
	7265119	Aug 03, 2022	DS DP U-56		M-183	Aug 12, 2019
					M-192	Dec 16, 2019
<b>VASOPRESSIN - VASOSTRICT</b>						
N 204485 001	9375478	Jan 30, 2035	U-1857			
	9687526	Jan 30, 2035	U-1857			
	9744209	Jan 30, 2035	U-1857			
	9744239	Jan 30, 2035	U-1857			
	9750785	Jan 30, 2035	DP			
<b>VASOPRESSIN - VASOSTRICT</b>						
N 204485 002	9375478	Jan 30, 2035	U-1857			
	9687526	Jan 30, 2035	U-1857			
	9744209	Jan 30, 2035	U-1857			
	9744239	Jan 30, 2035	U-1857			
	9750785	Jan 30, 2035	DP			
<b>VEMURAFENIB - ZELBORAF</b>						
N 202429 001	7504509	Oct 22, 2026	DS DP		I-757	Nov 06, 2020
	7863288	Jun 20, 2029	DS DP		M-184	Aug 31, 2019
	8143271	Jun 21, 2026	DS DP		ODE	Nov 06, 2024
	8470818	Aug 02, 2026	U-1418		ODE-13	Aug 17, 2018
	8470818	Aug 02, 2026	U-2164			
	8741920	Jul 27, 2030	DS DP			
	9447089	Jun 06, 2032	DP			
<b>VENETOCLAX - VENCLEXTA</b>						
N 208573 001	8546399	Jun 27, 2031	DS DP		NCE	Apr 11, 2021
	9174982	May 26, 2030	U-1835		ODE-114	Apr 11, 2023
<b>VENETOCLAX - VENCLEXTA</b>						
N 208573 002	8546399	Jun 27, 2031	DS DP		NCE	Apr 11, 2021
	9174982	May 26, 2030	U-1835		ODE-114	Apr 11, 2023
<b>VENETOCLAX - VENCLEXTA</b>						
N 208573 003	8546399	Jun 27, 2031	DS DP		NCE	Apr 11, 2021
	9174982	May 26, 2030	U-1835		ODE-114	Apr 11, 2023
<b>VILAZODONE HYDROCHLORIDE - VIIBRYD</b>						
N 022567 001	5532241	Sep 29, 2019	DS DP		D-146	Mar 16, 2018
	7834020	Jun 05, 2022	DS DP U-839			
	8193195	Jun 05, 2022	U-839			
	8236804	Jun 05, 2022	U-839			
	8673921	Jun 05, 2022	DS DP			
<b>VILAZODONE HYDROCHLORIDE - VIIBRYD</b>						
N 022567 002	5532241	Sep 29, 2019	DS DP		D-146	Mar 16, 2018
	7834020	Jun 05, 2022	DS DP U-839			
	8193195	Jun 05, 2022	U-839			
	8236804	Jun 05, 2022	U-839			
	8673921	Jun 05, 2022	DS DP			
<b>VILAZODONE HYDROCHLORIDE - VIIBRYD</b>						
N 022567 003	5532241	Sep 29, 2019	DS DP		D-146	Mar 16, 2018
	7834020	Jun 05, 2022	DS DP U-839			
	8193195	Jun 05, 2022	U-839			
	8236804	Jun 05, 2022	U-839			
	8673921	Jun 05, 2022	DS DP			
<b>VINCRISTINE SULFATE - MARQIBO KIT</b>						
N 202497 001	6723338	Mar 31, 2020	U-1271		ODE-28	Aug 09, 2019
	7247316	Sep 25, 2020	DP			
	7887836	Mar 31, 2020	U-1271			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>VISMODEGIB - ERIVEDGE</b>						
N 203388 001	7888364	Nov 11, 2028	DS DP			
	9278961	Dec 15, 2028		U-1825		
<b>VORAPAXAR SULFATE - ZONTIVITY</b>						
N 204886 001	7235567	Jun 13, 2021	DS DP		NCE	
	7304078	Apr 06, 2024	DS DP	U-1512		May 08, 2019
<b>VORICONAZOLE - VFEND</b>						
N 021267 001	6632803	Jun 02, 2018	DP			
<b>VORINOSTAT - ZOLINZA</b>						
N 021991 001	7399787	Feb 09, 2025		U-892		
	7456219	Mar 11, 2027	DS			
	7652069	Mar 04, 2023	DP			
	7732490	Mar 04, 2023		U-892		
	7851509	Feb 21, 2024	DP	U-892		
	8067472	Mar 04, 2023		U-892		
	8093295	May 16, 2026	DP			
	8101663	Mar 04, 2023		U-892		
	8450372	Mar 18, 2028		U-892		
<b>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</b>						
N 204447 001	7144884	Oct 02, 2022	DS DP	U-1439	NCE	
	8476279	Oct 02, 2022	DP	U-1439		Sep 30, 2018
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9227946	Jun 15, 2027		U-1668		
<b>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</b>						
N 204447 002	7144884	Oct 02, 2022	DS DP	U-1439	NCE	
	8476279	Oct 02, 2022	DP	U-1439		Sep 30, 2018
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9227946	Jun 15, 2027		U-1668		
<b>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</b>						
N 204447 003	7144884	Oct 02, 2022	DS DP	U-1439	NCE	
	8476279	Oct 02, 2022	DP	U-1439		Sep 30, 2018
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9227946	Jun 15, 2027		U-1668		
<b>VORTIOXETINE HYDROBROMIDE - TRINTELLIX</b>						
N 204447 004	7144884	Oct 02, 2022	DS DP	U-1439	NCE	
	8476279	Oct 02, 2022	DP	U-1439		Sep 30, 2018
	8722684	Jun 30, 2031	DS DP			
	8969355	Jun 15, 2027		U-1668		
	9227946	Jun 15, 2027		U-1668		
<b>ZICONOTIDE ACETATE - PRIALT</b>						
N 021060 001	8653033	Oct 01, 2024		U-48		
	8653033	Oct 01, 2024		U-55		
	8765680	Oct 01, 2024		U-48		
	8765680	Oct 01, 2024		U-55		
	9707270	Oct 01, 2024		U-2084		
<b>ZICONOTIDE ACETATE - PRIALT</b>						
N 021060 002	8653033	Oct 01, 2024		U-48		
	8653033	Oct 01, 2024		U-55		
	8765680	Oct 01, 2024		U-48		
	8765680	Oct 01, 2024		U-55		
	9707270	Oct 01, 2024		U-2084		
<b>ZICONOTIDE ACETATE - PRIALT</b>						
N 021060 003	8653033	Oct 01, 2024		U-48		
	8653033	Oct 01, 2024		U-55		
	8765680	Oct 01, 2024		U-48		
	8765680	Oct 01, 2024		U-55		
	9707270	Oct 01, 2024		U-2084		

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b>ZICONOTIDE ACETATE - PRIALT</b>						
N 021060 003	8653033	Oct 01, 2024	U-48			
	8653033	Oct 01, 2024	U-55			
	8765680	Oct 01, 2024	U-48			
	8765680	Oct 01, 2024	U-55			
	9707270	Oct 01, 2024	U-2084			
<b>ZICONOTIDE ACETATE - PRIALT</b>						
N 021060 004	8653033	Oct 01, 2024	U-48			
	8653033	Oct 01, 2024	U-55			
	8765680	Oct 01, 2024	U-48			
	8765680	Oct 01, 2024	U-55			
	9707270	Oct 01, 2024	U-2084			
<b>ZIPRASIDONE HYDROCHLORIDE - GEODON</b>						
N 020825 001	6150366	May 27, 2019	DP			
	6245766	Dec 18, 2018	U-601			
<b>ZIPRASIDONE HYDROCHLORIDE - GEODON</b>						
N 020825 002	6150366	May 27, 2019	DP			
	6245766	Dec 18, 2018	U-601			
<b>ZIPRASIDONE HYDROCHLORIDE - GEODON</b>						
N 020825 003	6150366	May 27, 2019	DP			
	6245766	Dec 18, 2018	U-601			
<b>ZIPRASIDONE HYDROCHLORIDE - GEODON</b>						
N 020825 004	6150366	May 27, 2019	DP			
	6245766	Dec 18, 2018	U-601			
<b>ZIPRASIDONE HYDROCHLORIDE - GEODON</b>						
N 021483 001	6150366	May 27, 2019	DP U-719			
	6245766	Dec 18, 2018	U-601			
	7175855	May 18, 2020	DP			
<b>ZOLEDRONIC ACID - ZOMETA</b>						
N 021223 002	8324189	May 29, 2025	U-1308			
	8324189	May 29, 2025	U-1309			
	8324189	May 29, 2025	U-53			
<b>ZOLEDRONIC ACID - ZOMETA</b>						
N 021223 003	7932241	Feb 05, 2028	DP			
	8324189	May 29, 2025	U-1308			
	8324189	May 29, 2025	U-1309			
	8324189	May 29, 2025	U-53			
<b>ZOLEDRONIC ACID - RECLAST</b>						
N 021817 001	7932241	Feb 05, 2028	DP			
	8052987	Oct 27, 2023	U-1199			
<b>ZOLMITRIPTAN - ZOMIG</b>						
N 021450 003	6750237	Nov 28, 2020	DP		NPP	Jun 12, 2018
	6750237*PED	May 28, 2021	DP			
	7220767	Nov 28, 2020	DP			
	7220767*PED	May 28, 2021	DP			
<b>ZOLMITRIPTAN - ZOMIG</b>						
N 021450 004	6750237	Nov 28, 2020	DP		NPP	Jun 12, 2018
	7220767	Nov 28, 2020	DP			
<b>ZOLPIDEM TARTRATE - AMBIEN CR</b>						
N 021774 001	6514531	Dec 01, 2019	DP			
<b>ZOLPIDEM TARTRATE - AMBIEN CR</b>						
N 021774 002	6514531	Dec 01, 2019	DP			
<b>ZOLPIDEM TARTRATE - EDLUAR</b>						
N 021997 001	6761910	Sep 24, 2019	DP U-674			
	8512747	Sep 24, 2019	U-674			
	9265720	Feb 25, 2031	U-674			
	9597281	Apr 06, 2027	U-674			

**PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST**

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<b><u>ZOLPIDEM TARTRATE - EDLUAR</u></b>						
N 021997 001	6761910	Sep 24, 2019	DP	U-674		
	8512747	Sep 24, 2019		U-674		
	9265720	Feb 25, 2031		U-674		
	9597281	Apr 06, 2027		U-674		
<b><u>ZOLPIDEM TARTRATE - EDLUAR</u></b>						
N 021997 002	6761910	Sep 24, 2019	DP	U-674		
	8512747	Sep 24, 2019		U-674		
	9265720	Feb 25, 2031		U-674		
	9597281	Apr 06, 2027		U-674		
<b><u>ZOLPIDEM TARTRATE - ZOLPIMIST</u></b>						
N 022196 001	7632517	Oct 01, 2017		U-70		
	8236285	Aug 07, 2032	DS	DP U-70		
<b><u>ZOLPIDEM TARTRATE - INTERMEZZO</u></b>						
N 022328 001	7658945	Apr 15, 2027	DP	U-1194		
	7682628	Feb 16, 2025		U-1194		
	8242131	Aug 20, 2029		U-1266		
	8252809	Feb 16, 2025	DP			
<b><u>ZOLPIDEM TARTRATE - INTERMEZZO</u></b>						
N 022328 002	7658945	Apr 15, 2027	DP	U-1194		
	7682628	Feb 16, 2025		U-1194		
	8242131	Aug 20, 2029		U-1266		
	8252809	Feb 16, 2025	DP			

## Footnote:

1. Patents are published upon receipt by the Orange book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).

2. Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. They may not be flagged with respect to other claims which may apply.

**PATENT AND EXCLUSIVITY TERMS**

ADB 1 of 117

**PATENT & EXCLUSIVITY ABBREVIATIONS**

CGT	COMPETITIVE GENERIC THERAPY
D	NEW DOSING SCHEDULE (SEE INDIVIDUAL REFERENCES)
GAIN	GAIN EXCLUSIVITY
I	NEW INDICATION (SEE INDIVIDUAL REFERENCES)
M	MISCELLANEOUS EXCLUSIVITY CODES (SEE INDIVIDUAL REFERENCES)
NC	NEW COMBINATION
NCE	NEW CHEMICAL ENTITY
NCE*	NEW CHEMICAL ENTITY (AN ENANTIOMER OF PREVIOUSLY APPROVED RACEMIC MIXTURE. SEE SECTION 505(U) OF THE FEDERAL FOOD AND DRUG COSMETIC ACT).
NDF	NEW DOSAGE FORM
NE	NEW ESTER OR SALT OF AN ACTIVE INGREDIENT
NP	NEW PRODUCT
NP*	NEW PRODUCT (MINT FLAVORED)
NPP	NEW PATIENT POPULATION
NR	NEW ROUTE
NS	NEW STRENGTH
ODE	ORPHAN DRUG EXCLUSIVITY (SEE INDIVIDUAL REFERENCES)
PC	PATENT CHALLENGE
PED	PEDIATRIC EXCLUSIVITY
RTO	RX TO OTC SWITCH OR OTC USE
RTO*	OTC USE FOR WOMEN AGES 15 AND 16
RTO**	OTC USE FOR WOMEN 14 AND BELOW
U	PATENT USE CODE (SEE INDIVIDUAL REFERENCES)
W	EXCLUSIVITY ON THIS APPLICATION EXPIRING ON THIS DATE HAS BEEN WAIVED BY SPONSOR - SEE SECTION 1.8 OF ORANGE BOOK PREFACE WAIVED EXCLUSIVITY

**EXCLUSIVITY DOSING SCHEDULE**

D-1	ONCE A DAY APPLICATION
D-2	ONCE DAILY DOSING
D-3	SEVEN DAYS/SEVEN DAYS/SEVEN DAYS DOSING SCHEDULE
D-4	SEVEN DAYS/FOURTEEN DAYS DOSING SCHEDULE
D-5	TEN DAYS/ELEVEN DAYS DOSING SCHEDULE
D-6	SEVEN DAYS/NINE DAYS/FIVE DAYS DOSING SCHEDULE
D-7	BID DOSING
D-8	INTRAVENOUS, EPIDURAL AND INTRATHECAL DOSING
D-9	NARCOTIC OVERDOSE IN ADULTS
D-10	NARCOTIC OVERDOSE IN CHILDREN
D-11	POSTOPERATIVE NARCOTIC DEPRESSION IN CHILDREN
D-12	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE DUODENAL ULCER
D-13	INCREASED MAXIMUM DAILY DOSAGE RECOMMENDATION
D-14	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
D-15	SINGLE DAILY DOSE OF 25MG/37.5MG
D-16	CONTINUOUS INTRAVENOUS INFUSION
D-17	400MG EVERY 12 HOURS FOR THREE DAYS FOR UNCOMPLICATED URINARY TRACT INFECTIONS
D-18	LOWER RECOMMENDED STARTING DOSE GUIDELINES
D-19	BOLUS DOSING GUIDELINES
D-20	SINGLE 32MG DOSE
D-21	ALTERNATIVE DOSAGE OF 300MG ONCE DAILY AFTER THE EVENING MEAL
D-22	REDUCTION IN INFUSION TIME FROM 24 TO 4 HOURS FOR THE 60MG DOSE
D-23	INCREASE MAXIMUM DOSE AND VARIATIONS IN THE DOSING REGIMEN
D-24	FOR OVARIAN CANCER THE RECOMMENDED REGIMEN IS 135MG/M2 OR 175MG/M2 INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS
D-25	ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN

**PATENT AND EXCLUSIVITY TERMS**

ADB 2 of 117

**EXCLUSIVITY DOSING SCHEDULE**

- D-26 ONCE WEEKLY APPLICATION
- D-27 BID DOSING IN PATIENTS 12 YEARS OF AGE AND OLDER FOR PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH MODERATE EMETOGENIC CANCER CHEMOTHERAPY
- D-28 USE OF ISOVUE-370 IN EXCRETORY UROGRAPHY AT EQUIVALENT GRAMS OF IODINE TO THE CURRENTLY APPROVED ISOVUE-250 AND ISOVUE-300
- D-29 INCREASE OF CUMULATIVE DOSE TO 0.3MMOL/KG FOR MRI OF CNS IN ADULTS
- D-30 5000 IU DOSE FOR PROPHYLAXIS AGAINST DEEP VEIN THROMBOSIS
- D-31 CHANGE IN RECOMMENDED TOTAL DAILY DOSE TO 80MG (40MG BID)
- D-32 REMOVAL OF THE RESTRICTIONS LIMITING TREATMENT TO TWO CONSECUTIVE WEEKS AND TO SMALL AREAS
- D-33 ONCE DAILY DOSING FOR PLAQUE PSORIASIS
- D-34 EVERY FOUR MONTHS DOSAGE REGIMEN
- D-35 FOR A ONE WEEK DOSING OF INTERDIGITAL TINEA PEDIS
- D-36 FOR A SINGLE 2MG DOSE AS AN ALTERNATIVE TO THE 1MG DOSE GIVEN TWICE DAILY
- D-37 DOSING REGIMEN FOR ADMINISTRATION EITHER ONCE DAILY (QD) OR TWICE DAILY (BID)
- D-38 CONTINUOUS INFUSION AS AN ALTERNATE METHOD OF ADMINISTRATION
- D-39 CHANGE IN TIME TO TAKE THE DRUG PRIOR TO A MEAL TO PREVENT MEAL-INDUCED HEARTBURN SYMPTOMS FROM "...1/2 TO 1 HOUR BEFORE EATING" TO "... RIGHT BEFORE EATING OR UP TO 60MIN BEFORE CONSUMING..."
- D-40 ONCE-A-DAY DOSING REGIMEN
- D-41 DRUG MAY BE DOSED RIGHT BEFORE A MEAL OR ANY TIME UP TO 30MIN BEFORE EATING OR DRINKING FOOD AND BEVERAGES THAT WOULD BE EXPECTED TO CAUSE SYMPTOMS
- D-42 TEN DAY DOSING REGIMEN FOR TRIPLE THERAPY, PREVACID IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN, FOR THE ERADICATION OF H.PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- D-43 INITIATION OF TREATMENT WITH 900MG/DAY BY DELETION OF THE REQUIREMENT TO TITRATE TO 900MG/DAY OVER A 3-DAY PERIOD
- D-44 IN A CLINICAL TRIAL, FEWER DISCONTINUATIONS DUE TO ADVERSE EVENTS, ESPECIALLY DIZZINESS AND VERTIGO, WERE OBSERVED WHEN TITRATING THE DOSE IN INCREMENTS OF 50MG/DAY EVERY 3 DAYS UNTIL AN EFFECTIVE DOSE (NOT EXCEEDING 400MG/DAY) WAS REACHED
- D-45 ONCE DAILY DOSING FOR MAINTENANCE ONLY
- D-46 NEW DOSING REGIMEN OF 80MG DAILY
- D-47 PREVENTION OF HEARTBURN SYMPTOMS WHEN ADMINISTERED FROM 15 MINUTES UP TO, BUT NOT INCLUDING, 1 HOUR PRIOR TO A PROVOCATIVE MEAL
- D-48 ADMINISTRATION OF CISATRACURIUM A NEUROMUSCULAR BLOCKING AGENT AT DOSES OF 3 AND 4X THE ED95 OF CISATRICURIUM FOLLOWING INDUCTION WITH THIOPENTAL
- D-49 PEDIATRIC DOSING GUIDELINES
- D-50 INFORMATION FOR USE OF CORVERT IN POST-CARDIAC SURGERY PATIENTS
- D-51 OPTIONAL STARTING DOSE OF 40MG/DAY
- D-52 ALTERNATE DOSING REGIMEN OF 1250MG TWICE DAILY
- D-53 USE IN PEDIATRIC PATIENTS FROM 1 MONTH TO 16 YEARS OF AGE
- D-54 USE OF ZYBAN FOR MAINTENANCE THERAPY. TREATMENT UP TO 6 MONTHS WAS SHOWN EFFICACIOUS
- D-55 ADDITION OF A HIGHER DOSE OF NUTROPIN FOR PUBERTAL PATIENTS (PUBERTAL DOSE LESS THAN OR EQUAL TO 0.7MG/KG/WEEK)
- D-56 ADDITION OF POSTPRANDIAL DOSING
- D-57 3-HOUR INFUSION OF TAXOL GIVEN EVERY THREE WEEKS AT A DOSE OF 175MG/M<sup>2</sup> FOLLOWED BY CISPLATIN AT A DOSE OF 75MG/M<sup>2</sup> FOR THE FIRST-LINE TREATMENT OF ADVANCED OVARIAN CANCER
- D-58 CHANGE IN DOSING INTERVAL TO ONCE-DAILY ADMINISTRATION
- D-59 REDUCTION OF ELEVATED LDL-C IN A NEW, HIGHER STRENGTH TABLET, 0.8MG, AND FOR EXTENSION OF THE DOSAGE RANGE TO 0.8MG DAILY

**PATENT AND EXCLUSIVITY TERMS**

ADB 3 of 117

**EXCLUSIVITY DOSING SCHEDULE**

- D-60 ADDITION OF A POST-OPERATIVE DOSING REGIMEN
- D-61 ONCE WEEKLY DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-62 ONCE WEEKLY DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-63 TO ALLOW A TITRATION DOSING REGIMEN USING A 25MG DOSE
- D-64 INCREASING DOSAGE FOR NERVE BLOCK ANESTHESIA USING NAROPIN 7.5MG/ML AND FOR EXTENDING THE DURATION OF TREATMENT FOR POSTOPERATIVE ANALGESIA USING NAROPIN 2MG/ML
- D-65 CHANGE DOSING AND ADMINISTRATION TO INDICATE MAINTENANCE OF WEIGHT LOSS OVER AN 18 MONTH PERIOD THUS EXTENDING THE USE OF THIS DRUG FROM ONE TO TWO YEARS
- D-66 DOSING RECOMMENDATIONS FOR PATIENTS UNDERGOING PCI
- D-67 SHORTER TREATMENT COURSE OF THREE DAYS IN THE TREATMENT OF RECURRENT EPISODES OF GENITAL HERPES
- D-68 CHANGE OF ADMIN RATE FOR INFUSION OF AREDIA FOR TREATMENT OF MODERATE AND SEVERE HYPERCALCEMIA OF MALIGNANCY FROM 24 HOURS TO 2 HOURS UP TO BUT NOT INCLUDING 24 HOURS
- D-69 SHORTENED DOSING REGIMEN TO 5 DAYS FOR THE TREATMENT OF ACUTE EXACERBATION OF CHRONIC BRONCHITIS
- D-70 80MG ONCE DAILY DOSING REGIMEN
- D-71 EIGHT WEEK DOSING REGIMEN
- D-72 INFORMATION REGARDING INCREASED RATE OF INFUSION FOR DEPACON
- D-73 ONCE A WEEK DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-74 ONCE A WEEK DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-75 INTERMITTENT DOSING REGIMEN, STARTING DAILY DOSE 14 DAYS PRIOR TO THE ANTICIPATED ONSET OF MENSTRUATION THROUGH THE FIRST FULL DAY OF MENSES AND REPEATING WITH EACH NEW CYCLE
- D-76 FOR USE ON AN "AS NEEDED" OR PRN BASIS FOR THE MANAGEMENT OF NASAL SYMPTOMS IN PATIENTS FOR WHOM THE DRUG IS INDICATED
- D-77 ADDITION OF 20MG AND 40MG DAILY AS OPTIONAL STARTING DOSES WITH 40MG INTENDED FOR PATIENTS WHO REQUIRE A LARGE REDUCTION IN LDL-C (MORE THAN 45%)
- D-78 USE OF FLEXERIL 5MG FOR THE RELIEF OF MUSCLE SPASM ASSOCIATED WITH ACUTE, PAINFUL, MUSCULOSKELETAL CONDITIONS
- D-79 NEW LOWER STARTING DOSE FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS AND/OR MODERATE TO SEVERE SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED W/ THE MENOPAUSE
- D-80 CHANGE OF DOSING SCHEDULE FOR LANTUS FROM ONCE DAILY AT BEDTIME TO FLEXIBLE DAILY DOSING
- D-81 NEW LOWER STARTING DOSE FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-82 USE OF PREMARIN 0.3 MG AND 0.45 MG FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-83 750 MG, ONCE DAILY FOR 5 DAYS FOR COMMUNITY ACQUIRED PNEUMONIA (CAP)
- D-84 ONCE-A-DAY DOSING OF FLOXACIN OTIC FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS(AGES 6 MO & OLDER) W/ OTITIS EXTERNA CAUSED BY SUSCEPTIBLE STRAINS OF E.COLI, P.AERUGINOSA AND S.AUREUS
- D-85 LOWER RECOMMENDED STARTING DOSE GUIDELINES FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE
- D-86 FOR USE IN SELECT EXTERNAL INSULIN PUMPS
- D-87 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- D-88 NEW DOSING RANGE OF 200-400MG PER DAY IN TWO DIVIDED DOSES FOR ADULTS WITH PARTIAL SEIZURES
- D-89 USE OF REYATAZ 300 MG/RITONAVIR 100 MG ONCE DAILY FOR TREATMENT IN HIV-INFECTED ANTIRETROVIRAL-EXPERIENCED PATIENTS
- D-90 ADDITION OF DAYTIME ADMINISTRATION TO TREAT VULVOVAGINAL CANDIDIASIS
- D-91 ALTERNATE INTERMITTENT DOSING REGIMEN

**PATENT AND EXCLUSIVITY TERMS**

ADB 4 of 117

**EXCLUSIVITY DOSING SCHEDULE**

- D-92 ALTERNATIVE DOSAGE OF 1000MG ONCE DAILY AT BEDTIME
- D-93 ALTERNATE TWO OR THREE TIMES DAILY DOSING REGIMENS
- D-94 NEW MAXIMUM DOSAGE OF 72 MG/DAY IN ADOLESCENTS 13-17 YEARS OF AGE WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- D-95 BROADENED INITIAL STARTING DOSE FOR HYPERTENSION FROM 50 MG TO 100 MG TO 25 MG TO 100 MG DOSE RANGE
- D-96 ONCE-MONTHLY TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS WITH BONIVA (IBANDRONATE SODIUM) 150 MG TABLETS
- D-97 PED CANCER PT POPULATION EXPANDED TO INCLUDE PTS 6 MOS UP TO BUT NOT INCLUDING 4 YRS AND DOSING INSTRUCTIONS TO ADMIN 30 MIN BEFORE CHEMO WITH SECOND AND THIRD DOSES 4 & 8 HOURS AFTER FIRST DOSE
- D-98 DOSING FOR PED SURGICAL PTS EXPANDED TO INCLUDE PTS 1 MONTH UP TO BUT NOT INCLUDING 2 YEARS OF AGE
- D-99 ONCE DAILY ADMINISTRATION FOR THE TREATMENT OF HIV INFECTION IN THERAPY NAIVE ADULT PATIENTS
- D-100 750 MG ONCE DAILY FOR FIVE DAYS FOR THE TREATMENT OF ACUTE BACTERIAL SINUSITIS
- D-101 ONCE DAILY IN CHRONIC IDIOPATHIC URTICARIA FOR ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- D-102 NEW DOSING REGIMENT OF ONE SPRAY TWICE DAILY FOR SEASONAL ALLERGIC RHINITIS IN PATIENTS 12 YRS OF AGE AND OLDER
- D-103 NEW DOSING RECOMMENDATION FOR THE TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT PATIENTS, SPECIFICALLY A REDUCTION IN COURSE OF THERAPY FROM FAMCICLOVIR 125 MG TWICE-A-DAY FOR 5 DAYS TO 1000 MG TWICE-A-DAY FOR 1 DAY.
- D-104 0.5MG/0.1MG FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE IN WOMEN WHO HAVE A UTERUS
- D-105 USE OF ACTONEL 75MG TWO CONSECUTIVE DAYS PER MONTH FOR THE PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-106 FIVE DAY TREATMENT OF SELECTED SUSCEPTIBLE STRAINS OF STREPTOCOCCUS PNEUMONIAE, HAEMOPHILUS INFLUENZA, MYCOPLASMA PNEUMONIAE, AND CHLAMYDIA PNEUMONIAE FOR COMMUNITY-ACQUIRED PNEUMONIA
- D-107 PROVIDES FOR THE COMBINATION TABLET OF 70MG ALENDRONATE AND 5600 IU OF VITAMIN D3 FOR THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- D-108 TREATMENT OF COMPLICATED URINARY TRACT INFECTION AND ACUTE PYELONEPHRITIS WITH LEVAQUIN 750MG ONCE DAILY FOR FIVE DAYS
- D-109 PROVIDE FOR THE USE OF A LOWER DOSE FOR THE TREATMENT OF ADULTS WITH CHRONIC PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE OR INTOLERANCE TO PRIOR THERAPY INCLUDING IMATINIB MESYLATE
- D-110 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGED 13-17
- D-111 PROVIDES FOR ONCE DAILY USE OF CIALIS, 2.5 MG AND 5 MG, FOR THE TREATMENT OF ERECTILE DYSFUNCTION
- D-112 PROVIDES FOR PEDIATRIC PUMP USE
- D-113 ONCE DAILY DOSING REGIMENT FOR PATIENTS WHO BECOME CONSTIPATED ON TWICE DAILY REGIMENT
- D-114 NEW DOSING RECOMMENDATIONS FOR USE OF SIROLIMUS IN COMBINATION WITH CYCLOSPORINE FOR THE PROPHYLAXIS OF REJECTION IN HIGH-RISK RENAL TRANSPLANT RECIPIENTS
- D-115 STARTING DOSE OF 15MG/DAY FOR MONOTHERAPY IN ACUTE TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED
- D-116 ALTERNATIVE DOSING REGIMENT ATAZANAVIR SULATE CO-ADMINISTERED WITH RITONAVIR FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT NAIVE PATIENTS
- D-117 50 MG TABLET FOR INITIATION OF DOSE TITRATION FOR BIPOLAR DISORDER
- D-118 TWO 400MG TABLETS ONCE DAILY, CO-ADMINISTERED WITH 100MG RITONAVIR
- D-119 DOSING RECOMMENDATIONS FOR HIV INFECTED PEDIATRIC PATIENTS 6 TO LESS THAN 18 YEARS OF AGE
- D-120 DOSING REGIMENT ADJUSTMENTS
- D-121 CHANGE TO REMOVE 20 MG MAXIMUM DOSAGE RESTRICTION

**PATENT AND EXCLUSIVITY TERMS**

ADB 5 of 117

**EXCLUSIVITY DOSING SCHEDULE**

- D-122 USE OF VAGIFEM 10 MCG FOR THE TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE
- D-123 ALTERNATIVE DOSING REGIMEN DOSE OF 20 MG/METER SQUARE BY CONTINUOUS INTRAVENOUS INFUSION OVER 1 HOUR REPEATED DAILY FOR 5 DAYS
- D-124 ONCE DAILY DOSING REGIMEN IN ADULT PATIENTS WITH LESS THAN THREE LOPINAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
- D-125 EXTEND CURRENT DOSING REGIMEN TO 900MG (2-450MG TABLETS) ONCE A DAY WITHIN 10 DAYS OF TRANSPLANTATION UNTIL 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN ADULT KIDNEY TRANSPLANT PATIENTS AT HIGH RISK.
- D-126 CHANGE DOSAGE REGIMEN FROM 250MG TO 500MG
- D-127 DOSING REGIMEN FOR ADULT PATIENTS WITH CHRONIC HEPATITIS B (CHB) AND DECOMPENSATED LIVER DISEASE
- D-128 SINGLE IV DOSE OF FOSAPREPITANT 150MG, DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID, FOR PREVENTION OF ACUTE & DELAYED NAUSEA & VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY EMETOGENIC CANCER CHEMO
- D-129 800/100 MG DARUNAVIR/RITONAVIR, ONCE DAILY, IN TREATMENT-EXPERIENCED HIV-1 INFECTED PATIENTS WITH NO DARUNAVIR RESISTANCE ASSOCIATED SUBSTITUTIONS
- D-130 DOSING RECOMMENDATIONS FOR TREATMENT OF HIV-1 INFECTION DURING PREGNANCY BASED ON DATA FROM STUDY AI424-182, A STUDY OF ATAZANAVIR/RITONAVIR IN COMBINATION WITH ZIDOVUDINE/LAMIVUDINE IN HIV INFECTED PREGNANT WOMEN
- D-131 EVERY 6 TO 8 WEEKS FOR THE 120MG STRENGTH FOR PATIENTS WHO ARE CONTROLLED ON SOMATULINE DEPOT 60MG OR 90MG
- D-132 45MG FOR 6 MONTH ADMINISTRATION
- D-133 NEW EFFICACY DATA AND DOSING REGIMEN FOR PREGNANCY IN NORMAL OVULATORY WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION AS PART OF AN IVF OR INTRACYTOPLASMIC SPERM INJECTION (ICSI) CYCLE
- D-134 INCREASING MAXIMUM DOSING OF PATIENTS WITH SCHIZOPHRENIA TO 160 MG/DAY
- D-135 UPDATE LABELING WITH ONCE DAILY DOSING IN HIV-1 INFECTED, TREATMENT-NAIVE PEDIATRIC PATIENTS 12 TO LESS THAN 18 YEARS OF AGE
- D-136 ALTERNATE DOSING REGIMEN FOR UNCOMPLICATED URETHRAL OR ENDOCERVICAL INFECTION CAUSED BY CHLAMYDIA TRACHOMATIS, ADMINISTER 200 MG BY MOUTH ONCE-A-DAY FOR 7 DAYS
- D-137 NEW LOWER DOSING REGIMEN FOR REVATIO IN THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP 1) IN ADULTS
- D-138 80 MG DOSING REGIMEN FOR THE RISK REDUCTION OF REBLEEDING OF GASTRIC AND DUODENAL ULCERS IN THE FIRST 72 HOURS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
- D-139 ADDITIONAL INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE LABELING REGARDING THE ADMINISTRATION OF BRAVELLE AND MENOPUR IN THE SAME SYRINGE TO OVULATORY WOMEN AS PART OF AN ART CYCLE
- D-140 REVISED DOSING SCHEDULE TO ADMINISTER AVANAFIL 15 MINUTES PRIOR TO SEXUAL ACTIVITY
- D-141 DOSING INFORMATION IN PREVIOUSLY UNTREATED MANTLE CELL LYMPHOMA
- D-142 DOSE MODIFICATION GUIDELINES FOR BORTEZOMIB WHEN GIVEN IN COMBINATION WITH RITUXIMAB, CYCLOPHOSPHAMIDE, DOXORUBICIN, AND PREDNISONE
- D-143 INITIATION OF VIMPAT THERAPY WITH A LOADING DOSE OF 200MG
- D-144 LOWER LIMIT OF 15 MINUTES FOR THE INFUSION DURATION
- D-145 UPDATES TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING TO REFLECT THE RESULTS OF TWO SHORT TERM STUDIES EVALUATING THE SAFETY AND EFFICACY OF INTUNIV IN CHILDREN AND ADOLESCENTS AGES 6 TO 17 WITH ADHD.
- D-146 CHANGE IN TARGET DOSING TO 20MG TO 40MG ORALLY ONCE DAILY
- D-147 ONCE DAILY DOSING IN PEDIATRIC PATIENTS 3 MONTHS OF AGE AND OLDER IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION
- D-148 EXTENDED THE DURATION OF THE DOSING REGIMEN FROM 100 DAYS TO 200 DAYS POST-TRANSPLANTATION FOR THE PREVENTION OF CMV DISEASE IN PEDIATRIC KIDNEY TRANSPLANT
- D-149 DOSING INFORMATION ADDED TO THE LABELING REGARDING PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH ITP

**PATENT AND EXCLUSIVITY TERMS**

ADB 6 of 117

**EXCLUSIVITY DOSING SCHEDULE**

- D-150 1600MG DAILY FOR PATIENTS ON ADJUNCTIVE THERAPY WHO DID NOT ACHIEVE A SATISFACTORY RESPONSE ON 1200MG DAILY DOSE
- D-151 DOSING RECOMMENDATIONS FOR THE TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS CO-INFECTED WITH HIV-1
- D-152 DOSING RECOMMENDATIONS AS NECESSARY FOR FEVER AND PAIN FOR AGES 6MO TO LESS THAN 12 YEARS AND 12 TO 17 YEARS.
- D-153 IN COMBINATION WITH RIBAVIRIN FOR 12 WEEKS, FOR THE TREATMENT OF GENOTYPE 1, CHRONIC HEPATITIS C TREATMENT EXPERIENCED PATIENTS WITH COMPENSATED CIRRHOSIS BASED UPON THE RESULTS OF THE SIRIUS STUDY
- D-154 ADDITION OF A 1500MG-SINGLE-DOSE REGIMEN FOR THE TREATMENT OF ADULT PATIENTS WITH ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS (ABSSI)
- D-155 SINGLE IV DOSE OF FOSAPREPITANT 150MG DOSED CONCOMITANTLY WITH 5HT3 RECEPTOR ANTAGONIST & CORTICOSTEROID FOR PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- D-156 DOSING INFORMATION ADDED TO THE LABELING PROVIDING INFORMATION ON TRANSITIONING FROM SUBCUTANEOUS OR INTRAVENOUS ROUTES OF ADMINISTRATION OF TREPROSTINIL
- D-157 UPDATED INFORMATION ADDED TO THE DOSAGE AND ADMINISTRATION SECTION OF THE LABELING PROVIDING DOSAGE RECOMMENDATIONS FOR INTERRUPTIONS AND DISCONTINUATION OF THERAPY
- D-158 REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 1 HCV INFECTION
- D-159 REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE LIVER TRANSPLANT RECIPIENTS WITH GENOTYPE 4 HCV INFECTION
- D-160 REVISED DOSING TO EXPAND PATIENT POPULATION TO INCLUDE PATIENTS WITH DECOMPENSATED CIRRHOSIS WITH GENOTYPE 1 HCV INFECTION
- D-161 DOSAGE RECOMMENDATIONS ADDED TO INCLUDE TREATMENT OF HCV GENOTYPE 3 SUBJECTS CO-INFECTED WITH HIV-1
- D-162 DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1 INFECTION WITH COMPENSATED (CHILD-PUGH A) OR DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS AND TREATMENT OF CHRONIC HCV GENOTYPE 3 INFECTION IN SUBJECTS WITH DECOMPENSATED (CHILD-PUGH B OR C) CIRRHOSIS
- D-163 DOSING TO INCLUDE PATIENTS WITH CHRONIC HCV GENOTYPE 1A INFECTION WITH COMPENSATED (CHILD-PUGH A) CIRRHOSIS AND GENOTYPE 1B WITH OR WITHOUT COMPENSATED (CHILD-PUGH A) CIRRHOSIS
- D-164 UPDATES TO THE DOSAGE AND ADMINISTRATION, DOSE MODIFICATIONS SECTION OF THE LABELING
- D-165 DOSING RECOMMENDATION ADDED TO THE LABELING FOR IMBRUVICA USE IN COMBINATION WITH BENDAMUSTINE AND RITUXIMAB FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)/SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- D-166 BROADEN INITIAL STARTING DOSE FOR BIPOLAR I DISORDER TO 5-10MG TWICE DAILY
- D-167 ADDITION OF 1200 MG ONCE DAILY DOSING FOR TREATMENT-NAÏVE PATIENTS OR PATIENTS WHO ARE VIROLOGICALLY SUPPRESSED ON AN INITIAL REGIMEN OF Raltegravir FILM-COATED TABLETS 400 MG TWICE DAILY
- D-168 NEW DOSING REGIMEN OF 10 MG ONCE DAILY FOR THE REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR DVT AND/OR PE AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
- D-169 ONCE-DAILY DOSING FOR PATIENTS 5 YEARS OF AGE AND OLDER WHO HAVE UNDETECTABLE SERUM AND URINE SUCCINYLACTIONE CONCENTRATIONS AFTER A MINIMUM OF 4 WEEKS ON A STABLE DOSAGE OF NITISINONE

**EXCLUSIVITY INDICATION**

- I-1 DYSMENORRHEA
- I-2 CHOLANGIOPANCREATOGRAPHY
- I-3 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
- I-4 PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)

**PATENT AND EXCLUSIVITY TERMS**

ADB 7 of 117

**EXCLUSIVITY INDICATION**

- I-5 HYSTEROSALPINGOGRAPHY  
I-6 TREATMENT OF JUVENILE ARTHRITIS  
I-7 BIOPSY PROVEN MINIMAL CHANGE NEPHROTIC SYNDROME IN CHILDREN  
I-8 ADULT INTRAVENOUS CONTRAST-ENHANCED COMPUTED TOMOGRAPHY OF THE HEAD AND BODY  
I-9 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING  
I-10 PREVENTION OF POSTOPERATIVE DEEP VENOUS THROMBOSIS AND PULMONARY EMBOLISM IN TOTAL HIP REPLACEMENT SURGERY  
I-11 RELIEF OF MILD TO MODERATE PAIN  
I-12 TREATMENT OF CUTANEOUS CANDIDIASIS  
I-13 URINARY TRACT INFECTION (UTI) PREVENTION FOR PERIODS UP TO FIVE MONTHS IN WOMEN WITH A HISTORY OF RECURRENT UTI  
I-14 SEBORRHEIC DERMATITIS  
I-15 PHOTOPHERESIS IN THE PALLIATIVE TREATMENT OF SKIN MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PERSONS NOT RESPONSIVE TO OTHER TREATMENT  
I-16 STIMULATE THE DEVELOPMENT OF MULTIPLE FOLLICLES/OOCYTES IN OVULATORY PATIENTS PARTICIPATING IN AN IN VITRO FERTILIZATION PROGRAM  
I-17 MANAGEMENT OF CONGESTIVE HEART FAILURE  
I-18 ENDOSCOPIC RETROGRADE PANCREATOGRAPHY  
I-19 HERNIOGRAPHY  
I-20 KNEE ARTHROGRAPHY  
I-21 HIGH DOSE METHOTREXATE WITH LEUCOVORIN RESCUE IN COMBINATION WITH OTHER CHEMOTHERAPEUTIC AGENTS TO DELAY RECURRENCE IN PATIENTS WITH NONMETASTATIC OSTEOSARCOMA WHO HAVE UNDERGONE SURGICAL RESECTION OR AMPUTATION FOR THE PRIMARY TUMOR  
I-22 RESCUE AFTER HIGH-DOSE METHOTREXATE THERAPY IN OSTEOSARCOMA  
I-23 SHORT-TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER  
I-24 TREATMENT OF RHEUMATOID ARTHRITIS  
I-25 ADULT INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY OF THE HEAD, NECK, ABDOMINAL, RENAL AND PERIPHERAL VESSELS  
I-26 TREATMENT OF LIVER FLUKES  
I-27 ADJUNCTIVE THERAPY TO DIET TO REDUCE THE RISK OF CORONARY ARTERY DISEASE  
I-28 SELECTIVE ADULT VISCELAR ARTERIOGRAPHY  
I-29 METASTATIC BREAST CANCER IN PREMENOPAUSAL WOMEN AS AN ALTERNATIVE TO OOPHORECTOMY OR OVARIAN IRRADIATION  
I-30 TREATMENT OF TINEA PEDIS  
I-31 CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS IN THE SPINE AND ASSOCIATED TISSUES  
I-32 PEDIATRIC MYELOGRAPHY  
I-33 ORAL USE OF DILUTED OMNIPAQUE INJECTION IN ADULTS FOR CONTRAST ENHANCED COMPUTED TOMOGRAPHY OF THE ABDOMEN  
I-34 ORAL USE IN ADULTS FOR PASS-THROUGH EXAMINATION OF THE GASTROINTESTINAL TRACT  
I-35 PEDIATRIC CONTRAST ENHANCEMENT OF COMPUTED TOMOGRAPHIC HEAD IMAGING  
I-36 ARTHROGRAPHY OF THE SHOULDER JOINTS IN ADULTS  
I-37 RADIOGRAPHY OF THE TEMPOROMANDIBULAR JOINT IN ADULTS  
I-38 CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS OF THE CENTRAL NERVOUS SYSTEM IN CHILDREN (2 YEARS OF AGE AND OLDER)  
I-39 TREATMENT OF ACUTE MYOCARDIAL INFARCTION  
I-40 PRIMARY NOCTURNAL ENURESIS  
I-41 MIGRAINE HEADACHE PROPHYLAXIS  
I-42 HERPES ZOSTER

**PATENT AND EXCLUSIVITY TERMS**

ADB 8 of 117

**EXCLUSIVITY INDICATION**

- I-43 HERPES SIMPLEX ENCEPHALITIS
- I-44 MAINTENANCE THERAPY IN HEALED DUODENAL ULCER PATIENTS AT DOSE OF 1 GRAM TWICE DAILY
- I-45 ACUTE TREATMENT OF VARICELLA ZOSTER VIRUS
- I-46 USE IN PEDIATRIC COMPUTED TOMOGRAPHIC HEAD AND BODY IMAGING
- I-47 TREATMENT OF PEDIATRIC PATIENTS WITH SYMPTOMATIC HUMAN IMMUNODEFICIENCY VIRUS (HIV) DISEASE
- I-48 PEDIATRIC ANGIOCARDIOGRAPHY
- I-49 TREATMENT OF TRAVELERS' DIARRHEA DUE TO SUSCEPTIBLE STRAINS OF ENTEROTOXIGENIC ESCHERICHIA COLI
- I-50 FOR USE IN WOMEN WITH AXILLARY NODE-NEGATIVE BREAST CANCER
- I-51 TREATMENT OF PRIMARY DYSMENORRHEA AND FOR THE TREATMENT OF IDIOPATHIC HEAVY MENSTRUAL BLOOD LOSS
- I-52 PEDIATRIC EXCRETORY UROGRAPHY
- I-53 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA
- I-54 RENAL CONCENTRATION CAPACITY TEST
- I-55 HYPERTENSION
- I-56 EROSION GASTROESOPHAGEAL REFLUX DISEASE
- I-57 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER
- I-58 INITIAL TREATMENT OF ADVANCED OVARIAN CARCINOMA IN COMBINATION WITH OTHER APPROVED CHEMOTHERAPEUTIC AGENTS
- I-59 ENDOSCOPICALLY DIAGNOSED ESOPHAGITIS, INCLUDING EROSION AND ULCERATIVE ESOPHAGITIS, AND ASSOCIATED HEARTBURN DUE TO GASTROESOPHAGEAL REFLUX DISEASE
- I-60 SINGLE APPLICATION TREATMENT OF HEAD LICE IN CHILDREN TWO MONTHS TO TWO YEARS IN AGE
- I-61 FEMALE ANDROGENETIC ALOPECIA
- I-62 PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-63 ONCE DAILY TREATMENT AS INITIAL THERAPY IN THE TREATMENT OF HYPERTENSION
- I-64 PREVENTION OF SUPRAVENTRICULAR TACHYCARDIAS
- I-65 PREVENTION OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- I-66 UNCOMPLICATED GONORRHEA
- I-67 TREATMENT OF ACUTE ASTHMATIC ATTACKS IN CHILDREN SIX YEARS OF AGE AND OLDER
- I-68 CENTRAL PRECOCIOUS PUBERTY
- I-69 SHORT TERM TREATMENT OF PATIENTS WITH SYMPTOMS OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), AND FOR THE SHORT TERM TREATMENT OF ESOPHAGITIS DUE TO GERD INCLUDING ULCERATIVE DISEASE DIAGNOSED BY ENDOSCOPY
- I-70 USE IN COMBINATION WITH 5-FLUOROURACIL TO PROLONG SURVIVAL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED COLORECTAL CANCER
- I-71 VARICELLA INFECTIONS (CHICKENPOX)
- I-72 PREVENTION OF CMV DISEASE IN TRANSPLANT PATIENTS AT RISK FOR CMV DISEASE
- I-73 INITIATE AND MAINTAIN MONITORED ANESTHESIA CARE (MAC) SEDATION DURING DIAGNOSTIC PROCEDURES
- I-74 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
- I-75 TREATMENT OF ENDOSCOPICALLY DIAGNOSED EROSION ESOPHAGITIS
- I-76 PREVENTION OF OSTEOPOROSIS
- I-77 DERMAL INFECTIONS-TINEA PEDIS, TINEA CORPORIS, TINEA CRURIS DUE TO EPIDERMOPHYTON FLOCCOSUM
- I-78 CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY AND INTRAVENOUS EXCRETORY UROGRAPHY
- I-79 MANAGEMENT OF CHRONIC STABLE ANGINA AND ANGINA DUE TO CORONARY ARTERY SPASM
- I-80 DIAGNOSIS AND LOCALIZATION OF ISCHEMIA AND CORONARY HEART DISEASE

**PATENT AND EXCLUSIVITY TERMS**

ADB 9 of 117

**EXCLUSIVITY INDICATION**

- I-81 PROPHYLAXIS IN DESIGNATED IMMUNOCOMPROMISED CONDITIONS TO REDUCE THE INCIDENCE OF OROPHARYNGEAL CANDIDIASIS
- I-82 TREATMENT OF TRAVELERS' DIARRHEA
- I-83 ANGIOCARDIOGRAPHY, CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY IN CHILDREN
- I-84 INTRAOPERATIVE AND POSTOPERATIVE TACHYCARDIA AND/OR HYPERTENSION
- I-85 TREATMENT OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- I-86 TREATMENT OF SECONDARY CARNITINE DEFICIENCY
- I-87 RENAL IMAGING AGENT FOR USE IN CHILDREN
- I-88 MANAGEMENT OF ENDOMETRIOSIS
- I-89 EPIDURAL USE IN LABOR AND DELIVERY AS AN ANALGESIC ADJUNCT TO BUPIVACAINE
- I-90 INTENSIVE CARE UNIT SEDATION
- I-91 MONOTHERAPY USE FOR HYPERTENSION
- I-92 ADJUNCTIVE THERAPY IN THE MANAGEMENT OF HEART FAILURE
- I-93 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN CHILDREN AGES 4-11 YEARS
- I-94 USE WITH MRI IN ADULTS TO PROVIDE CONTRAST ENHANCEMENT AND FACILITATE VISUALIZATION OF LESIONS IN THE BODY [EXCLUDING THE HEART]
- I-95 TREATMENT OF LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-96 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA
- I-97 ORAL OR RECTAL USE IN CHILDREN FOR THE EXAMINATION OF THE GASTROINTESTINAL TRACT
- I-98 TREATMENT OF CHILDREN WHO HAVE GROWTH FAILURE ASSOCIATED WITH CHRONIC RENAL INSUFFICIENCY
- I-99 PEDIATRIC ANESTHESIA IN CHILDREN 3 YEARS AND OLDER
- I-100 TO DECREASE THE INCIDENCE OF CANDIDIASIS IN PATIENTS UNDERGOING BONE MARROW TRANSPLANTATION WHO RECEIVE CYTOTOXIC CHEMOTHERAPY AND/OR RADIATION THERAPY
- I-101 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN-DEPENDENT DIABETES MELLITUS AND RETINOPATHY
- I-102 TREATMENT OF OBSESSIVE-COMPULSIVE DISORDER
- I-103 PROPHYLAXIS AGAINST PNEUMOCYSTIS CARINII PNEUMONIA IN INDIVIDUALS WHO ARE IMMUNOCOMPROMISED AND CONSIDERED TO BE AT RISK OF DEVELOPING PNEUMOCYSTIS CARINII PNEUMONIA
- I-104 TREATMENT OF PULMONARY AND EXTRAPULMONARY ASPERGILLOSIS IN PATENTS WHO ARE INTOLERANT OF OR WHO ARE REFRACTORY TO AMPHOTERICIN B THERAPY
- I-105 TREATMENT OF METASTATIC CARCINOMA OF THE BREAST AFTER FAILURE OF FIRST-LINE OR SUBSEQUENT CHEMOTHERAPY
- I-106 TREATMENT OF ACROMEGALY
- I-107 VAGINAL CANDIDIASIS
- I-108 EXPANDED USE-FOR ICU PATIENTS UNDERGOING LONG-TERM INFUSION DURING MECHANICAL VENTILATION
- I-109 TYPHOID FEVER
- I-110 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIOTHERAPY
- I-111 TREATMENT OF PAGET'S DISEASE OF BONE
- I-112 MANAGEMENT OF MODERATE TO SEVERE PAIN
- I-113 TREATMENT OF PROSTATITIS
- I-114 USE IN CHILDREN TO VISUALIZE LESIONS WITH ABNORMAL VASCULARITY IN THE BRAIN (INTRACRANIAL LESIONS), SPINE, AND ASSOCIATED TISSUE
- I-115 USE IN MRI IN ADULTS TO VISUALIZE LESIONS IN THE HEAD AND NECK
- I-116 MAINTENANCE OF HEALING OF EROSIVE ESOPHAGITIS
- I-117 TO SLOW THE PROGRESSION FO CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE
- I-118 PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM

**PATENT AND EXCLUSIVITY TERMS**

ADB 10 of 117

**EXCLUSIVITY INDICATION**

FOLLOWING KNEE REPLACEMENT SURGERY

- I-119 TREATMENT OF ANEMIA CAUSED BY UTERINE LEIOMYOMATA IN WOMEN WHO FAIL IRON THERAPY
- I-120 MAINTENANCE THERAPY FOR GASTRIC ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING ACUTE ULCERS
- I-121 EXPANDED PATIENT POPULATION -- USE IN ICU PATIENTS
- I-122 PSORIASIS OF THE SCALP
- I-123 RELIEF OF MILD TO MODERATE PAIN IN PATIENTS AGED 6 MONTHS AND OLDER
- I-124 LEUCOCYTE LABELED SCINTIGRAPHY AS AN ADJUNCT IN THE LOCALIZATION OF INTRA-ABDOMINAL INFECTION AND INFLAMMATORY BOWEL DISEASE
- I-125 EXPANSION OF CONSCIOUS SEDATION INDICATION TO INCLUDE SHORT THERAPEUTIC PROCEDURES
- I-126 ADJUNCT TO THALLIUM- 201 MYOCARDIAL PERfusion IN PATIENTS UNABLE TO EXERCISE ADEQUATELY
- I-127 TREATMENT OF ACYCLOVIR-RESISTANT HERPES IN IMMUNOCOMPROMISED PATIENTS
- I-128 IN PT W/ CH DISEASE AND HYPERCHOLESTEROLEMIA: REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH; REDUCE RISK NON-FATAL MI; REDUCE RISK UNDERGOING MYOCARDIAL REVASCULARIZATION PROCEDURES; REDUCTION ELEVATED TOTAL AND LDL CHOL LEVELS...
- I-129 TREATMENT OF ALCOHOL DEPENDENCE
- I-130 MAINTENANCE OF HEALING OF EROsIVE ESOPHAGITIS
- I-131 PERIPHERAL ARTERIOGRAPHY
- I-132 TREATMENT OF MANIC PHASE OF BIPOLAR DISORDER
- I-133 MANAGEMENT OF CHRONIC STABLE ANGINA
- I-134 HEART FAILURE POST MYOCARDIAL INFARCTION
- I-135 BONE METASTASES ASSOCIATED WITH MULTIPLE MYELOMA
- I-136 IDIOPATHIC CHRONIC URTICARIA
- I-137 PREVENTION OF METAL-INDUCED HEART BURN, ACID INDIGESTION, AND SOUR STOMACH WHEN TAKEN 30 MINUTES PRIOR TO CONSUMING FOOD OR BEVERAGES
- I-138 TREATMENT OF ACUTE RECURRENT GENITAL HERPES
- I-139 PALLIATIVE TREATMENT OF ADVANCED BREAST CANCER IN PRE- AND PERIMENOPAUSAL WOMEN
- I-140 PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN INDIVIDUALS WITH HIV INFECTION AT RISK FOR DEVELOPING CMV DISEASE
- I-141 TREATMENT OF HEMODYNAMICALLY STABLE PATIENTS WITHIN 24 HOURS OF ACUTE MYOCARDIAL INFARCTION TO IMPROVE SURVIVAL
- I-142 LOCALIZE MYOCARDIAL ISCHEMIA(REVERSIBLE DEFECT) AND INFARCTION (NON-REVERSIBLE DEFECTS) IN EVALUATING MYOCARDIAL FUNCTION
- I-143 EPISODIC TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
- I-144 ENHANCEMENT OF MRI OF THE ADULT BODY INTERNAL ORGANS
- I-145 0.1MMOL/KG AS A SINGLE INTRAVEOUS BOLUS FOR MRI OF THE CNS IN CHILDREN
- I-146 CONTRAST ENHANCEMENT AND FACILITATION OF VISUALIZATION OF EXTRACRANIAL HEAD AND NECK LESIONS
- I-147 PREVENTION OF GALLSTONE FORMATION IN OBESE PATIENTS EXPERIENCING RAPID WEIGHT LOSS
- I-148 TREATMENT OF ACUTE PNEUMOCYSTIC CARINI PNEUMONIA (PCP) IN HIV-INFECTED PATIENTS WHOSE ALVEOLAR-ARTERIAL OXYGEN DIFFERENCE (AaDO<sub>2</sub>) IS LESS THAN OR EQUAL TO 55 TORR
- I-149 TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER
- I-150 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER AND PANIC DISORDER
- I-151 PREVENTION OF AND PREVENTION OF FURTHER POSTOPERATIVE NAUSEA AND VOMITING IN PEDIATRIC PATIENTS RECEIVING GENERAL ANESTHESIA
- I-152 SLOWING THE PROGRESSION OF CORONARY ATHEROSCLEROSIS AND REDUCING THE RISK OF ACUTE CORONARY EVENTS

**PATENT AND EXCLUSIVITY TERMS**

ADB 11 of 117

**EXCLUSIVITY INDICATION**

- I-153 MANAGEMENT OF SEVERE SPASTICITY [ENCOMPASSES SPINAL AND CEREBRAL ORIGIN]
- I-154 PATIENT POPULATION ALTERED TO INCLUDE PEDIATRIC USE
- I-155 TREATMENT OF ONCHOMYCOSIS DUE TO DERMATOPHYTES (TINEA UNGUIUM) OF THE TOENAIL WITH OR WITHOUT FINGERNAIL INVOLVEMENT
- I-156 ADDITIONAL DATA REGARDING THE SAFE USE OF NORVASC IN PATIENTS WITH HEART FAILURE
- I-157 TREATMENT OF ACUTE UNCOMPLICATED CYSTITIS IN FEMALES
- I-158 TREATMENT OF OSTEOLYTIC BONE METASTASES OF BREAST CANCER
- I-159 FOR HYPERCHOLESTEROLEMIC PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE REDUCE THE RISK OF MYOCARDIAL INFARCTION, REVASCULARIZATION, AND DEATH DUE TO CARDIOVASCULAR CAUSES WITH NO INCREASE IN DEATH FROM NON-CARDIOVASCULAR CAUSES
- I-160 TREATMENT OF BACTERIAL CORNEAL ULCERS
- I-161 TREATMENT OF ADULT-ONSET OR CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENCY
- I-162 FOR USE IN PATIENTS 6-11 YEARS OF AGE
- I-163 TREATMENT OF PHOTOPHOBIA
- I-164 CHRONIC BACTERIAL PROSTATITIS
- I-165 MANAGEMENT OF ADULTS WITH ACTIVE, CLASSIC AND DEFINITIVE RHEUMATOID ARTHRITIS WHO HAVE HAD INSUFFICIENT THERAPEUTIC RESPONSE TO OR ARE INTOLERANT OF AN ADEQUATE TRIAL OF FULL DOSES OF ONE OR MORE NON-STEROIDAL ANTI-INFLAMMATORY DRUGS
- I-166 TREATMENT OF BULIMIA
- I-167 COMPLICATED INTRA-ABDOMINAL INFECTIONS (USED IN COMBINATION WITH METRONIDAZOLE) CAUSED BY MIXED AEROBIC/ANAEROBIC PATHOGENS
- I-168 MANAGEMENT OF LOCALLY CONFINED STAGE B2-C METASTATIC CARCINOMA OF THE PROSTATE (IN COMBINATION WITH LHRH AGONISTS)
- I-169 USE IN COMBINATION WITH CORTICOSTEROIDS AS INITIAL CHEMOTHERAPY FOR THE TREATMENT OF PATIENTS WITH PAIN RELATED TO ADVANCED HORMONE-REFRACTORY PROSTATE CANCER
- I-170 PROPHYLACTIC USE DURING HEAD LICE EPIDEMICS
- I-171 RELIEF OF SYMPTOMS OF THE COMMON COLD
- I-172 TREATMENT OF INITIAL EPISODE OF GENITAL HERPES
- I-173 PREOPERATIVELY FOR THE PREVENTION OF INFECTION IN TRANSRECTAL PROSTATE BIOPSY
- I-174 PELVIC INFLAMMATORY DISEASE
- I-175 TREATMENT OF TINEA CORPORIS AND TINEA CRURIS
- I-176 TREATMENT OF POSTOPERATIVE INFLAMMATION IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION
- I-177 TX OF MODERATE ACNE VULGARIS IN FEMALES, GREATER OR EQUAL TO 15YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, DESIRE CONTRACEPTION, HAVE ACHIEVED MENARCHE AND ARE UNRESPONSIVE TO TOPICAL ANTI-ACNE MEDICATIONS
- I-178 TREATMENT OF ONCHOMYCOSIS OF THE FINGERNAIL WITHOUT CONCOMITANT ONCHOMYCOSIS OF THE TOENAIL WITH A PULSE DOSING REGIMEN
- I-179 NOSOCOMIAL PNEUMONIA-MILD TO MODERATE AND SEVERE CAUSED BY HAEMOPHILUS INFLUENZAE OR KLEBSIELLA PNEUMONIAE
- I-180 TREATMENT OF PLANTAR TINEA PEDIS (MOCCASIN TYPE)
- I-181 TREATMENT OF PATIENTS WITH COMPLEX PARTIAL SEIZURES WITH AND WITHOUT SECONDARY GENERALIZATION
- I-182 TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME
- I-183 MAINTENANCE THERAPY IN THE MANAGEMENT OF MILD TO MODERATE ASTHMA IN PEDIATRIC PATIENTS AGES 6-11
- I-184 TREATMENT OF PANIC DISORDER AT A RECOMMENDED DOSE RANGE OF 1 TO 2MG/DAY (MAXIMUM OF 4MG)
- I-185 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-186 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR CAUSED BY OR PRESUMED TO BE CAUSED BY

**PATENT AND EXCLUSIVITY TERMS**

ADB 12 of 117

**EXCLUSIVITY INDICATION**

- I-187 PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE) PREVENTION OF FRACTURES IN THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-188 TREATMENT OF ACUTE SINUSITIS AND ACUTE EXACERBATION OF CHRONIC SINUSITIS
- I-189 TREATMENT OF ACUTE OTITIS MEDIA IN PEDIATRIC PATIENTS
- I-190 PLANAR IMAGING AS A SECOND LINE DIAGNOSTIC DRUG AFTER MAMMOGRAPHY TO ASSIST IN THE EVALUATION OF BREAST LESIONS IN PATIENTS WITH AN ABNORMAL MAMMOGRAM OR A PALPABLE BREAST MASS
- I-191 ENDOMETRIAL THINNING AGENT PRIOR TO ENDOMETRIAL ABLATION FOR DYSFUNCTIONAL UTERINE BLEEDING
- I-192 THE PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS AND A NEW DOSAGE REGIMEN, 40MG ONCE DAILY, FOR THIS INDICATION
- I-193 TREATMENT OF PANIC DISORDER IN A RECOMMENDED DOSE RANGE OF 50 TO 200MG/DAY
- I-194 CONGESTIVE HEART FAILURE
- I-195 FOR USE OF LANSOPRAZOLE IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF DUODENAL ULCER
- I-196 ACUTE TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- I-197 MAINTENANCE OF HEALING OF DUODENAL ULCER
- I-198 FOR THE USE OF LANSOPRAZOLE IN COMBINATION WITH AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF A DUODENAL ULCER
- I-199 MONOTHERAPY AND COMBINATION THERAPY WITH SULFONYLUREA IN THE TREATMENT OF TYPE II DIABETES
- I-200 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR
- I-201 EMPIRICAL THERAPY FOR FEBRILE NEUTROPENIC PATIENTS
- I-202 SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA
- I-203 MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
- I-204 USE IN PEDIATRIC PATIENTS BETWEEN THE AGES OF 6 AND 11 FOR THE TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- I-205 INITIAL ANTICONVULSANT TREATMENT OF STATUS EPILEPTICUS
- I-206 TREATMENT OF EDEMA ASSOCIATED WITH CHRONIC RENAL FAILURE
- I-207 FOR THE SUPPRESSION OF RECURRENT EPISODES OF GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
- I-208 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER IN THE PEDIATRIC POPULATION
- I-209 PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA (PSVT)
- I-210 TO SLOW THE PROGRESSION OF CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL AND LDL CHOLESTEROL TO TARGET LEVELS
- I-211 FOR USE IN PEDIATRIC POPULATION
- I-212 TREATMENT OF SYMPTOMS OF DRY MOUTH IN PATIENTS WITH SJOGREN'S SYNDROME
- I-213 TEMPORARY RELIEF OF PAIN AND PHOTOPHOBIA IN PATIENTS UNDERGOING CORNEAL REFRACTIVE SURGERY
- I-214 TREATMENT OF OSTEOPOROSIS
- I-215 PRE-PROCEDURAL APPLICATION TO ADULT MALE GENITAL SKIN PRIOR TO SITE-SPECIFIC SUBCUTANEOUS INFILTRATION WITH LIDOCAINE FOR THE REMOVAL OF GENITAL WARTS
- I-216 FOR THE LONG-TERM TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-217 PREVENTION (DURING AND FOLLOWING HOSPITALIZATION) OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
- I-218 USE OF LIPITOR AS AN ADJUNCTIVE THERAPY TO DIET FOR THE TREATMENT OF PATIENTS WITH ELEVATED SERUM TRIGLYCERIDE LEVELS (FREDERICKSON TYPE IV)

**PATENT AND EXCLUSIVITY TERMS**

ADB 13 of 117

**EXCLUSIVITY INDICATION**

- I-219 USE OF LIPITOR BY PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDERICKSON TYPE III) WHO DO NOT RESPOND ADEQUATELY TO DIET
- I-220 TREATMENT OF EPISODIC- HEARTBURN, ACID INDIGESTION AND SOUR STOMACH
- I-221 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA (BPH) IN MEN WITH AN ENLARGED PROSTATE TO IMPROVE SYMPTOMS, REDUCE THE RISK OF ACUTE URINARY RETENTION AND REDUCE THE RISK OF THE NEED OF SURGERY
- I-222 PREVENTION OF ISCHEMIC COMPLICATIONS OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION, WHEN CONCURRENTLY ADMINISTERED WITH ASPIRIN
- I-223 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH ALLERGIC AND NONALLERGIC-PERENNIAL RHINITIS IN CHILDREN AGE 6-11 YEARS
- I-224 FOR THE USE IN PEDIATRIC PATIENTS 4 TO 11 YEARS OF AGE FOR THE MANAGEMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- I-225 USE IN PATIENTS WITH PREVIOUS MI AND NORMAL CHOLESTEROL LEVELS, TO REDUCE RISK OF RECURRENT MI, MYOCARDIAL REVASCULARIZATION, AND CEREBROVASCULAR DISEASE EVENTS
- I-226 FIRST-LINE THERAPY FOR THE TREATMENT OF ADVANCED CARCINOMA OF THE OVARY IN COMBINATION WITH CISPLATIN
- I-227 SHORT-TERM TREATMENT OF SYMPTOMATIC GASTROESOPHAGEAL REFLUX DISEASE (GERD)
- I-228 PREVENTION OF MEAL INDUCED HEARTBURN AT A DOSE OF 75MG TAKEN 30-60MIN PRIOR TO A MEAL
- I-229 PRILOSEC (OMEPRAZOLE), AMOXICILLIN, AND CLARITHROMYCIN FOR THE ERADICATION OF H. PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- I-230 IN COMBINATION WITH CIS-PLATIN, FOR THE FIRST LINE TREATMENT OF NON-SMALL CELL LUNG CANCER IN PATIENTS WHO ARE NOT CANDIDATES FOR POTENTIALLY CURATIVE SURGERY AND/OR RADIATION
- I-231 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
- I-232 TREATMENT OF RECURRENT MUCOCUTANEOUS HERPES SIMPLEX INFECTIONS IN HIV-AFFECTED PATIENTS AT A DOSE OF 500MG TWICE DAILY
- I-233 PROPHYLACTIC USE TO REDUCE PERIOPERATIVE BLOOD LOSS AND THE NEED FOR BLOOD TRANSUFSION IN PATIENTS UNDERGOING CARDIOPULMONARY BYPASS IN THE COURSE OF CORONARY ARTERY BYPASS GRAFT SURGERY
- I-234 FOR USE IN COMBINATION WITH CISPLATIN FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED (STAGE IIIA OR IIIB) OR METASTATIC (STAGE IV) NON-SMALL CELL LUNG CANCER
- I-235 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 12 YEARS OF AGE AND OLDER
- I-236 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-237 MAINTENANCE TREATMENT OF ASTHMA AND PREVENTION OF BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-238 ADJUNCTIVE TREATMENT OF LENNOX-GASTAUT SYNDROME IN PEDIATRIC AND ADULT PATIENTS
- I-239 TREATMENT OF PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- I-240 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM AND RESULTANT METABOLIC BONE DISEASE IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL FAILURE (CCR 15 TO 55ML/MIN) NOT YET ON DIALYSIS
- I-241 USE IN PHOTODYNAMIC THERAPY (PDT) FOR REDUCTION OF OBSTRUCTION AND PALLIATION OF SYMPTOMS IN PATIENTS WITH COMPLETELY OR PARTIALLY OBSTRUCTING ENDOBRONCHIAL NONSMALL CELL LUNG CANCER
- I-242 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE AND IN THE TREATMENT OF VULVAR AND VAGINAL ATROPHY IN WOMEN WITH AN INTACT UTERUS
- I-243 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH THE COMMON COLD IN CHILDREN AGE 5 TO 11 YEARS
- I-244 REDUCE THE INCIDENCE OF BREAST CANCER IN WOMEN AT HIGH RISK FOR BREAST CANCER
- I-245 TREATMENT OF ACUTE SINUSITIS
- I-246 TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS

**PATENT AND EXCLUSIVITY TERMS**

ADB 14 of 117

**EXCLUSIVITY INDICATION**

- I-247 USE IN CONVERSION TO MONOTHERAPY IN ADULTS WITH PARTIAL SEIZURES WHO ARE RECEIVING TREATMENT WITH A SINGLE ENZYME-INDUCING ANTIEPILEPTIC DRUG
- I-248 INPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITH/WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIM AND OUTPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM
- I-249 TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS WITH COMPENSATED LIVER DISEASE PREVIOUSLY UNTREATED WITH ALPHA INTERFERON THERAPY
- I-250 PRIMARY PREVENTION OF CORONARY HEART DISEASE IN PATIENTS WITHOUT SYMPATOMATIC CARDIOVASCULAR DISEASE WHO HAVE AVERAGE TO MODERATELY ELEVATED TOTAL-C AND LDL-C AND BELOW AVERAGE HDL-C
- I-251 TREATMENT OF GENERALIZED ANXIETY DISORDER
- I-252 NEW COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS METFORMIN
- I-253 COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS INSULIN
- I-254 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS (LOSS OF BONE MASS)
- I-255 PREVENTION OF PNEUMOCYSTIS CARINII PNEUMONIA (PCP)
- I-256 USE IN TREATMENT OF SMALL CELL LUNG CANCER SENSITIVE DISEASE AFTER FAILURE OF FIRST-LINE CHEMOTHERAPY
- I-257 TREATMENT OF CHRONIC HEPATITIS B ASSOCIATED WITH EVIDENCE OF HEPATITIS B VIRAL REPLICATION AND ACTIVE LIVER INFLAMATION
- I-258 FOR PERENNIAL NONALLERGIC RHINITIS FOR AGES 4 AND ABOVE
- I-259 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
- I-260 EXPANDED PEDIATRIC USE IN CHILDREN YOUNGER THAN ONE MONTH OF AGE TO BIRTH (WITH A GESTATIONAL AGE OF 37 WEEKS OR GREATER)
- I-261 TREATMENT OF SOCIAL ANXIETY DISORDER
- I-262 TREATMENT OR PREVENTION OF BRONCHOSPASM WITH REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE AND FOR THE PREVENTION OF EXERCISE INDUCED BRONCHOSPASM IN CHILDREN AGES 4-12
- I-263 TREATMENT OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION FOR THE PREVENTION OF ISCHEMIC COMPLICATIONS IN PATIENTS ON CONCURRENT ASPIRIN THERAPY
- I-264 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIATION, INCLUDING TOTAL BODY IRRADIATION (TBI) AND FRACTIONATED ABDOMINAL RADIATION
- I-265 TREATMENT OF ATOPIC DERMATITIS IN PEDIATRIC PATIENTS 6 YEARS AND OLDER
- I-266 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN PEDIATRIC PATIENTS AGES 2-16 YEARS WITH PARTIAL ONSET SEIZURES
- I-267 USE IN PEDIATRIC PATIENTS 3 MONTHS OLD AND OLDER - FOR CORTICOSTEROID-RESPONSIVE DERMATOSES
- I-268 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 7-11 YEARS OF AGE
- I-269 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HIGHLY EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING CISPLATIN
- I-270 ADJUVANT TREATMENT OF NODE-POSITIVE BREAST CANCER ADMINISTRERED SEQUENTIALLY TO STANDARD DOXORUBICIN-CONTAINING COMBINATION CHEMOTHERAPY
- I-271 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-272 TREATMENT OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN MEN AND WOMEN RECEIVING GLUCOCORTICOIDS IN A DAILY DOSE EQUIVALENT TO 7.5MG OR GREATER OF PREDNISONE AND WHO HAVE LOW BONE MINERAL DENSITY
- I-273 ADJUNCT TO DIET TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NON FAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-274 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
- I-275 USE IN COMBINATION WITH METFORMIN AND SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES
- I-276 USE OF REZULIN IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH

**PATENT AND EXCLUSIVITY TERMS**

ADB 15 of 117

**EXCLUSIVITY INDICATION**

## TYPE 2 DIABETES

- I-277 TREATMENT OF TYPE III HYPERLIPOPROTEINEMIA
- I-278 TREATMENT OF PATIENTS WITH ISOLATED HYPERTRIGLYCERIDEMIA (FREDERICKSON TYPE IV)
- I-279 TREATMENT OF POST-TRAUMATIC STRESS DISORDER
- I-280 USE OF CARNITOR INJECTION FOR THE PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS
- I-281 INCREASING HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NONFAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-282 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- I-283 TO REDUCE THE INCIDENCE OF MODERATE TO SEVERE XEROSTOMIA IN PATIENTS UNDERGOING POST-OPERATIVE RADIATION TREATMENT FOR HEAD AND NECK CANCER, WHERE THE RADIATION PORT INCLUDES A SUBSTANTIAL PORTION OF THE PAROTID GLANDS
- I-284 TO REDUCE THE NUMBER OF ADENOMATOUS COLORECTAL POLYPS IN FAMILIAL ADENOMATOUS POLYPOSIS PATIENTS AS AN ADJUNCT TO USUAL CARE
- I-285 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN ADULTS AND CHILDREN 3 YEARS OF AGE AND OLDER
- I-286 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE III
- I-287 USE OF PRAVASTATIN IN PATIENTS WITH EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF TOTAL MORTALITY BY REDUCING CORONARY DEATH
- I-288 CHANGES IN SEVERAL SECTIONS OF THE INSERT TO INCORPORATE STATEMENTS CONCERNING THE USE OF HIGH DOSES OF LISINOPRIL TO REDUCE THE RISK OF THE COMBINED OUTCOMES OF MORTALITY AND HOSPITALIZATION IN PATIENTS WITH CONGESTIVE HEART FAILURE
- I-289 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN DIET AND EXERCISE WITH EITHER SINGLE AGENT DOES NOT ACHIEVE ADEQUATE GLYCEMIC CONTROL
- I-290 PREVENTION OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-291 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-292 TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-293 TREATMENT OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-294 TREATMENT OF UNCOMPLICATED ACUTE ILLNESS DUE TO INFLUENZA A AND B IN PEDIATRIC PATIENTS 7 YEARS AND OLDER WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 2 DAYS
- I-295 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR WOMEN WITH AN INTACT UTERUS
- I-296 LONG-TERM INTRAVENOUS TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH THE SCLERODERMA SPECTRUM OF DISEASE IN NYHA CLASS III AND CLASS IV PATIENTS WHO DO NOT RESPOND TO CONVENTIONAL THERAPY
- I-297 SHORT-TERM TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-298 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IIA AND IIB HYPERLIPOPROTEINEMIA
- I-299 USE OF CAMPTOSAR AS A COMPONENT OF FIRST-LINE THERAPY IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR PATIENTS WITH METASTATIC CARCINOMA OF THE COLON OR RECTUM
- I-300 PROPHYLAXIS FOR ASTHMA IN CHILDREN 2-5 YEARS OF AGE
- I-301 TREATMENT OF SIGNS AND SYMPTOMS OF ALLERGIC CONJUNCTIVITIS
- I-302 TREATMENT OF PEDIATRIC PATIENTS WITH PRADER-WILLI SYNDROME
- I-303 INCREASING HDL-CHOLESTEROL IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIAS
- I-304 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IV
- I-305 TREATMENT OF LEVOFLOXACIN SUSCEPTIBLE STRAINS OF PENICILLIN-RESISTANT STREPTOCOCCUS PNEUMONIAE IN PATIENTS WITH COMMUNITY ACQUIRED PNEUMONIA
- I-306 INDUCTION OF SPERMATOGENESIS IN MEN WITH PRIMARY AND SECONDARY HYPOGONADOTROPIC HYPOGONADISM IN WHOM THE CAUSE OF INFERTILITY IS NOT DUE TO PRIMARY TESTICULAR FAILURE
- I-307 NEW COMBINATION USE OF METFORMIN AND INSULIN IN TYPE 2 DIABETES

**PATENT AND EXCLUSIVITY TERMS**

ADB 16 of 117

**EXCLUSIVITY INDICATION**

- I-308 TREATMENT OF PEDIATRIC PATIENTS WITH POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS WHO RESPONDED INADEQUATELY TO SALICYLATES OR OTHER NSAIDS
- I-309 USE OF ACTONEL 35MG ONCE A WEEK TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- I-310 REDUCTION IN RISK OF MYOCARDIAL INFARCTION, STROKE, AND DEATH FROM CARDIOVASCULAR CAUSES
- I-311 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS AGE 3 TO 12 YEARS
- I-312 FIRST LINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER
- I-313 EXTENSION OF INDICATION TO PROVIDE FOR MAINTENANCE OF RESPONSE
- I-314 TOPICAL ANESTHETIC FOR SUPERFICIAL MINOR SURGERY OF GENITAL MUCOUS MEMBRANES AND AS AN ADJUNCT FOR LOCAL INFILTRATION ANESTHESIA IN GENITAL MUCOUS MEMBRANES
- I-315 THROMBOPROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
- I-316 TREATMENT OF NSAID-ASSOCIATED GASTRIC ULCER PATIENTS WHO CONTINUE NSAID USE AND REDUCING RISK OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS WITH HISTORY OF DOCUMENTED GASTRIC ULCER WHO REQUIRE USE OF AN NSAID
- I-317 PROPHYLAXIS OF INFLUENZA IN ADULTS AND ADOLESCENTS 13 YEARS AND OLDER
- I-318 FIRSTLINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER
- I-319 USE FOR SUSPECTED OR CONFIRMED METHANOL POISONING, EITHER ALONE OR IN COMBINATION WITH HEMODIALYSIS
- I-320 TREATMENT OF TYPE 2 DIABETES IN PEDIATRIC PATIENTS (AGES 10-16 YEARS)
- I-321 JUVENILE RHEUMATOID ARTHRITIS
- I-322 USE OF DIPRIVAN IN PATIENTS 3 MONTHS TO 16 YEARS
- I-323 COLORECTAL CANCER
- I-324 REDUCING NEUROLOGIC DISABILITY AND/OR FREQUENCY OF CLINICAL RELAPSES IN PATIENTS WITH SECONDARY (CHRONIC) PROGRESSIVE, PROGRESSIVE RELAPSING, OR WORSENING RELAPSING-REMITTING MULTIPLE SCLEROSIS
- I-325 PREVENTION OF RELAPSE AND RECURRENCE OF DEPRESSION
- I-326 GENERALIZED ANXIETY DISORDER
- I-327 SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN PATIENTS 5 YEARS AND OLDER
- I-328 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 5-6 YEARS OF AGE
- I-329 UNCOMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS
- I-330 MAINTENANCE OF HEALING OF EROSIVE ESOPHAGITIS AND CONTROL OF DAYTIME AND NIGHTTIME HEARTBURN SYMPTOMS IN PATIENTS WITH GERD
- I-331 TREATMENT OF MODERATE ACNE VULGARIS
- I-332 EMPIRIC THERAPY IN FEBRILE NEUTROPENIC PATIENTS WITH SUSPECTED FUNGAL INFECTIONS (EFTN)
- I-333 TOPICAL TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR DUE TO MALASSEZIA FURFUR (FORMERLY PITYROSPORUM ORBICULARE)
- I-334 LONG-TERM TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE WHO FAIL TO MANIFEST CATCH-UP GROWTH BY TWO YEARS OF AGE
- I-335 ADJUNCTIVE THERAPY IN PATIENTS TWO YEARS AND OLDER WITH SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
- I-336 EXPANSION OF INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH PREDOMINATELY CLASSIC SUBFOVEAL CHOROIDAL NEOVASCULARIZATION DUE TO PATHOLOGIC MYOPIA OR PRESUMED OCULAR HISTOPLASMOSIS
- I-337 PATHOLOGICAL HYPERSECRETION ASSOCIATED WITH ZOLLINGER-ELLISON SNYDROME
- I-338 MANAGEMENT OF ACUTE PAIN IN ADULTS AND TREATMENT OF PRIMARY DYSMENORRHEA
- I-339 TREATMENT OF HEPATITIS B IN PEDIATRIC PATIENTS AGES 2-17 YEARS
- I-340 ATOPIC DERMATITIS IN PEDIATRIC PATIENTS AGES 2-5

**PATENT AND EXCLUSIVITY TERMS**

ADB 17 of 117

**EXCLUSIVITY INDICATION**

- I-341 BREAST CANCER COMBINATION THERAPY
- I-342 USE OF FORADIL FOR LONG-TERM, TWICE DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHO-CONSTRICKTION IN PATIENTS WITH COPD INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-343 USE OF COREG FOR SEVERE HEART FAILURE
- I-344 ACNE VULGARIS
- I-345 TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- I-346 TREATMENT OF SYMPTOMATIC GASTRO ESOPHAGEAL REFLUX DISEASE (GERD)
- I-347 TREATMENT OR PREVENTION OF BRONCHOSPASM IN CHILDREN 6 YEARS OF AGE AND OLDER WITH OBSTRUCTIVE AIRWAY DISEASE
- I-348 LONG-TERM, TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD (INCLUDING EMPHYSEMA AND CHRONIC BRONCHITIS)
- I-349 ACUTE CORONARY SYNDROME
- I-350 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND GIRLS AT LEAST ONE YEAR POSTMENARCHAL, AGES 10 TO 17 YEARS, WITH A RECOMMENDED DOSING RANGE OF 10 TO 40MG ONCE DAILY
- I-351 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR ALL STRENGTHS
- I-352 ANTICOAGULANT IN PATIENTS WITH OR AT RISK FOR HEPARIN-INDUCED THROMBOCYTOPENIA UNDERGOING PERCUTANEOUS CORONARY INTERVENTIONS (PCI)
- I-353 TREATMENT OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS
- I-354 MANAGEMENT OF POST HERPETIC NEURALGIA
- I-355 PREMENSTRUAL DYSPHORIC DISORDER
- I-356 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS, INCLUDING ZOLLINGER-ELLISON SYNDROME
- I-357 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS
- I-358 TREATMENT OF PANIC DISORDER
- I-359 TREATMENT OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE
- I-360 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN CHILDREN AGES TWO UP TO AGE THREE
- I-361 TREATMENT OF MULTIPLE MYELOMA AND DOCUMENTED BONE METASTASES FROM SOLID TUMORS, IN CONJUNCTION WITH STANDARD ANTINEOPLASTIC THERAPY. PROSTATE CANCER SHOULD HAVE PROGRESSED AFTER TREATMENT WITH AT LEAST ONE HORMONAL THERAPY
- I-362 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA
- I-363 ADJUVANT TREATMENT OF POST MENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER
- I-364 TREATMENT OF COMMUNITY-ACQUIRED PNEUMONIA IN ADULTS
- I-365 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV) IN PATIENTS WHO ARE INTOLERANT TO AN ACE INHIBITOR
- I-366 PREVENTION OF RELAPSE FOLLOWING LONG-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER
- I-367 COMBINATION THERAPY WITH THIAZOLIDINEDIONE TO LOWER BLOOD GLUCOSE IN PTS WHOSE HYPERGLYCEMIA CANNOT BE CONTROLLED BY DIET/EXERCISE PLUS MONOTHERAPY WITH ANY OF THE FOLLOWING AGENTS: METFORMIN, SULFONYLUREAS, REPAGLINIDE, OR THIAZOLIDINEDIONES
- I-368 USE OF GLUCOVANCE WITH A THIAZOLIDINEDIONE WHEN GLYCEMIC CONTROL IS NOT OBTAINED WITH GLUCOVANCE ALONE
- I-369 PREVENTION AND TREATMENT OF POSTOPERATIVE NAUSEA AND VOMITING
- I-370 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN CHILDREN, AGES 8-13 YEARS, WITH RECOMMENDED DOSE OF 20MG ONCE DAILY AND IN ADOLESCENTS, AGES 14-18 WITH A RECOMMENDED DOSE OF 40MG ONCE DAILY
- I-371 HELICOBACTER PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- I-372 NOSOCOMIAL PNEUMONIA
- I-373 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY

**PATENT AND EXCLUSIVITY TERMS**

ADB 18 of 117

**EXCLUSIVITY INDICATION**

- I-374 SHORT TERM TOPICAL TREATMENT OF MILD TO MODERATE PLAQUE-TYPE PSORIASIS OF NON SCALP REGIONS
- I-375 FIRST LINE THERAPY FOR THE REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- I-376 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (CML)
- I-377 USE OF BRAVELLE FOR MULTIPLE FOLLICULAR DEVELOPMENT (CONTROLLED OVARIAN STIMULATION) DURING ASSISTED REPRODUCTIVE TECHNOLOGY CYCLES IN PATIENTS WHO HAVE PREVIOUSLY RECEIVED PITUITARY SUPPRESSION
- I-378 RELIEF OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-379 USE TAXOTERE IN COMBINATION WITH CISPLATIN FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHO HAVE NOT PREVIOUSLY RECEIVED CHEMOTHERAPY FOR THIS CONDITION
- I-380 TO TREAT PATIENTS WITH SCHIZOPHRENIA OR SCHIZOAFFECTIVE DISORDER AT RISK FOR EMERGENT SUICIDAL BEHAVIOR
- I-381 TREATMENT OF COLD SORES (HERPES LABIALIS) IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER
- I-382 FOR NEWLY-DIAGNOSED HIGH GRADE MALIGNANT GLIOMA PATIENTS AS AN ADJUNCT TO SURGERY AND RADIATION
- I-383 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-384 USE IN COMBINATION WITH INSULIN FOR THE TREATMENT OF PATIENTS WITH TYPE 2 DIABETES MELLITUS
- I-385 MODIFICATION OF THE INDICATION FOR COMMUNITY ACQUIRED PNEUMONIA TO ADD "INCLUDING PENICILLIN-RESISTANT STRAINS, MIC PENICILLIN $\geq$ 2MCG/ML TO STREPTOCOCCUS PNEUMONIAE"
- I-386 RAPAMUNE (SIROLIMUS) WITHIN AN IMMUNOSUPPRESSIVE REGIMEN THAT WOULD ALLOW FOR THE WITHDRAWAL OF CYCLOSPORINE 2 TO 4 MONTHS AFTER RENAL TRANSPLANTATION IN PATIENTS CONSIDERED AT LOW TO MODERATE IMMUNOLOGIC RISK FOR RENAL TRANSPLANT REJECTION
- I-387 ADJUNCTIVE THERAPY OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE
- I-388 TREATMENT OF PATIENTS WITH LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-389 SUPPRESSION OF RECURRENT GENITAL HERPES IN HIV-INFECTED INDIVIDUALS
- I-390 USE IN PTS AT HIGH RISK CORONARY EVENTS DUE TO EXISTING CORONARY HEART DISEASE, DIABETES, PERIPHERAL VESSEL DISEASE, STROKE HISTORY, OTHER CV DISEASE TO REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH, REDUCE NONFATAL MI & STROKE.....
- I-391 ABLATION OF HIGH-GRADE DYSPLASIA IN BARRETT'S ESOPHAGUS PATIENTS WHO DO NOT UNDERGO ESOPHAGECTOMY
- I-392 TX OF PED PATIENTS W/PH+ CHRONIC PHASE CML DISEASE RECURRANCE AFTER STEM CELL TRANSPLANT OR RESISTANCE TO INTERFERON ALPHA THERAPY. NO CONTROLLED TRIALS DEMONSTRATING A CLINICAL BENEFIT SUCH AS IMPROVEMENT IN DISEASE RELATED SX OR INCREASED SURVIVAL
- I-393 CHRONIC BACTERIAL PROSTATITIS
- I-394 USE IN PATIENTS WITH CORONARY HEART DISEASE TO REDUCE THE RISK OF UNDERGOING CORONARY REVASCULARIZATION PROCEDURES
- I-395 TO IMPROVE PHYSICAL FUNCTION
- I-396 EXPANDED INDICATION TO INCLUDE THE ASSESSMENT OF VENTRICULAR FUNCTION IN SUBJECTS BEING EVALUATED FOR HEART DISEASE AND/OR VENTRICULAR FUNCTION
- I-397 EXTENDED PROPHYLAXIS IN PATIENTS UNDERGOING HIP FRACTURE SURGERY
- I-398 IDIOPATHIC SHORT STATURE
- I-399 TREATMENT OF CANDIDEMIA AND THE FOLLOWING CANDIDA INFECTIONS: INTRA-ABDOMINAL ABSCESSSES, PERITONITIS AND PLEURAL SPACE INFECTIONS
- I-400 USE OF OLANZAPINE IN COMBINATION WITH LITHIUM OR VALPROATE FOR THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR DISORDER

**PATENT AND EXCLUSIVITY TERMS**

ADB 19 of 117

**EXCLUSIVITY INDICATION**

- I-401 LONGER-TERM EFFICACY OF ARIPIPRAZOLE IN THE TREATMENT OF SCHIZOPHRENIA
- I-402 DIABETIC FOOT INFECTIONS WITHOUT CONCOMITANT OSTEOMYELITIS
- I-403 USE OF VALTREX IN COMBINATION WITH SAFER SEX PRACTICES FOR THE REDUCTION OF THE RISK OF TRANSMISSION OF GENITAL HERPES DURING SUPPRESSIVE THERAPY OF THE SOURCE PARTNER IN A HETEROSEXUAL COUPLE
- I-404 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES (DEPRESSION, MANIA, HYPMANIA, MIXED EPISODES) IN PATIENTS TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- I-405 TREATMENT OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) USING AN INTERMITTENT DOSING REGIMEN
- I-406 PREVENTION OF CYTOMEGALOVIRUS DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
- I-407 IMPROVE SURVIVAL OF STABLE PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (EJECTION FRACTION<=40%) AND CLINICAL EVIDENCE OF CONGESTIVE HEART FAILURE AFTER AN ACUTE MYOCARDIAL INFARCTION
- I-408 STIMULATION OF PANCREATIC SECRETIONS TO FACILITATE THE IDENTIFICATION OF THE AMPULLA OF VATER AND ACCESSORY PAPILLA DURING ENDOSCOPIC RETROGRADE CHOLANGIO-PANCREATOGRAPHY (ERCP)
- I-409 ESOPHAGEAL CANDIDIASIS
- I-410 USE OF ADVAIR DISKUS 250/50 FOR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) ASSOCIATED WITH CHRONIC BRONCHITIS
- I-411 EXPANDED INDICATION FOR USE IN COMBINATION WITH ANTIDIABETIC DRUGS IN THE THIAZOLIDINEDIONE CLASS
- I-412 MONOTHERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-413 ADJUNCTIVE THERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-414 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM (PE) IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
- I-415 SEVERE HYPERTENSION WHEN THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY
- I-416 THE USE OF CIPRO XR FOR COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
- I-417 USE IN THE LONG TERM TREATMENT OF BIPOLAR I DISORDER
- I-418 ADJUNCTIVE THERAPY W/ MOOD STABILIZERS (LITHIUM OR DIVALPROEX) IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDERS
- I-419 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-420 TOPICAL TREATMENT OF CLINICALLY TYPICAL, NONHYPERKERATOTIC, NONHYPERTROPHIC ACTINIC KERATOSES ON THE FACE OR SCALP IN IMMUNOCOMPETENT ADULTS
- I-421 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND PYELONEPHRITIS DUE TO E.COLI FOR PED PATIENTS (1-17) NOT AS FIRST CHOICE
- I-422 INDICATED FOR THE IN-HOSPITAL SHORT-TERM (UP TO 4 HOURS) REDUCTION IN BLOOD PRESSURE IN PEDIATRIC PATIENTS
- I-423 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS
- I-424 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL INSUFFICIENCY NOT YET ON DIALYSIS
- I-425 ELOXATIN IN COMBINATION WITH INFUSIONAL 5-FLUOROURACIL (5-FU) AND LEUCOVORIN (LV) FOR THE TREATMENT OF PATIENTS PREVIOUSLY UNTREATED FOR ADVANCED COLORECTAL CANCER
- I-426 TREATMENT OF ACUTE PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-427 TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-428 FOR USE IN COMBINATION WITH PACLTAXEL FOR THE FIRST-LINE TREATMENT OF PATIENTS

**PATENT AND EXCLUSIVITY TERMS**

ADB 20 of 117

**EXCLUSIVITY INDICATION**

WITH METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR ANTHRACYCLINE CONTAINING ADJUVANT CHEMOTHERAPY UNLESS ANTHRACYCLINES WERE CLINICALLY CONTRAINDICATED

- I-429 FOR USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER
- I-430 FOR USE IN THE RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS
- I-431 NOSOCOMIAL PNEUMONIA AND COMMUNITY-ACQUIRED PNEUMONIA CAUSED BY STREPTOCOCCUS PNEUMONIAE INDICATION EXPANDED TO INCLUDE MULTI-DRUG RESISTANT STRAINS
- I-432 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA CAUSED BY MULTI-DRUG RESISTANT STREPTOCOCCUS PNEUMONIAE
- I-433 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA IN IMMUNOCOMPETENT ADULTS, WITH A MAXIMUM TUMOR DIAMETER OF 2.0CM, LOCATED ON THE TRUNK (EXCLUDING ANOGENITAL SKIN), NECK, OR EXTREMITIES (EXCLUDING HANDS AND FEET)
- I-434 PREVENTION OF CARDIOVASCULAR DISEASE IN ADULT PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE, BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE TO REDUCE RISK OF MI AND RISK FOR REVASCULARIZATION PROCEDURES AND ANGINA
- I-435 CHRONIC IDIOPATHIC CONSTIPATION
- I-436 FOR USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR THE ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- I-437 TREATMENT OF ACUTE MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-438 EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS
- I-439 USED TO TREAT ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-440 FOR THE REPLACEMENT OF ENDOGENOUS GROWTH HORMONE IN ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-441 USE COMBINATION WITH INFUSIONAL 5-FU/LV FOR ADJUVANT TREATMENT STAGE III COLON CANCER PTS WHO HAVE UNDERGONE COMPLETE RESECTION PRIMARY TUMOR-BASED ON IMPROVEMENT IN DISEASE FREE SURVIVAL, NO DEMONSTRATED BENEFIT OVERALL SURVIVAL AFTER 4YRS
- I-442 USED FOR CANDIDEMIA IN NONNEUTROPENIC PATIENTS AND THE FOLLOWING CANDIDA INFECTIONS: DISSEMINATED INFECTIONS IN SKIN & INFECTIONS IN ABDOMEN, KIDNEY, BLADDER WALL, AND WOUNDS
- I-443 TREATMENT OF NASAL POLYPs IN PATIENTS 18 YEARS OF AGE AND OLDER
- I-444 USE OF PROTONIX IV FOR INJECTION AS STAND ALONE THERAPY FOR THE SHORT-TERM TREATMENT OF PATIENTS HAVING GASTROESOPHAGEAL REFLUX (GERD) WITH A HISTORY OF EROSIONAL ESOPHAGITIS
- I-445 TO IMPROVE (COMPARED TO 4.25% DEXTROSE) LONG-DWELL ULTRAFILTRATION AND CLEARANCE OF CREATININE AND UREA NITROGEN IN PATIENTS WITH HIGH AVERAGE OR GREATER TRANSPORT CHARACTERISTICS, AS DEFINED USING THE PERITONEAL EQUILIBRATION TEST (PET)
- I-446 EXTENDED ADJUVANT TREATMENT OF EARLY BREAST CANCER IN POSTMENOPAUSAL WOMEN WHO HAVE RECEIVED 5 YRS ADJUVANT TAMOXIFEN THERAPY-EFFECTIVENESS BASED ON AN ANALYSIS OF DISEASE FREE SURVIVAL IN PATIENTS TREATED FOR A MEDIAN 24 MONTHS
- I-447 USE OF COPEGUS (RIBAVIRIN) FOR TREATMENT OF CHRONIC HEPATITIS C IN ADULT PATIENTS COINFECTED WITH HIV IN COMBINATION WITH PEGASYS (PEGINTERFERON ALFA-2A)
- I-448 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV AND EJECTION FRACTION <=40%) TO REDUCE THE RISK OF DEATH FROM CARDIOVASCULAR CAUSES AND TO REDUCE HOSPITALIZATIONS FOR HEART FAILURE
- I-449 TO IMPROVE WAKEFULNESS IN TWO NEW PATIENT POPULATIONS WITH EXCESSIVE SLEEPINESS: THOSE WITH OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME AND THOSE WITH SHIFT WORK SLEEP DISORDER
- I-450 TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED HIGH GRADE GLIOMAS CONCOMITANTLY WITH RADIOTHERAPY AND THEN AS ADJUVANT TREATMENT
- I-451 MANAGEMENT OF ENDOMETRIOSIS ASSOCIATED PAIN
- I-452 EXPANDED INDICATION TO INCLUDE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- I-453 USE IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN WHEN DIET, EXERCISE AND

**PATENT AND EXCLUSIVITY TERMS**

ADB 21 of 117

**EXCLUSIVITY INDICATION**

BOTH AGENTS DO NOT RESULT IN ADEQUATE GLYCEMIC CONTROL (TRIPLE THERAPY)

- I-454 MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON FOR UP TO 3 MONTHS
- I-455 MODIFIED HEART FAILURE INDICATION TO INCLUDE TREATMENT OF HEART FAILURE IN PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (NYHA CLASS II-IV; EJECTION FRACTION LESS THAN OR EQUAL TO 40%)
- I-456 TO REDUCE CARDIOVASCULAR DEATH AND TO REDUCE HEART FAILURE HOSPITALIZATIONS. INCLUDES ADDITIONAL INFORMATION ON THE ADDED EFFECT ON THESE OUTCOMES WHEN USED WITH AN ACE INHIBITOR
- I-457 TREATMENT OF PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS
- I-458 USE OF BIVALIRUDIN FOR INJECTION WITH PROVISIONAL USE OF GLYCOPROTEIN IIB/IIA INHIBITOR (GPI) AS LISTED IN THE CLINICAL TRIALS REPLACE-2 SECTION FOR USE AS AN ANTICOAGULANT IN PATIENTS UNDERGOING PERCUTANEOUS CORONARY INTERVENTION (PCI)
- I-459 NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE (NDD-CKD) PATIENTS RECEIVING OR NOT RECEIVING AN ERYTHROPOIETIN
- I-460 TREATMENT OF DIARRHEA CAUSED BY CRYPTOSPORIDIUM PARVUM IN NON-HIV INFECTED PATIENTS 12 YEARS OF AGE AND OLDER
- I-461 USE AS A SINGLE AGENT FOR ADJUVANT TREATMENT IN PATIENTS WITH DUKE'S' C COLON CANCER WHO HAVE UNDERGONE COMPLETE RESECTION OF THE PRIMARY TUMOR WHEN TREATMENT WITH FLUOROPYRIMIDINE THERAPY ALONE IS PREFERRED
- I-462 LONG TERM TREATMENT OF IDIOPATHIC SHORT STATURE
- I-463 TREATMENT OF PATIENTS POST MYOCARDIAL INFARCTION
- I-464 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME
- I-465 PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER
- I-466 FOR RELIEF OF THE SIGNS AND SYMPTOMS OF ANKYLOSING SPONDYLITIS
- I-467 USE OF TOPIRAMATE AS INITIAL MONOTHERAPY IN PATIENTS 10 YEARS OF AGE AND OLDER WITH PARTIAL ONSET OR PRIMARY GENERALIZED TONIC CLONIC SEIZURES
- I-468 USE IN PATIENTS WITH STABLE CORONARY ARTERY DISEASE TO REDUCE THE RISK OF CARDIOVASCULAR MORTALITY OR NON-FATAL MYOCARDIAL INFECTON
- I-469 RELIEF OF THE SIGNS AND SYMPTOMS OF PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- I-470 DIABETIC PERIPHERAL NEUROPATHIC PAIN
- I-471 INDICATED TO REDUCE THE RISK OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH TYPE 2 DIABETES AND WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE
- I-472 USE IN PATIENTS WITH ANGIOGRAPHICALLY DOCUMENTED CORONARY ARTERY DISEASE
- I-473 USE IN COMBINATION WITH GEMCITABINE FOR THE FIRST LINE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER
- I-474 TREATMENT OF IRON DEFICIENCY ANEMIA IN PERITONEAL DIALYSIS DEPENDANT CHRONIC KIDNEY DISEASE IN PATIENTS RECEIVING AN ERYTHROPOIETIN
- I-475 PREVENTION OF NAUSEA AND VOMITTING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- I-476 TREATMENT OF DIABETIC FOOT INFECTIONS WITHOUT OSTEOMYELITIS
- I-477 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY METHICILLIN SUSCEPTIBLE STAPHYLOCOCCUS AUREUS, ESCHERICHIA COLI, KLEBSIELLA PNEUMONIAE, OR ENTEROBACTER CLOACAE
- I-478 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN CHILDREN WITH EPILEPSY AGED 2-4 YEARS
- I-479 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS CAUSED BY E.COLI, B. FRAGILIS, S.ANGINOSUS, S.CONSTELLATUS, E. FAECALIS, P. MIRABILIS, C. PERFRINGENS, B. THETAIOTAOMICRON OR PEPTOSTREPTOCOCCUS SPECIES
- I-480 PROPHYLAXIS OF INFLUENZA FOR PATIENTS BETWEEN 1-12 YEARS OF AGE
- I-481 INDICATED FOR THE ADJUVANT TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER

**PATENT AND EXCLUSIVITY TERMS**

ADB 22 of 117

**EXCLUSIVITY INDICATION**

- I-482 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH OR WITHOUT PSYCHOTIC FEATURES
- I-483 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-484 FOR THE RISK REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS
- I-485 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION
- I-486 ANGIOMAX IS INDICATED FOR PATIENTS WITH, OR AT RISK OF, HIT/HITTS UNDERGOING PCI
- I-487 INDICATED FOR THE RELIEF OF THE INFAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YRS OF AGE OR OLDER
- I-488 MAINTENANCE THERAPY IN BIPOLAR I DISORDER
- I-489 FOR USE IN PEDIATRIC PATIENTS WITH TYPE I DIABETES
- I-490 FOR USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE TREATMENT OF PATIENTS WITH ADVANCED GASTRIC ADENOCARCINOMA, INCLUDING ADENOCARCINOMA OF GASTROESOPHAGEAL JUNCTION, WHO HAVE NOT RECEIVED PRIOR CHEMOTHERAPY FOR ADVANCED DISEASE
- I-491 INFLUENZA PROPHYLAXIS
- I-492 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC OR MIXED EPISODES IN BIPOLAR I DISORDER, WITH OR WITHOUT PSYCHOTIC FEATURES
- I-493 ADMINISTERED IN COMBINATION WITH FENOFLIBRATE, AS ADJUNCTIVE THERAPY TO DIET FOR THE REDUCTION OF ELEVATED TOTAL-C, LDL-C, APO B, AND NON-HDL-C IN PATIENTS WITH MIXED HYPERLIPIDEMIA
- I-494 CLINICAL DATA IN SUPPORT OF AVANDAMET AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH DUAL ROSIGLITAZONE AND METFORMIN THERAPY IS APPROPRIATE
- I-495 ADJUVANT TX OF POSTMENOPAUSAL WOMEN WITH ESTROGEN-RECEPTOR POSITIVE EARLY BREAST CANCER WHO HAVE RECEIVED 2 TO 3 YRS OF TAMOXIFEN AND ARE SWITCHED TO AROMASIN FOR COMPLETION OF A TOTAL OF 5 CONSECUTIVE YRS OF ADJUVANT HORMONAL THERAPY
- I-496 LONG TERM TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME IN PATIENTS WHO HAVE OPEN EPIPHYSES
- I-497 PREVENTION OF SEASONAL MAJOR DEPRESSIVE EPISODES IN PATIENTS WITH SEASONAL AFFECTIVE DISORDER
- I-498 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
- I-499 USE OF GEMZAR IN COMBINATION WITH CARBOPLATIN FOR THE TREATMENT OF PATIENTS WITH ADVANCED OVARIAN CANCER THAT HAS RELAPSED AT LEAST 6 MONTHS AFTER COMPLETION OF PLATINUM-BASED THERAPY
- I-500 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN IMMUNOCOMPETANT PATIENTS WITH A SINGLE DOSE OF FAMCICLOVIR 1500 MG.
- I-502 FOR PTS WITH ST-SEGMENT ELEVATION ACUTE MYOCARDIAL INFARCTION, PLAVIX TO REDUCE RATE OF DEATH FROM ANY CAUSE AND THE RATE OF A COMBINED ENDPOINT OF DEATH, REINFARCTION OR STROKE. NOT KNOWN TO PERTAIN TO PTS WHO RECEIVE PRIMARY ANGIOPLASTY
- I-503 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-504 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME
- I-505 TREATMENT OF STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA), INCLUDING THOSE WITH RIGHT SIDED INFECTIVE ENDOCARDITIS, CAUSED BY METHICILLIN-SUSCEPTIBLE AND METHICILLIN-RESISTANT ISOLATES
- I-506 ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 12 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY
- I-507 ADJUNCT TO DIET TO REDUCE TOTAL-C, LDL-C AND APO B LEVELS IN ADOLESCENT BOYS AND GIRLS WHO ARE AT LEAST ONE YEAR POST-MENARCHE, 10-16 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- I-508 PREMENSTRUAL DYSPHONIC DISORDER
- I-509 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER

**PATENT AND EXCLUSIVITY TERMS**

ADB 23 of 117

**EXCLUSIVITY INDICATION**

- I-510 ADULT DERMAFIBROSARCOMA PROTUBERANS (DFSP)
- I-511 ADULT MYELODYSPLASTIC SYNDROME/MYELOPROLIFERATIVE DISEASES (MDS/MDP)
- I-512 ADULT PH+ ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) MONOTHERAPY
- I-513 ADULT AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM)
- I-514 ADULT HYPEREOSINOPHILIC SYNDROME/CHRONIC EOSINOPHILIC LEUKEMIA (HES/CEL)
- I-515 PROPHYLAXIS OF SURGICAL SITE INFECTION FOLLOWING ELECTIVE COLORECTAL SURGERY
- I-516 PRIMARY GENERALIZED TONIC CLONIC SEIZURES IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-517 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEG SYNDROME (RLS)
- I-518 TREATMENT OF SHORT STATURE OR GROWTH FAILURE IN CHILDREN WITH SHOX (SHORT STATURE HOMEobox CONTAINING GENE) DEFICIENCY WHOSE EPIPHYES ARE NOT CLOSED
- I-519 USE OF TAXOTERE (DOCETAXEL) INJECTION CONCENTRATE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)
- I-520 USE OF EXENATIDE IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE USING A THIAZOLIDINEDIONE ALONE OR IN COMBINATION WITH METFORMIN BUT HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL
- I-521 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 YEAR PRIOR THERAPY
- I-522 TREATMENT OF MODERATE ACNE VULGARIS IN WOMEN AT LEAST 14 YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, AND HAVE ACHIEVED MENARCHE, IF THE PATIENT DESIRES AN ORAL CONTRACEPTIVE FOR BIRTH CONTROL.
- I-523 USE IN ADULT PATIENTS WITH CLINICALLY EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF NONFATAL MYOCARDIAL INFARCTION, FATAL AND NONFATAL STROKE, ANGINA, REVASCULARIZATION PROCEDURES AND HOSPITALIZATION FOR CONGESTIVE HEART FAILURE
- I-524 GENERALIZED ANXIETY DISORDER (GAD)
- I-525 USE OF 0.5MG/0.1MG FOR PREVENTION OF POST-MENOPAUSAL OSTEOPOROSIS
- I-526 TREATMENT OF HYponatremia IN HOSPITALIZED PATIENTS
- I-527 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY
- I-528 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH MENOPAUSE
- I-529 TREATMENT OF DEMENTIA OF THE ALZHEIMER'S TYPE IN PATIENTS WITH SEVERE ALZHEIMER'S DISEASE
- I-530 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION IN PATIENTS 15 YEARS OF AGE AND OLDER
- I-531 MAINTENANCE TREATMENT OF SCHIZOPHRENIA
- I-532 TREATMENT OF BACTERIAL VAGINOSIS IN NON-PREGNANT FEMALES
- I-533 ACUTE ST-SEGMENT ELEVATION MYOCARDIAL INFARCTION (STEMI)
- I-534 EXTENDED TREATMENT OF SYMPTOMATIC VENOUS THROMBOEMBOLISM (VTE) AND/OR PULMONARY EMBOLISM TO REDUCE THE RECURRENCE OF VTE IN PATIENTS WITH CANCER
- I-535 MANAGEMENT OF FIBROMYALGIA
- I-536 FOR THE TREATMENT OF SHORT STATURE IN CHILDREN WITH NOONAN SYNDROME
- I-537 LONG TERM TREATMENT OF PANIC DISORDER
- I-538 SHORT TERM TREATMENT OF PANIC DISORDER
- I-539 REDUCTION IN RISK OF INVASIVE BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS OR AT HIGH RISK FOR INVASIVE BREAST CANCER
- I-540 TREATMENT OF SCHIZOPHRENIA IN ADOLESCENTS AGES 13-17
- I-541 TREATMENT OF BIPO极 I DISORDER IN CHILDREN AGES 10-12 AND ADOLESCENTS AGES 13-17
- I-542 EXPANSION OF PATIENT POPULATION FOR HEAD AND NECK CANCER FROM "INOPERABLE" PATIENTS TO ALL PATIENTS

**PATENT AND EXCLUSIVITY TERMS**

ADB 24 of 117

**EXCLUSIVITY INDICATION**

- I-543 USE IN COMBINATION WITH CISPLATIN AND FLUOROURACIL FOR THE INDUCTION TREATMENT OF PATIENTS WITH LOCALLY ADVANCED SQUAMOUS CELL CARCINOMA OF THE HEAD AND NECK (SCCHN)
- I-544 ADJUNCTIVE THERAPY OF MYOCLONIC SEIZURES IN ADULTS AND ADOLESCENTS AGE 16 AND OVER WITH JUVENILE MYOCLONIC EPILEPSY
- I-545 ADJUNCTIVE TREATMENT TO TREAT PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- I-546 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA
- I-547 ADJUNCTIVE THERAPY TO DIET TO SLOW THE PROGRESSION OF ARTEROSCLEROSIS IN ADULT PATIENTS AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL-C AND LDL-C TO TARGET LEVELS
- I-548 SEASONAL ALLERGIC RHINITIS IN PATIENTS 6 THROUGH LESS THAN 12 YEARS OF AGE
- I-549 USE OF AVALIDE TABLETS AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS
- I-550 TREATMENT OF HYPERTENSION IN PEDIATRIC PATIENTS 6-16 YEARS OF AGE
- I-551 TREATMENT OF SHORT STATURE IN CHILDREN WITH TURNER'S SYNDROME
- I-552 ADJUNCTIVE TREATMENT FOR RADIOIODINE ABLATION OF THYROID TISSUE REMNANTS IN PATIENTS WHO HAVE UNDERGONE THYROIDECTOMY FOR WELL-DIFFERENTIATED THYROID CANCER AND WHO DO NOT HAVE EVIDENCE OF METASTATIC THYROID CANCER
- I-553 FOR USE AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- I-554 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESSSES
- I-555 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER IN PEDIATRIC PATIENTS AGED 10-17 YEARS
- I-556 PREVENTION OF POST OPERATIVE NAUSEA AND VOMITING FOR UP TO 24 HOURS FOLLOWING SURGERY
- I-557 USE OF AMITIZA (LUBIPROSTONE) 8 MCG TWICE DAILY FOR TREATMENT OF IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN WOMEN GREATER THAN OR EQUAL TO 18 YEARS OLD
- I-558 MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION AND REDUCING EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-559 ADJUNCTIVE THERAPY ADDED TO LITHIUM OR VALPROATE IN SHORT TERM TREATMENT OF BIPOLAR DISORDER, MANIC OR MIXED
- I-560 MAINTENANCE TREATMENT FOR BIPOLAR I DISORDER, AS ADJUNCTIVE THERAPY TO LITHIUM OR DIVALPROEX
- I-561 LONG-TERM TREATMENT OF SOCIAL ANXIETY DISORDER
- I-562 MAINTENANCE TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD) IN CHILDREN AND ADOLESCENTS
- I-563 ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SIEZURES IN ADULTS AND CHILDREN 16 YEARS OF AGE AND OLDER WITH IDIOPATHIC GENERALIZED EPILEPSY
- I-564 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA
- I-565 USE OF DUTASTERIDE IN COMBINATION WITH TAMSULOSIN FOR THE TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-566 MANAGEMENT OF FIBROMYALGIA
- I-567 INITIAL THERAPY IN PATIENTS LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS
- I-568 USE OF APTIVUS, CO-ADMINISTERED W/RITONAVIR, FOR COMBINATION ANTIRETROVIRAL TREATMENT OF HIV-1 INFECTED PED (AGE 2-18 YRS) PATIENTS WHO ARE TREATMENT-EXPERIENCED AND INFECTED W/HIV-1 STRAINS RESISTANT TO MORE THAN ONE PROTEASE INHIBITOR
- I-569 TREATMENT OF CHRONIC HEPATITIS B
- I-570 TREATMENT OF CHICKEN POX IN IMMUNOCOMPETENT PEDIATRIC PATIENTS 2 TO <18 YEARS OF AGE
- I-571 NON-SMALL CELL LUNG CANCER IN COMBINATION WITH CISPLATIN AND AS SINGLE AGENT FOR NONSQAMOUS NON-SMALL CELL LUNG CANCER

**PATENT AND EXCLUSIVITY TERMS**

ADB 25 of 117

**EXCLUSIVITY INDICATION**

- I-572 TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE (SGA) WITH NO CATCH-UP BY AGE 2-4 YRS.
- I-573 TO TREAT PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDRICKSON TYPE III HYPERLIPOPROTEINEMIA) AS AN ADJUNCT TO DIET
- I-574 MONOTHERAPY IN THE TREATMENT OF BIPOLAR DEPRESSION
- I-575 MONOTHERAPY IN THE TREATMENT OF BIPOLAR MANIA
- I-576 ADJUNCTIVE THERAPY IN THE TREATMENT OF BIPOLAR MANIA
- I-577 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES
- I-578 EXPANSION OF INDICATION TO INCLUDE TREATMENT OF HIV IN TREATMENT NAIVE ADULTS
- I-579 TREATMENT OF MODERATE TO SEVERE DYSPAREUNIA, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO MENOPAUSE AND NEW TWICE WEEKLY DOSING REGIMEN FOR THIS INDICATION
- I-580 INDOLENT B-CELL NON-HODGKINS LYMPHOMA (NHL) THAT HAS PROGRESSED DURING OR WITHIN SIX MONTHS OF TREATMENT WITH RITUXIMAB OR A RITUXIMAB CONTAINING REGIMEN
- I-581 TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- I-582 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- I-583 ADJUVANT TREATMENT OF ADULT PATIENTS FOLLOWING COMPLETE GROSS RESECTION OF KIT (CD117) POSITIVE GASTOINTESTINAL STROMAL TUMORS (GIST)
- I-584 TREATMENT AND PREVENTION OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN PATIENTS EXPECTED TO BE ON GLUCOCORTICOIDS FOR AT LEAST 12 MONTHS
- I-585 TREATMENT OF SHORT STATURE IN PEDIATRIC PATIENTS SMALL FOR GESTATIONAL AGE WHO DO NOT MANIFEST CATCH UP GROWTH BY AGE 2 TO 4 YEARS
- I-586 COMMUNITY ACQUIRED BACTERIAL PNEUMONIA
- I-587 ADDITIONAL PATHOGENS TO COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS INDICATION
- I-588 ADDITIONAL PATHOGENS TO COMPLICATED INTRA-ABDOMINAL INFECTIONS INDICATION
- I-589 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH OLANZAPINE
- I-590 ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH OLANZAPINE)
- I-591 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD) IN COMBINATION WITH FLUOXETINE
- I-592 ACUTE TREATMENT OF DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR DISORDER (IN COMBINATION WITH FLUOXETINE)
- I-593 TREATMENT OF TREATMENT RESISTANT DEPRESSION (TRD)
- I-594 INDICATION EXPANDED TO INCLUDE PATIENTS WHO HAVE EXPERIENCED A FIRST CLINICAL EPISODE AND HAVE MRI FEATURES CONSISTENT WITH MULTIPLE SCLEROSIS
- I-595 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-596 USE AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
- I-597 MONOTHERAPY FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
- I-598 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION INDICATION EXPANDED TO INCLUDE DELAY IN CLINICAL WORSENING
- I-599 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS
- I-600 FOR USE AS INITIAL THERAPY IN PATIENTS WHO ARE LIKELY TO NEED MULTIPLE DRUGS TO ACHIEVE THEIR BLOOD PRESSURE GOALS
- I-601 MAINTENANCE TREATMENT IN PATIENTS WITH ADVANCED OR METASTATIC NONSQUAMOUS NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST LINE CHEMOTHERAPY
- I-602 TREATMENT OF MEN AND WOMEN WITH OSTEOPOROSIS ASSOCIATED WITH SUSTAINED SYSTEMIC GLUCOCORTICOID THERAPY AT HIGH RISK FOR FRACTURE
- I-603 GOUT FLARES
- I-604 PREVENTION OF CMV DISEASE IN KIDNEY AND HEART TRANSPLANT PATIENTS 4 MONTHS TO 16

**PATENT AND EXCLUSIVITY TERMS**

ADB 26 of 117

**EXCLUSIVITY INDICATION**

## YEARS AT HIGH RISK

- I-605 ADJUNCT TO MOOD STABILIZERS AND/OR ANTIDEPRESSANTS FOR SCHIZOAFFECTIVE DISORDER
- I-606 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY
- I-607 INDICATION EXPANDED TO INCLUDE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (WHO GROUP I) IN PATIENTS WITH CLASS II SYMPTOMS
- I-608 REDUCE LDL-C LEVELS IN BOYS AND POSTMENARHAL GIRLS, 10 TO 17 YEARS OF AGE, WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA AS MONOTHERAPY OR IN COMBINATION WITH A STATIN AFTER FAILING AN ADEQUATE TRIAL OF DIET THERAPY
- I-610 TREATMENT OF HEAVY MENSTRUAL BLEEDING FOR WOMEN WHO CHOOSE TO USE INTRAUTERINE CONTRACEPTION AS THEIR METHOD OF CONTRACEPTION
- I-611 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND POSTMENARHAL GIRLS, AGES 10 TO 17 YEARS, WITH A RECOMMENDATION DOSING RANGE OF 5 TO 20 MG ONCE DAILY
- I-612 MICARDIS 80 MG FOR REDUCTION OF THE RISK OF MYOCARDIAL INFARCTION, STROKE, OR DEATH FROM CARDIOVASCULAR CAUSES IN PATIENTS 55 YEARS OF AGE OR OLDER AT HIGH RISK OF DEVELOPING MAJOR CARDIOVASCULAR EVENTS WHO ARE UNABLE TO TAKE ACE INHIBITORS
- I-613 MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE TO LESS THAN 18 YEARS OF AGE
- I-614 SHORT TERM TREATMENT OF EROSIONAL ESOPHAGITIS ASSOCIATED WITH GERD IN PEDIATRIC PATIENTS AGES FIVE YEARS AND OLDER
- I-615 MAINTENANCE TREATMENT OF BIPOLAR DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-616 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17 YEARS OF AGE
- I-617 MAINTENANCE OF GENERALIZED ANXIETY DISORDER (GAD)
- I-618 ADJUNCTIVE THERAPY IN THE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- I-619 INTRAVENOUS CONTRAST ENHANCED COMPUTER TOMOGRAPHY OF THE HEAD AND BODY
- I-620 FOR USE IN COMBINATION WITH LETROZOLE FOR THE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER THAT OVEREXPRESSES THE HER2 RECEPTOR FOR WHOM HORMONAL THERAPY IS INDICATED
- I-621 PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE, BASED ON THE RESULTS OF JUSTIFICATION FOR THE USE OF STATINS IN PRIMARY PREVENTION; AN INTERVENTION TRIAL EVALUATING ROSUVASTATIN (JUPITER)
- I-622 ADJUNCTIVE THERAPY FOR PRIMARY GENERALIZED TONIC-CLONIC SEIZURES IN PATIENTS THIRTEEN YEARS OF AGE AND OLDER
- I-623 TREATMENT OF SIGNS AND SYMPTOMS OF ADVANCED IDIOPATHIC PARKINSON'S DISEASE
- I-624 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES OF PLATINUM-BASED FIRST-LINE CHEMOTHERAPY
- I-625 PANCREATIC INSUFFICIENCY DUE TO CHRONIC PANCREATITIS AND PANCREATECTOMY
- I-626 RELIEF OF NASAL CONGESTION ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATENTS 2 YEARS OF AGE AND OLDER
- I-627 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH & CML) IN CHRONIC PHASE.
- I-628 MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS
- I-629 ADJUNCTIVE THERAPY WITH EITHER LITHIUM OR VALPROATE FOR THE ACUTE TREATMENT OF MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-630 TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS (TS) WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION.
- I-631 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE FOLLOWING OPIOID DETOXIFICATION
- I-632 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN
- I-633 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER AS AN ADJUNCT TO LITHIUM OR VALPROATE
- I-634 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY

**PATENT AND EXCLUSIVITY TERMS**

ADB 27 of 117

**EXCLUSIVITY INDICATION**

- I-635 ADJUNCTIVE TREATMENT WITH LONG-ACTING ORAL PSYCHOSTIMULANTS FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- I-636 TREATMENT OF EXTERNAL GENITAL AND PERIANAL WARTS/CONDYLOMA ACUMINATA IN PATIENTS 12 YEARS OR OLDER
- I-637 USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER
- I-638 FOR PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC.
- I-639 TREATMENT OF PROGRESSIVE, WELL-DIFFERENTIATED PANCREATIC NEUROENDOCRINE TUMORS IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED, OR METASTATIC DISEASE
- I-640 MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
- I-641 TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-642 TREATMENT OF ERECTILE DYSFUNCTION (ED) AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
- I-643 REDUCE THE RISK OF STROKE AND SYSTEMIC EMBOLISM IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION.
- I-644 MONOTHERAPY IN PATIENTS 13 YEARS OF AGE AND OLDER WITH PARTIAL SEIZURES WHO ARE RECEIVING THERAPY WITH A SINGLE ANTIEPILEPTIC DRUG (AED)
- I-645 MAINTENANCE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS
- I-646 SIGNS AND SYMPTOMS OF ADVANCED PARKINSON'S DISEASE (APD)
- I-647 SIGNS AND SYMPTOMS OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)
- I-648 TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
- I-649 TREATMENT OF PATIENTS WITH ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
- I-650 TREATMENT OF ADULTS WITH RENAL ANGIOMYOLIPOMA AND TUBEROUS SCLEROSIS COMPLEX (TSC), NOT REQUIRING IMMEDIATE SURGERY
- I-651 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH SPINAL CORD INJURY
- I-652 MANAGEMENT OF POSTHERPETIC NEURALGIA
- I-653 TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
- I-654 MAGNETIC RESONANCE ANGIOGRAPHY (MRA) TO EVALUATE ADULTS WITH KNOWN OR SUSPECTED RENAL OR AORTO-ILIO-FEMORAL OCCLUSIVE VASCULAR DISEASE
- I-655 TREATMENT OF POSTMENOPAUSAL WOMEN WITH ADVANCED HORMONE RECEPTOR-POSITIVE, HER2-NEGATIVE BREAST CANCER (ADVANCED HR+BC) IN COMBINATION WITH EXEMESTANE, AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
- I-656 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY (DPN) IN ADULTS WHEN A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME
- I-657 PLAQUE PSORIASIS OF THE SCALP
- I-658 FIRST-LINE TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER, IN COMBINATION WITH CARBOPLATIN, IN PATIENTS WHO ARE NOT CANDIDATES FOR CURATIVE SURGERY OR RADIATION THERAPY
- I-659 PLAQUE PSORIASIS OF THE BODY
- I-660 TREATMENT OF DEEP VEIN THROMBOSIS
- I-661 TREATMENT OF PULMONARY EMBOLISM
- I-662 REDUCTION IN RISK FOR DEEP VEIN THROMBOSIS AND THE REDUCTION IN RISK FOR PULMONARY EMBOLISM
- I-663 IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- I-664 TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY
- I-665 TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS OF AGE AND OLDER WITH (NTDT) SYNDROMES AND WITH A (LIC) OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT (MG FE/G DW) AND SERUM FERRITIN GREATER THAN 300MCG/L

**PATENT AND EXCLUSIVITY TERMS**

ADB 28 of 117

**EXCLUSIVITY INDICATION**

- I-666 TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) IN COMBINATION WITH CHEMOTHERAPY
- I-667 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE
- I-668 PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT
- I-669 SCINTIGRAPHIC ASSESSMENT OF SYMPATHETIC INNERVATION OF THE MYOCARDIUM BY MEASUREMENT OF THE HEART TO MEDIASTINUM (H/M) RATIO OF RADIOACTIVITY UPTAKE IN PATIENTS WITH NYHA CLASS II OR CLASS III HEART FAILURE AND LVEF LESS THAN 35%
- I-670 TREATMENT OF OPIOID-INDUCED CONSTIPATION (OIC) IN ADULTS WITH CHRONIC, NON-CANCER PAIN
- I-671 FIRSTLINE TREATMENT OF PATIENTS WITH METASTATIC NON- SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21(L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-672 USE IN PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- I-673 TREATMENT OF HOSPITAL-ACQUIRED BACTERIAL PNEUMONIA/VENTILATOR-ASSOCIATED BACTERIAL PNEUMONIA (HABP/VABP) CAUSED BY SUSCEPTIBLE ISOLATES OF S. AUREUS (INCLUDING METHICILLIN-SUSCEPTIBLE AND RESISTANT ISOLATES) WHEN ALTERNATIVE TREATMENTS ARE NOT SUITABLE
- I-674 TREATMENT OF PATIENTS WITH DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER (BIPOLAR DEPRESSION) AS MONOTHERAPY AND AS ADJUNCTIVE THERAPY WITH LITHIUM OR VALPROATE
- I-675 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER
- I-676 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS, IN COMBINATION WITH GEMCITABINE
- I-677 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DTC) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- I-678 TRAMETINIB, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- I-679 RISK REDUCTION OF REBLEEDING OF GASTRIC OR DUODENAL ULCERS FOLLOWING THERAPEUTIC ENDOSCOPY IN ADULTS
- I-680 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- I-681 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT) WHICH MAY LEAD TO PULMONARY EMBOLISM (PE), IN ADULT PATIENTS WHO HAVE UNDERGONE HIP OR KNEE REPLACEMENT
- I-682 TREATMENT OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM (PE) IN PATIENTS WHO HAVE BEEN TREATED WITH A PARENTERAL ANTICOAGULANT FOR 5-10 DAYS
- I-683 TO REDUCE THE RISK OF RECURRENCE OF DVT AND PE IN PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED
- I-684 PREVENTION OF ACUTE NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING HIGHLY EMETOGENIC CANCER CHEMOTHERAPY IN PEDIATRIC PATIENTS AGED 1 MONTH TO LESS THAN 17 YEARS
- I-685 EXPANDED INDICATION OF RASAGILINE AS AN ADD-ON THERAPY TO STABLE DOSES OF DOPAMINE AGONISTS IN THE TREATMENT OF EARLY PARKINSON'S DISEASE
- I-686 INDICATED FOR THE TREATMENT OF DIABETIC MACULAR EDEMA IN PATIENTS WHO ARE PSEUDOPHAKIC OR ARE PHAKIC AND SCHEDULED FOR CATARACT SURGERY
- I-687 GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
- I-688 GADAVIST IS INDICATED WITH MRI TO DETECT THE PRESENCE AND EXTENT OF MALIGNANT BREAST DISEASE
- I-689 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION
- I-690 INDICATED FOR THE TREATMENT OF DEEP VEIN THROMBOSIS (DVT)
- I-691 INDICATED TO REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT) AND

**PATENT AND EXCLUSIVITY TERMS**

ADB 29 of 117

**EXCLUSIVITY INDICATION**

PULMONARY EMBOLISM (PE) FOLLOWING INITIAL THERAPY

I-692 INDICATED FOR MANAGEMENT OF OSTEOARTHRITIS PAIN.

I-693 TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC)

I-694 TREATMENT OF PATIENTS WITH MODERATE TO SEVERE PLAQUE PSORIASIS WHO ARE CANDIDATES FOR PHOTOTHERAPY OR SYSTEMIC THERAPY

I-695 REVISED INDICATION FOR BORTEZOMIB IN THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA

I-696 USE AS MONOTHERAPY IN THE TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGE 17 YEARS AND OLDER

I-697 FOR USE IN COMBINATION WITH SOFOSBUVIR FOR THE TREATMENT OF PATIENTS WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1 INFECTION

I-698 SCHIZOAFFECTIVE DISORDER AS MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS

I-699 FOR TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA

I-700 TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S DISORDER (6-18 YEARS)

I-701 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL-OR MODERATELY-DIFFERENTIATED, LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS (GEP-NETS) TO IMPROVE PROGRESSION FREE SURVIVAL

I-702 FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM MACROGLOBULINEMIA

I-703 MODERATE TO SEVERE BINGE EATING DISORDER (BED)

I-704 EXPANDED INDICATION TO INCLUDE PATIENTS WHO ARE VIROLOGICALLY-SUPPRESSED (HIV-1 RNA &lt;50 COPIES/ML) ON A STABLE ANTIRETROVIRAL REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TREATMENT FAILURE IN ORDER TO REPLACE THEIR CURRENT REGIMEN

I-705 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS 6 YEARS AND OLDER WHO HAVE AN R117H MUTATION IN THE CFTR GENE

I-706 EXPANDED INDICATION FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA

I-707 POMALYST, IN COMBINATION WITH DEXAMETHASONE, IS INDICATED FOR PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY

I-708 DAILY TREATMENT OF ASTHMA IN PATIENTS AGED 18 YEARS AND OLDER

I-709 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS

I-710 ADJUNCTIVE THERAPY FOR THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC (PG TC) SEIZURES IN PATIENTS WITH EPILEPSY 12 YEARS OF AGE OR OLDER.

I-711 INCLUSION OF PEDIATRIC PATIENTS AGES 6 YRS AND OLDER FOR THE TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC ITP WHO HAVE HAD AN INSUFFICIENT RESPONSE TO CORTICOSTEROIDS, IMMUNOGLOBULINS, OR SPLENECTOMY.

I-712 EXPANDED INDICATION FOR USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE PRIOR LINES OF THERAPY

I-713 REVISIONS TO THE LABELING TO PERMIT THE USE OF ZUBSOLV AS INITIAL ("INDUCTION") TREATMENT OF OPIOID DEPENDENCE

I-714 EXTENDS THE 2011 APPROVAL OF BRILINTA FOR USE BEGINNING WITH ACS TO USE BEGINNING MORE REMOTE FROM MYOCARDIAL INFARCTION

I-715 FOR THE ADDITION OF THE INDICATION FOR MONOTHERAPY TREATMENT IN PARTIAL-ONSET SEIZURES IN ADULTS.

I-716 REVISED INDICATION TO INCLUDE LANGUAGE ABOUT THE BENEFITS OF USING LETAIRIS IN COMBINATION WITH Tadalafil TO REDUCE THE RISK OF DISEASE PROGRESSION AND HOSPITALIZATION FOR WORSENING PAH AND TO IMPROVE EXERCISE ABILITY, BASED ON THE AMBITION STUDY

I-717 EXPANDED INDICATION TO INCLUDE THE TREATMENT OF CHRONIC HEPATITIS C GENOTYPE 4

I-718 EXPANDED INDICATION TO INCLUDE SUBJECTS INFECTED WITH CHRONIC HEPATITIS C, GENOTYPE 6 VIRUS INFECTION BASED UPON THE RESULTS OF THE ELECTRON- 2 STUDY

I-719 EXPANDED INDICATION TO INCLUDE THE TREATMENT OF SUBJECTS WITH GENOTYPE 5 CHRONIC HEPATITIS C VIRUS INFECTION BASED ON THE RESULTS FROM STUDY GS-US-337-119.

**PATENT AND EXCLUSIVITY TERMS**

ADB 30 of 117

**EXCLUSIVITY INDICATION**

- I-720 EXPANDED INDICATION TO INCLUDE TREATMENT OF GENOTYPE 4, CHRONIC HEPATITIS C VIRUS INFECTION BASED UPON THE RESULTS FROM STUDIES ION-4 AND GS-US-337-119.
- I-721 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN.
- I-722 REVISED INDICATION FOR USE IN COMBINATION WITH DEXAMETHASONE OR WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY.
- I-723 AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE OR MORE LINES OF THERAPY
- I-724 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL DIFFERENTIATED, NON-FUNCTIONAL NEUROENDOCRINE TUMORS (NET) OF GI OR LUNG ORIGIN WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- I-725 TREATMENT OF HORMONE RECEPTOR (HR)-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION THERAPY WITH PALBOCICLIB AND FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY.
- I-726 EXPANSION OF THE PATIENT POPULATION TO INCLUDE PATIENTS WITH RECURRENT CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1 OR 3 AFTER LIVER TRANSPLANTATION
- I-727 EXPANSION OF THE INDICATION TO INCLUDE TREATMENT OF SUBJECTS WITH GENOTYPE-1 CHRONIC HEPATITIS C VIRUS INFECTION, INCLUDING SUBJECTS WHO ARE CO-INFECTED WITH THE HUMAN IMMUNODEFICIENCY VIRUS (HIV-1) BASED ON THE RESULTS FROM THE ALLY-2 CLINICAL TRIAL
- I-728 EXPANDED INDICATION FOR USE IN ULTRASONOGRAPHY OF THE LIVER FOR CHARACTERIZATION OF FOCAL LIVER LESIONS IN ADULT AND PEDIATRIC PATIENTS
- I-729 PROVIDES FOR THE FRONTLINE INDICATION FOR THE TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA
- I-730 NEW INDICATION FOR THE TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- I-731 FOR USE IN MAGNETIC RESONANCE ANGIOGRAPHY IN ADULT AND PEDIATRIC PATIENTS (INCLUDING TERM NEONATES) TO EVALUATE KNOWN OR SUSPECTED SUPRA-AORTIC OR RENAL ARTERY DISEASE
- I-732 TREATMENT OF PEDIATRIC PATIENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA TO REDUCE LDL-C, TOTAL C, NONHDL-C AND APOB AS AN ADJUNCT TO DIET, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS
- I-733 USE OF CANAGLIFLOZIN FOR INITIAL THERAPY IN COMBINATION WITH METFORMIN
- I-734 EXPANDED INDICATION FOR THE USE OF LENVIMA IN COMBINATION WITH EVEROLIMUS FOR THE TREATMENT OF PATIENTS WITH ADVANCED RCC FOLLOWING ONE PRIOR ANTI-ANGIOGENIC THERAPY.
- I-735 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH CANAGLIFLOZIN AND METFORMIN IS APPROPRIATE
- I-736 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL)
- I-737 REVISED INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LEUKEMIA (SLL) WITH 17P DELETION
- I-738 REVISIONS TO THE INDICATIONS AND USAGE SECTION WITH RESPECT TO COMPLICATED INTRA-ABDOMINAL INFECTIONS
- I-739 TO REDUCE THE RISK OF CARDIOVASCULAR DEATH IN ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-740 EXPANDED INDICATION FOR THE TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 6 YEARS AND OLDER TO INCLUDE THE G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N, OR S549R MUTATION IN THE CFTR GENE
- I-741 TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
- I-742 TREATMENT OF NODAL MARGINAL ZONE LYMPHOMA
- I-743 INFORMATION ADDED TO THE LABELING FOR THE ADDITION OF THE TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 4 (GT4) INFECTED PATIENTS WITH COMPENSATED CIRRHOSIS BASED ON RESULTS FROM STUDY M11-665
- I-744 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC) WHO HAVE BEEN PREVIOUSLY TREATED WITH SORAFENIB

**PATENT AND EXCLUSIVITY TERMS**

ADB 31 of 117

**EXCLUSIVITY INDICATION**

- I-745 MEKINIST, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.
- I-746 NEW INDICATION OF MAINTENANCE MONOTHERAPY TREATMENT OF BIPOLAR I DISORDER IN ADULTS
- I-747 FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED WITH HIGH-DOSE BUSULFAN AND CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENEIC HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION FOR PEDIATRIC PATIENTS WITH CLASS 3 BETA-THALASSEMIA
- I-748 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS FIVE YEARS OF AGE AND OLDER
- I-749 MONOTHERAPY FOR THE TREATMENT OF HORMONE RECEPTOR (HR) POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE, ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- I-750 REDUCE THE RISK OF MAJOR ADVERSE CARDIOVASCULAR EVENTS IN ADULTS WITH TYPE 2 DIABETES MELLITUS AND ESTABLISHED CARDIOVASCULAR DISEASE
- I-751 TREATMENT OF TARDIVE DYSKINESIA
- I-752 CORONARY COMPUTED TOMOGRAPHY ANGIOGRAPHY (CCTA) TO ASSIST DIAGNOSTIC EVALUATION OF PATIENTS WITH SUSPECTED CORONARY ARTERY DISEASE
- I-753 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD) AFTER FAILURE OF ONE OR MORE LINES OF SYSTEMIC THERAPY
- I-754 TO REDUCE THE FREQUENCY OF SHORT-ACTING SOMATOSTATIN ANALOG RESCUE THERAPY WHEN USED FOR THE TREATMENT OF ADULTS WITH CARCINOID SYNDROME
- I-755 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RENAL CELL CARCINOMA (RCC) FOLLOWING NEPHRECTOMY
- I-756 EXPANDED THE APPROVED INDICATION BY REMOVING THE RESTRICTION FOR USE ONLY IN PATIENTS WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- I-757 TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- I-758 FOR USE WITH RILPIVIRINE AS A COMPLETE REGIMEN TO REPLACE THE CURRENT ARV REGIMEN IN VIROLOGICALLY SUPPRESSED PATIENTS ON A STABLE ARV REGIMEN FOR AT LEAST 6 MONTHS WITH NO HISTORY OF TX FAILURE OR KNOWN SUBSTITUTIONS ASSOC. WITH RESISTANCE TO EITHER ARV
- I-759 TREATMENT OF ADULT PATIENTS WITH NEWLY-DIAGNOSED CHRONIC PHASE (CP) PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOGENOUS LEUKEMIA (Ph+CM)
- I-760 FOR THE TREATMENT OF PATIENTS WITH ADVANCED RENAL CELL CARCINOMA
- I-761 TREATMENT OF ADULT PATIENTS WITH ACTIVE PSORIATIC ARTHRITIS WHO HAVE HAD AN INADEQUATE RESPONSE OR INTOLERANCE TO METHOTREXATE OR OTHER NON-BIOLOGIC DISEASE-MODIFYING ANTIRHEUMATIC DRUGS

**EXCLUSIVITY MISCELLANEOUS**

- M-1 INFORMATION REGARDING SUPERIORITY CLAIM OVER RANITIDINE FOR DAY AND NIGHT HEARTBURN ADDED TO CLINICAL STUDIES SECTION
- M-2 APPROVAL FOR ADDITION TO CLINICAL PHARMACOLOGY SECTION OF THE LABEL REGARDING (1) IMPROVEMENT IN BONE MINERAL DENSITY IN CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENT PATIENTS AND (2) INCREASES IN SERUM ALKALINE PHOSPHATASE
- M-3 ADDITION OF EFFICACY AND SAFETY INFORMATION IN WHICH FOSAMAX WAS USED CONCOMITANTLY WITH ESTROGEN ALONE OR WITH ESTROGEN PLUS PROGESTIN
- M-4 CHANGES TO PEDIATRIC USE SECTION TO PROVIDE INFORMATION REGARDING SAFETY AND EFFICACY IN PEDIATRIC PATIENTS AS YOUNG AS 2 YEARS OLD
- M-5 INFORMATION REGARDING EFFECTS IN PATIENTS WITH ASTHMA ON CONCOMITANT INHALED CORTICOSTEROIDS IN CLINICAL PHARMACOLOGY SECTION
- M-6 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH GLUCOPHAGE/GLYBURIDE COMBINATION ADDED TO CLINICAL PHARMACOLOGY AND DOSING AND ADMINISTRATION
- M-7 CLINICAL PHARMACOLOGY IN PEDIATRIC PATIENTS; DOSAGE AND ADMINISTRATION INFORMATION
- M-8 ADDITIONAL INFORMATION FOR THE USE OF SONATA CAPSULES FOR UP TO 5 WEEKS (35 NIGHTS) OF TREATMENT IN A CONTROLLED TRIAL SETTING

**PATENT AND EXCLUSIVITY TERMS**

ADB 32 of 117

**EXCLUSIVITY MISCELLANEOUS**

- M-9 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING OF TEXT AND TWO TABLES CONTAINING INFORMATION FOR THE PRESCRIBING PHYSICIAN ON BLOOD PRESSURE, HEART RATE, AND HEART RATE VARIABILITY
- M-10 INFORMATION REGARDING MAINTENANCE OF AN ANTIDEPRESSANT EFFECT UP TO 1 YEAR OF DOSING
- M-11 USE FOR LONG-TERM TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- M-12 NEW LANGUAGE FOR PEDIATRIC USE
- M-13 INFORMATION FROM PEDIATRIC STUDIES ADDED TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION
- M-14 ADDITIONAL CLINICAL TRIAL INFORMATION ADDED TO PEDIATRIC USE SUBSECTION
- M-15 LONGER TERM EFFICACY INFORMATION FOR RISPERIDONE IN THE TREATMENT OF SCHIZOPHRENIA
- M-16 CHANGE IN WORDING OF THE PEDIATRIC SECTION OF THE PACKAGE INSERT
- M-17 INFORMATION REGARDING USE OF ULTANE IN PEDIATRIC PATIENTS WITH CONGENITAL HEART DISEASE
- M-18 INFORMATION DENOTING THE EFFICACY OF REMERON IN MAINTAINING A RESPONSE IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER (MDD)
- M-19 INFORMATION REGARDING USE IN PEDIATRIC PATIENTS TWO YEARS OF AGE AND OLDER
- M-20 LABELING REVISIONS RELATED TO MCCUNE ALBRIGHT SYNDROME
- M-21 COMPARISON DATA ON THE ANTIHYPERTENSIVE EFFECTS OF ATACAND AND COZAAR
- M-22 CHANGE IN TIME TO ONSET OF ACTION
- M-23 INFORMATION REGARDING ELIMINATION ADDED TO CLINICAL PHARMACOLOGY, STUDY RESULTS IN PATIENTS WITH HEPATIC AND RENAL IMPAIRMENT
- M-24 INFORMATION ON RESULTS OF A LONG TERM LONGITUDINAL GROWTH STUDY AND PEDIATRIC SAFETY INFORMATION
- M-25 ADDITIONAL SAFETY AND PHARMACOKINETICS INFORMATION IN CHILDREN 6 MONTHS TO LESS THAN 6 YEARS OF AGE ADDED TO PACKAGE INSERT
- M-26 INCORPORATION OF INFORMATION CONTAINED IN THE PEG-INTRON PACKAGE INSERT INTO THE REBETOL PACKAGE INSERT AND MEDGUIDE-PEG-INTRON WAS APPROVED FOR USE IN COMBINATION WITH REBETOL FOR TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION ON 8/7/01
- M-27 INFORMATION DESCRIBING ASPIRIN ENDOSCOPY STUDY AND THE MAXIMUM RECOMMENDED DOSE FOR PATIENTS WITH MODERATE HEPATIC INSUFFICIENCY
- M-28 INFORMATION FROM A STUDY IN PEDIATRIC PATIENTS IN ASSOCIATION WITH A NEUROLOGICAL CONDITION
- M-29 LABELING CHANGES TO PROVIDE INFORMATION IN THE MANAGEMENT OF OBESITY IN ADOLESCENTS AGED 12 TO 16 YEARS
- M-30 CHANGES TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION SECTIONS OF LABELING CONCERNING USE OF LOTENSIN IN PEDIATRIC PATIENTS WITH HYPERTENSION
- M-31 INFORMATION FOR USE IN PEDIATRIC PATIENTS WITH CHRONIC KIDNEY DISEASE STAGE 5 (END-STAGE RENAL DISEASE)
- M-32 ADDITIONAL LANGUAGE TO CLINICAL PHARMACOLOGY AND CLINICAL STUDIES
- M-33 INFORMATION FOR USE OF ADVAIR DISKUS 100/50 IN CHILDREN 4 TO 11 YEARS OF AGE WITH ASTHMA
- M-34 EXPANDED INFORMATION TO PEDIATRIC USE SUBSECTION OF LABELING IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-35 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH ACTOS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, OR INSULIN ADDED TO CLINICAL PHARMACOLOGY
- M-36 ADDITION OF INFORMATION TO CLINICAL STUDIES REGARDING PREVENTION OF CARDIOVASCULAR DISEASE
- M-37 INFORMATION ADDED TO THE LABELING THAT DETAILS INFORMATION RELATIVE TO STUDIES DONE IN PEDIATRIC POPULATIONS IN THE CLINICAL PHARMACOLOGY AND PEDIATRIC USE SUBSECTIONS
- M-38 SAFETY AND IOP-LOWERING EFFECTS OF TRUSOPT HAVE BEEN DEMONSTRATED IN PEDIATRIC PATIENTS IN A 3 MONTH, MULTI-CENTER DOUBLE MASKED ACTIVE-TREATMENT-CONTROLLED

**PATENT AND EXCLUSIVITY TERMS**

ADB 33 of 117

**EXCLUSIVITY MISCELLANEOUS****TRIAL**

- M-39 FOR LABELING CHANGES BASED ON RESULTS OF THE SPD422-202 CLINICAL STUDY REPORT (CSR) SUBMITTED IN RESPONSE TO THE WRITTEN REQUEST
- M-40 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED IN PEDIATRIC PATIENTS WITH LEUKEMIA ADDED TO PRECAUTIONS
- M-41 REVISION TO THE PEDIATRIC USE PRECAUTIONS OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM THE CAPPS-169 STUDY ENTITLED "THE EFFECT OF ORTHO TRICYCLEN ON BONE MINERAL DENSITY IN PEDIATRIC SUBJECTS WITH ANOREXIA NERVOSA"
- M-42 ADDITION OF A GERIATRIC USE SUBSECTION TO THE PRECAUTIONS SECTION OF THE PACKAGE INSERT AND GERIATRIC DOSING INFORMATION
- M-43 INCLUSION OF RESULTS OF STUDY "PLACEBO-CONTROLLED STUDY TO EVALUATE SAFETY AND PILOT EFFICACY OF ILOPROST AS ADD ON THERAPY WITH BOSENTAN IN SUBJECTS WITH PULMONARY ARTERIAL HYPERTENSION"
- M-44 CLINICAL INFORMATION ADDED TO THE PEDIATRIC USE SUBSECTION OF PRECAUTIONS REGARDING THE USE OF NOVOLOG IN ADOLESCENTS WITH TYPE I DIABETES AGE 6 TO 18
- M-45 INFORMATION ADDED TO CLINICAL TRIALS SECTION OF LABELING, "EFFECTS OF HUMATROPE TREATMENT IN ADULTS WITH GROWTH HORMONE DEFICIENCY"
- M-46 PROVISION OF RESULTS OF STUDY AND PROPOSED REVISIONS TO PACKAGE INSERT SEE SECTION ON CARDIAC ELECTROPHYSIOLOGY
- M-47 PROVIDES FOR USE OF ANTARA WITHOUT REGARD TO MEALS
- M-48 CHANGES TO THE LABELING DESCRIBING THE RESULTS OF A STUDY OF THE USE OF NOVOLOG MIX 70/30 WITH ORAL ANTIDIABETIC AGENTS IN PATIENTS WITH TYPE 2 DIABETES
- M-49 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING EFFECT OF SINGULAIR ON GROWTH RATES IN PREPUBERTAL CHILDREN
- M-50 NEW INFO TO THE CLINICAL STUDIES, ADULT GROWTH HORMONE DEFICIENCY (GHD) SUBSECTION OF THE NUTROPIN AQ PACKAGE INSERT DESCRIBING THE EFFECTS OF SOMATROPIN ON VISCERAL ADIPOSE TISSUE IN THE ADULT GROWTH HORMONE DEFICIENT PATIENT POPULATION
- M-51 INFORMATION ADDED TO LABELING REGARDING OSTEOGENESIS IMPERFECTA STUDY
- M-52 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY/CLINICAL STUDIES SECTION REGARDING THE USE OF RISEDRONATE ADMINISTERED ONCE A WEEK IN THE PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- M-53 FOR LABELING CHANGES TO THE QUALITY OF LIFE (QOL) STATEMENT IN THE APPROVED PACKAGE INSERT
- M-54 INFORMATION FROM PEDIATRIC STUDIES ADDED TO LABEL
- M-55 INFORMATION ON RESULTS OF A STUDY OF THE USE OF SANDOSTATIN LAR DEPOT IN PEDIATRIC PATIENTS WITH HYPOTHALAMIC OBESITY.
- M-56 INFORMATION ADDED TO CLINICAL TRIAL SECTION WITH INFORMATION ON "GEMINI" TRIAL
- M-57 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING THE PHARMACOKINETICS OF EZETIMIBE IN ASIAN SUBJECTS
- M-58 CHANGES TO THE CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, VYTORIN SUBSECTION OF THE PACKAGE INSERT TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR AN ATORVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PRMTS
- M-59 RESULTS OF THE T20-310 STUDY WHICH EVALUATED THE PHARMACOKINETICS, SAFETY, AND ANTIVIRAL ACTIVITY OF FUZEON IN TREATMENT EXPERIENCED PEDIATRIC SUBJECTS AND ADOLESCENTS WAS ADDED TO THE PEDIATRIC SUBSECTION OF PRECAUTIONS
- M-60 CHANGES TO CLINICAL STUDIES, PRIMARY HYPERCHOLESTEROLEMIA, TO ADD EFFICACY DATA FOR THE EZETIMIBE/SIMVASTATIN COMBINATION PRODUCT AND FOR A ROUVASTATIN PRODUCT ON LDL-C AND OTHER LIPID PARAMETERS IN PATIENTS WTH HYPERCHOLESTEROLEMIA
- M-61 REVISIONS TO LABELING BASED ON DATA SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-62 CLINICAL INFORMATION FROM ONE CLINICAL STUDY INVESTIGATING THE USE OF AVANDAMET PLUS INSULIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE NOT ACHIEVED ADEQUATE GLYCEMIC CONTROL WITH PREVIOUS ANTI-DIABETIC THERAPIES
- M-63 DETAILED INFORMATION ON AN INCONCLUSIVE PEDIATRIC STUDY
- M-64 CHANGES TO CLINICAL PHARMACOLOGY DETAILING STUDY RESULTS

**PATENT AND EXCLUSIVITY TERMS**

ADB 34 of 117

**EXCLUSIVITY MISCELLANEOUS**

- M-65 ADDITION OF INFORMATION TO LABEL TO INCLUDE INFORMATION REGARDING USE IN PATIENTS WITH HIV-ASSOCIATED ADIPOSE REDISTRIBUTION SYNDROME (HARS)
- M-66 USE IN SPECIFIC POPULATIONS - PATIENTS WITH CONCOMITANT ILLNESS SUBSECTION OF THE LABELING REGARDING USE OF STRATTERA IN PATIENTS WITH ADHD WHO HAVE COMORBID TIC DISORDER
- M-67 INDICATION EXPANDED TO INCLUDE PATIENTS ON PERITONEAL DIALYSIS
- M-68 DESCRIPTION OF RESULTS OF STUDY OF INITIAL THERAPY IN COMBINATION WITH METFORMIN WHEN DIET AND EXERCISE DO NOT PROVIDE GLYCEMIC CONTROL
- M-69 RESULTS OF STUDY OF COMBINATION THERAPY AND NON-INFERIORITY STUDY
- M-70 PROVISION OF INFORMATION OF THE RESULTS OF A PHASE 2 RANDOMIZED TRIAL OF SPRYCEL 70MG TWICE DAILY OR IMATINIB 800MG DAILY
- M-71 REVISIONS TO PROVIDE FOR RESULTS OF MAINTENANCE DATA IN ADULT PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- M-72 INFORMATION ABOUT USE OF INSPRA (EPLERENONE) FOR HYPERTENSION IN PEDIATRIC PATIENTS
- M-73 NEW INFORMATION ADDED REGARDING THE TUMOR SHRINKING POTENTIAL OF SANDOSTATIN LAR DEPOT INJECTION ON GH - SECRETING PITUITARY ADENOMAS
- M-74 REVISIONS TO CLINICAL STUDIES - CHILDREN AND ADOLESCENTS BASED ON CLINICAL TRIAL DATA TO SUPPORT A DURATION OF ACTION CLAIM UP TO 12 HOURS
- M-75 PROVISION FOR USE OF ARGATROBAN IN CERTAIN PEDIATRIC PATIENTS WITH HEPARIN-INDUCED THROMBOCYTOPENIA (HIT) OR HEPARIN-INDUCED THROMBOCYTOPENIA WITH THROMBOSIS (HITTS)
- M-76 REMOVAL OF SCREEN REQUIREMENT IN PTS WITH G6PD DEFICIENCY PRIOR TO INITIATING ACZONE TREATMENT; REMOVAL OF BLOOD COUNT & RETICULOCYTE MONITORING DURING TREATMENT IN G6PD DEFICIENT PTS AND IN PATIENTS WITH HISTORY OF ANEMIA
- M-77 USE IN COMBINATION WITH THE NEW AKTILITE CL128 LAMP FOR THE TREATMENT OF THIN AND MODERATELY THICK, NON-HYPERKERATOTIC, NON-PIGMENTED ACTINIC KERATOSES OF THE FACE AND SCALP IN IMMUNOCOMPETENT PATIENTS
- M-78 CLINICAL TRIAL INFO ON USE OF STRATTERA IN PATIENTS WITH ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) AND COMORBID ANXIETY DISORDER WITHOUT CAUSING WORSENING OF ANXIETY
- M-79 LABELING REVISIONS RELATED TO SMOKING AND ERLOTINIB EXPOSURE
- M-80 ADDITIONAL TIME POINT OF 30 MINUTES (0.5 HOUR) IN CHILDREN AGED 6-12 YEARS WITH A DIAGNOSIS OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER (ADHD)
- M-81 ADDITIONAL INFO FOR PEDIATRIC USE FOR CASODEX (STUDIED IN COMBINATION WITH ARIMIDEX) IN THE PEDIATRIC POPULATION, SPECIFICALLY BOYS WITH FAMILIAL MALE-LIMITED PRECOCIOUS PUBERTY (TESTOXICOSIS)
- M-82 LABELING REVISIONS RELATED TO CLINICAL STUDIES
- M-83 ADDITIONAL INFORMATION ADDED TO LABELING REGARDING ESTABLISHMENT OF EFFICACY IN ADDITIONAL CLINICAL TRIALS AND ONE MAINTENANCE TRIAL
- M-84 STUDY INFORMATION ADDED TO LABEL REGARDING BONE MINERAL DENSITY
- M-85 INFORMATION ADDED TO LABELING REGARDING USE OF PREVACID IN PATIENTS LESS THAN 1 YEAR WITH SYMPTOMATIC GERD
- M-86 LABELING CHANGES SUBMITTED IN RESPONSE TO PEDIATRIC WRITTEN REQUEST FOR INFANTS AGES BIRTH TO 11 MONTH INCLUSIVE REFLECTING LACK OF EFFICACY FOR GERD INDICATION FOR THIS PATIENT POPULATION
- M-87 INCLUSION OF RESULTS FROM TWO DRUG INTERACTION STUDIES WITH LIPITOR AND CRESTOR IN CLINICAL PHARMACOLOGY SECTION
- M-88 ADDITION OF INFORMATION REGARDING ABUSE POTENTIAL OF CONCERTA VERSUS IMMEDIATE-RELEASE METHYLPHENIDATE
- M-89 PROVIDES FOR REVISIONS TO MULTIPLE SECTIONS OF THE PACKAGE INSERT TO REFLECT RESULTS OF CLINICAL TRIALS 205.235 (UPLIFT) AND 205.266 (VA STUDY) IN SUPPORT OF EXACERBATION CLAIM
- M-90 LABELING CHANGES BASED ON DATA FROM CLINICAL STUDIES NV20235 AND NV20236 STUDIES OF SEASONAL PROPHYLAXIS OF INFLUENZA IN IMMUNOCOMPROMISED PATIENTS AND CHILDREN AGES 1-12
- M-91 UPDATED LABELING BASED UPON STUDY: A SINGLE-DOSE, SINGLE-BLIND, PLACEBO-AND MOXIFLOXACIN-CONTROLLED 2-PERIOD, RANDOMIZED, CROSSOVER, 3RD PERIOD SEQUENTIAL

**PATENT AND EXCLUSIVITY TERMS**

ADB 35 of 117

**EXCLUSIVITY MISCELLANEOUS**

STUDY OF SIDE EFFECTS OF TEMSIROLIMUS ON CARDIAC REPOLARIZATION IN HEALTHY SUBJECTS

- M-92 UPDATES TO THE PACKAGE INSERT BASED UPON THE TRIAL ENTITLED "A PHASE I PHARMACOKINETIC AND PHARMACODYNAMIC STUDY OF TEMSIROLIMUS IN PATIENTS WITH ADVANCED MALIGNANCIES AND NORMAL AND IMPAIRED LIVER FUNCTION"
- M-93 EXPANSION OF LABELING TO INCLUDE INFORMATION ON SAFETY AND EFFICACY OF CREON IN PATIENTS AGES 7 YEARS THROUGH 11 YEARS WITH PANCREATIC EXOCRINE INSUFFICIENCY DUE TO CYSTIC FIBROSIS
- M-94 INFO ADDED TO LABEL RELATED TO NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE (Ph+) CHRONIC MYELOID LEUKEMIA IC CHRONIC PHASE
- M-95 INFORMATION FOR TREATMENT OF CHRONIC HEPATITIS B (CHB) IN ADULT PATIENTS WITH DECOMPENSATED LIVER DISEASE BASED ON DATA FROM CLINICAL TRIAL GS-US-174-0108
- M-96 UPDATED INFORMATION IN THE CLINICAL STUDIES SECTION RELATED TO THE LOSS AND RECOVERY OF BONE MINERAL DENSITY IN ADOLESCENT GIRLS DURING AND FOLLOWING THE USE OF DEPO-PROVERA CONTRACEPTIVE INJECTION
- M-97 LABELING CHANGES IN RESPONSE TO PEDIATRIC STUDIES - NOT INDICATED FOR USE IN PEDIATRIC POPULATION
- M-98 NEW INFORMATION FROM A STUDY WHICH EVALUATED THE SAFETY AND EFFICACY OF FAMVIR IN TREATING RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT BLACK/AFRICAN AMERICAN SUBJECTS.
- M-99 ADDITION OF FINDINGS FROM A SINGLE PEDIATRIC CLINICAL TRIAL (P04292) OF NASONEX NASAL SPRAY IN THE TREATMENT OF NASAL POLYPs IN PATENTS 6 TO <18 YEARS OF AGE TO THE PACKAGE INSERT.
- M-100 INFORMATION ADDED TO LABEL BASED UPON COMPLETED CLINICAL TRIAL REPORTS
- M-101 INCLUSION OF DATA FROM AN ADDITIONAL 19 SUBJECTS WITH HYPERCALCEMIA FROM PARATHYROID CARCINOMA TO THE INFORMATION CURRENTLY PRESENTED IN THE LABEL
- M-102 INFORMATION FROM PEDIATRIC STUDY REPORT ML16633, "INTRAVENOUS GRANisetron (KYTRIL) IN THE PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING (PONV) IN PEDIATRIC SUBJECTS UNDERGOING TONSILLECTOMY OR ADENOTONSILLECTOMY."
- M-103 SAFETY, EFFICACY AND PHARMACOKINETIC INFO FOR FASLODEX IN THE PEDIATRIC POPULATION, SPECIFICALLY FOR GIRLS WITH PROGRESSIVE PRECOCIOUS PUBERTY ASSOCIATED WITH MCCUNE-ALBRIGHT SYNDROME ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING
- M-104 INFORMATION ADDED TO DOSING AND ADMINISTRATION REGARDING A 26 WEEK STUDY
- M-105 NEW LANGUAGE ADDED TO CLINICAL STUDIES REGARDING USE IN SMOKERS WITH CARDIOVASCULAR DISEASE, CHRONIC OBSTRUCTIVE PULMONARY DISEASE, AND USE ACCORDING TO AN ALTERNATIVE SET OF DIRECTIONS FOR SETTING A QUIT DATE
- M-106 ADDITION OF THE T1-WEIGHTED GD-ENHANCED LESION EFFICACY VARIABLE IN THE CLINICAL STUDIES SECTION 14 OF THE PACKAGE INSERT
- M-107 INFORMATION TO THE CLINICAL STUDIES SECTION OF THE LUPRON DEPOT-PED 1-MONTH BASED UPON THE PHASE 3/4 COMPLETED CLINICAL STUDY REPORT FOR STUDY M90-516 ENTITLED "STUDY OF LUPRON DEPOT IN THE TREATMENT OF CENTRAL PRECOCIOUS PUBERTY".
- M-108 CHANGES ARE BASED ON RESULTS FROM STUDY CV181057
- M-109 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-110 CHANGES TO THE PACKAGE INSERT TO REFLECT THE RESULTS OF THE STUDY OF HEART AND RENAL PROTECTION (SHARP) TRIAL
- M-111 LABELING CHANGES BASED ON STUDY HW80-EW-GWCI ENTITLED A PLACEBO AND POSITIVE CONTROLLED STUDY OF THE ELECTROPHYSIOLOGICAL EFFECTS OF A SINGLE 10 MCG DOSE OF EXENATIDE ON THE 12 LEAD ELECTROCARDIOGRAM QT INTERVAL IN HEALTHY SUBJECTS
- M-112 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO ADD INFORMATION FROM A PEDIATRIC STUDY IN PATIENTS AGED 12 YEARS TO LESS THAN 18 YEARS OF AGE WITH RECURRENT HERPES LABIALIS
- M-113 LABELING CHANGES BASED ON STUDY H80-US-GWCO ENTITLED A RANDOMIZED TRIAL COMPARING EXENATIDE WITH PLACEBO IN SUBJECTS WITH TYPE 2 DIABETES ON INSULIN GLARGINE WITH OR WITHOUT ORAL ANTIHYPERGLYCEMIC MEDICATIONS
- M-114 CHANGES IN SECTION 14 OF THE PACKAGE INSERT TO INCLUDE DATA FROM THE SWITCHMRK STUDIES (SWITCH OF SUPPRESSED SUBJECTS FROM LOPINAVIR/RITONAVIR TO Raltegravir)

**PATENT AND EXCLUSIVITY TERMS**

ADB 36 of 117

**EXCLUSIVITY MISCELLANEOUS**

- M-115 REVISIONS TO THE PI BASED ON RESULTS FROM STUDY NN2211-1842, ENTITLED THE EFFECT OF INSULIN DETEMIR IN COMBINATION WITH LIRAGLUTIDE AND METFORMIN COMPARED TO LIRAGLUTIDE AND METFORMIN IN SUBJECTS WITH TYPE 2 DIABETES
- M-116 LABELING CHANGES BASED ON RESULTS FROM CLINICAL STUDY 01-06-TL-OPIMET-008
- M-117 ADDITION OF RESULTS OF PEDIATRIC TRIAL TO LABEL
- M-118 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.36
- M-119 LABELING CHANGES REGARDING MISSED DOSES
- M-120 CHANGES TO CLINICAL TRIALS DETAILING STUDY RESULTS
- M-121 LABELING CHANGES BASED UPON SAFETY AND EFFICACY RESULTS FROM TRIAL 1218.43
- M-122 LABELING CHANGES TO INCLUDE THE RESULTS OF THE PARAMOUNT TRIAL
- M-123 UPDATED RESULTS OF OVERALL SURVIVAL FROM 'CONFIRM' STUDY
- M-124 LONG TERM SAFETY AND EFFICACY DATA FROM STUDY CLDT600A2303 FOR SUBJECTS PREVIOUSLY ENROLLED IN THE ORIGINAL TWO YEAR GLOBE (NV-02B-007/CLDT600A2302) AND NV02B-015 STUDIES WHO CONTINUED TELBIVUDINE TREATMENT FOR UP TO 208 WEEKS
- M-125 LABELING CHANGES TO INCLUDE LACK OF EFFICACY IN CHILDREN 6 MONTHS TO 4 YEARS OF AGE
- M-126 UPDATES TO THE CLINICAL STUDIES SECTION 14, OF THE PACKAGE INSERT (PI), WITH THE RESULTS OF CLINICAL TRIAL P06086
- M-127 REVISIONS TO THE PEDIATRIC USE SECTION OF THE PACKAGE INSERT TO REFLECT THE RESULTS FROM CLINICAL STUDY C-10-004
- M-128 CLINICAL TRIAL STUDY RESULTS
- M-129 RESULTS OF A CLINICAL STUDY REPORT WHICH ASSESSES THE SAFETY AND EFFICACY IN CHILDREN AGES 6 TO 12 YEARS OF AGE
- M-130 ADDITION OF INFORMATION ON LONG-TERM TREATMENT WITH VPRIV IN THE CLINICAL TRIALS SECTION OF THE PACKAGE INSERT
- M-131 INFORMATION FROM STUDIES CONDUCTED IN PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED NON-DISSEMINATED DIFFUSED INTRINSIC BRAINSTEM GLIOMAS
- M-132 REVISIONS TO THE CLINICAL TRIALS SECTION IN THE INOMAX LABEL TO REFLECT RESULTS FROM THE PEDIATRIC STUDY REPORTS
- M-133 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF SILDENAFIL TO BOSENTAN THERAPY
- M-134 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED WITH SAXagliptin IN COMBINATION WITH METFORMIN AND A SULFONYLUREA ADDED TO THE LABELING
- M-135 ADDITION OF INFORMATION TO THE CLINICAL STUDIES - RADIOGRAPHIC RESPONSE SECTION OF THE PACKAGE INSERT
- M-136 ADDITIONAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING REGARDING POST-OPERATIVE NAUSEA AND VOMITING STUDIES IN PEDIATRIC PATIENTS
- M-137 LABELING REVISIONS RESULTING FROM A MAINTENANCE TRIAL IN PEDIATRIC PATIENTS WITH IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- M-138 INFORMATION ADDED TO THE 8.4 PEDIATRIC USE SECTION ON THE USE OF MEMANTINE IN CHILDREN AGES 6-12 YEARS WITH AUTISM SPECTRUM DISORDER
- M-139 INFORMATION ADDED TO THE DOSING AND ADMINISTRATION SECTION OF THE PACKAGE INSERT REGARDING RETREATMENT WITH VELCADE FOR PATIENTS WITH MULTIPLE MYELOMA
- M-140 INFORMATION ADDED TO THE PEDIATRIC USE SECTION OF THE LABELING REGARDING USE OF LATISSE IN PATIENTS WHO WERE POST-CHEMOTHERAPY OR HAD ALOPECIA AREATA, AND ADOLESCENTS WHO HAD HYPERTRICHOSIS WITH NO ASSOCIATED MEDICAL CONDITION
- M-141 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING TO INCORPORATE STUDY RESULTS FOR TREATMENT OF MAJOR DEPRESSIVE DISORDER IN ADOLESCENTS (AGES 12-17)
- M-142 ADDITIONS TO THE LABELING DESCRIBING RESULTS FROM STUDY H6P-MC-HDAY
- M-143 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSION IN PATIENTS WITH CURRENT OR PAST HISTORY OF MAJOR DEPRESSIVE DISORDER
- M-144 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSION IN PATIENTS WHO HAD BEEN PREVIOUSLY TREATED WITH VARENICLINE

**PATENT AND EXCLUSIVITY TERMS**

ADB 37 of 117

**EXCLUSIVITY MISCELLANEOUS**

- M-145 ADDITION OF INFORMATION ABOUT LONG-TERM TREATMENT OF PULMONARY ARTERIAL HYPERTENSION TO THE CLINICAL STUDIES SECTION OF THE LABELING
- M-146 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION ON INITIAL COMBINATION THERAPY WITH LINAGLIPTIN AND METFORMIN VS. LINAGLIPTIN MONOTHERAPY IN TREATMENT NAIVE PATIENTS
- M-147 OTC USE FOR TEMPORARY RELIEF OF OCULAR SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES
- M-148 LABELING CHANGES BASED ON STUDY H80-EW-GWDM
- M-149 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE MONOTHERAPY FOR ADHD
- M-150 ADDITION OF THE RESULTS OF A CONTROLLED CLINICAL STUDY TREATING ADULT PATIENTS WITH SCHIZOPHRENIA EXPERIENCING AN ACUTE RELAPSE
- M-151 REVISIONS TO THE LABELING BASED ON THE OUTCOMES OF PEDIATRIC STUDIES CONDUCTED TO ASSESS THE SAFETY AND EFFICACY OF XOPENEX IN SUBJECTS LESS THAN 6 YEARS OF AGE
- M-152 INFORMATION ADDED TO THE CLINICAL PHARMACOLOGY SECTION OF THE LABELING REGARDING A SAFETY STUDY IN PEDIATRIC SUBJECTS AGES 6 MONTHS TO 4 YEARS OF AGE WITH AN ACTIVE HEAD LICE INFESTATION
- M-153 ADDITION OF INFORMATION REGARDING THE INTRANASAL ABUSE POTENTIAL OF OXYCONTIN
- M-154 UPDATE TO THE LABELING TO REFLECT THE RESULTS OF A LONG-TERM MAINTENANCE TREATMENT STUDY OF ADHD IN CHILDREN AND ADOLESCENTS AGES 6-17.
- M-155 ADDITION OF CLINICAL FINDINGS FROM AN OBSERVATIONAL STUDY IN A PEDIATRIC AGE GROUP GREATER THAN 2 MONTHS TO 18 YEARS IN SECTION 8.4 PEDIATRIC USE OF THE PACKAGE INSERT
- M-156 UPDATE TO THE LABELING WITH INFORMATION REGARDING A CLINICAL TRIAL IN CHILDREN LESS THAN 4 YEARS OF AGE.
- M-157 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN 10MG ONCE DAILY IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND SULFONYLUREA
- M-158 UPDATES TO THE LABELING TO REFLECT SAFETY RESULTS FROM CLINICAL TRIALS IN SCHIZOPHRENIA ADOLESCENT PATIENTS AGED 12 TO 17 YEARS
- M-159 ADDITION OF PED SAFETY INFORMATION DERIVED FROM A MAINTENANCE TREATMENT STUDY OF BIPOLAR 1 DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES IN PATIENTS (> THAN OR = TO 13 YRS OF AGE) TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- M-160 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND ACTIVE-CONTROLLED STUDY COMPARING EMPAGLIFLOZIN TO GLIMEPIRIDE IN PATIENTS WITH TYPE 2 DIABETES AND INSUFFICIENT GLYCEMIC CONTROL DESPITE METFORMIN TREATMENT
- M-161 UPDATED LABELING WITH DATA FROM A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY OF EMPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES MELLITUS AND INSUFFICIENT GLYCEMIC CONTROL ON A MULTIPLE DAILY INJECTION INSULIN REGIMEN ALONE OR WITH METFORMIN
- M-162 INCLUSION OF EFFICACY AND SAFETY DATA TO THE PRESCRIBING INFORMATION OF BYDUREON BASED ON STUDY GWDE
- M-163 INFORMATION ADDED TO THE LABELING REGARDING PREVIOUSLY UNTREATED ALK-POSITIVE METASTATIC NON SMALL CELL LUNG CANCER (NSCLC)
- M-164 REVISES THE CLINICAL TRIALS SECTION OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM STUDY E7273-G000-401 ENTITLED "PHASE IV RANDOMIZED STUDY OF TWO DOSE LEVELS OF TARGRETIN CAPSULES IN SUBJECTS WITH REFRACTORY CUTANEOUS T-CELL LYMPHOMA"
- M-165 PROVIDES FOR UPDATES TO THE PEDIATRIC USE SECTION BASED ON THE PEDIATRIC STUDY REPORT ENTITLED, "A PHASE II PILOT TRIAL OF BORTEZOMIB IN COMBINATION WITH INTENSIVE RE-INDUCTION THERAPY IN CHILDREN WITH RELAPSED ACUTE LYMPHOBLASTIC LYMPHOMA (LL)"
- M-166 UPDATE TO LABELING WITH WEEK 48 RESULTS FROM VIKING-4 IN ANTIRETROVIRAL THERAPY (ART) - EXPERIENCED INTEGRASE STRAND TRANSFER INHIBITOR (INSTI) - RESISTANT SUBJECTS
- M-167 APPROVED FOR REVISIONS TO THE LABELING BASED ON THE CLINICAL STUDY ENTITLED "BRONCHOPULMONARY DYSPLASIA (BPD) IN PRETERM INFANTS REQUIRING MECHANICAL VENTILATION OR POSITIVE PRESSURE SUPPORT ON DAYS 5 TO 14 AFTER BIRTH".

**PATENT AND EXCLUSIVITY TERMS**

ADB 38 of 117

**EXCLUSIVITY MISCELLANEOUS**

- M-168 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE RE-NOVATE AND RE-NOVATE LL STUDIES (PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM FOLLOWING HIP REPLACEMENT SURGERY)
- M-169 UPDATES TO LABELING DESCRIBING RESPONSE TO A REPEAT COURSE OF PICATO GEL 0.015% ON THE FACE OR SCALP IF AN INCOMPLETE RESPONSE IS OBSERVED AT A FOLLOW-UP EXAMINATION.
- M-170 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION REGARDING USE FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- M-171 UPDATES TO LABELING WITH RESULTS TO THE TIGER CLINICAL TRIAL
- M-172 UPDATES TO THE CLINICAL TRIALS SECTION OF THE LABELING TO INCLUDE RESULTS OF STUDIES PERFORMED TO EVALUATE THE BENEFIT OF ADDING INCRUNE ELLIPTA TO PATIENTS WHO ARE ON BACKGROUND THERAPY WITH BREO ELLIPTA AND ADVAIR DISKUS
- M-173 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING DESCRIBING THE EFFECTS OF STIOLTO RESPIMAT ON COPD PATIENTS
- M-174 INFORMATION ADDED TO CLINICAL STUDIES SECTION OF THE LABELING REGARDING INITIAL COMBINATION THERAPY OF EMPAGLIFLOZIN WITH METFORMIN
- M-175 INFORMATION ADDED TO THE LABELING DESCRIBING SAVOR, A PHASE IV TRIAL EVALUATING THE EFFECT OF SAXagliptin ON THE INCIDENCE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION OR ISCHAEMIC STROKE IN PATIENTS WITH TYPE 2 DIABETES
- M-176 INFORMATION ADDED TO THE LABELING DESCRIBING TRIAL NN2211-3916, A TRIAL EVALUATING THE SAFETY AND EFFICACY OF LIRAGLUTIDE IN SUBJECTS WITH TYPE 2 DIABETES AND MODERATE RENAL IMPAIRMENT
- M-177 INFORMATION ADDED TO THE LABELING DESCRIBING EXAMINE, A TRIAL EVALUATING CARDIOVASCULAR ISCHEMIC RISKS ASSOCIATED WITH ALOGLIPTIN USE IN PATIENTS WITH TYPE 2 DIABETES AT HIGH RISK OF ISCHEMIC CARDIOVASCULAR DISEASE
- M-178 INFORMATION ADDED TO THE LABELING REGARDING MAINTENANCE OF REMISSION IN CROHN'S DISEASE IN PEDIATRIC PATIENTS
- M-179 UPDATES TO THE PRODUCT LABELING WITH STUDY REPORTS FROM THE OPTIMIST-1 AND OPTIMIST-2 CLINICAL TRIALS
- M-180 INFORMATION ADDED TO THE LABELING REGARDING THE ADDITION OF MAINTENANCE TREATMENT IN PATIENTS WITH SCHIZOPHRENIA
- M-181 UPDATE TO THE DOSAGE AND ADMINISTRATION, PATIENT SELECTION (2.1), SECTION OF THE PACKAGE INSERT TO INCLUDE THE USE OF AN FDA-APPROVED PLASMA TEST FOR THE IDENTIFICATION OF EGFR EXON 19 DELETION OR EXON 21 (L858R) SUBSTITUTION MUTATIONS
- M-182 UPDATES TO THE PRODUCT LABELING BASED ON THE RESULTS OF STUDY H7T-MC-TADO TITLED, "A PHASE 3 DOUBLE-BLIND, RANDOMIZED, MULTICENTER, EFFICACY AND SAFETY STUDY OF PRASUGREL COMPARED TO PLACEBO IN PEDIATRIC PATIENTS WITH SICKLE CELL DISEASE"
- M-183 CHANGES TO THE DOSAGE AND ADMINISTRATION AND CLINICAL STUDIES SECTIONS OF THE LABELING TO SUPPORT THE REDUCE-TO-QUIT PARADIGM
- M-184 UPDATES MADE TO THE LABELING TO INCLUDE INFORMATION FROM STUDY MO25743 ON THE ANTI-TUMOR ACTIVITY OF VEMURAFENIB IN THE TREATMENT OF PATIENTS WITH BRAF V600E MUTATION-POSITIVE MELANOMA WITH BRAIN METASTASES
- M-185 UPDATES TO THE LABELING TO INCLUDE RESULTS OF A TRIAL TO EVALUATE THE SAFETY OF MOXIFLOXACIN IN PEDIATRIC PATIENTS WITH COMPLICATED INTRA-ABDOMINAL INFECTIONS
- M-186 UPDATES TO THE PRODUCT INFORMATION REGARDING MAINTENANCE TREATMENT OF SCHIZOPHRENIA IN ADULTS BASED UPON THE RESULTS FROM STUDY 331-10-232
- M-187 ADDITION OF CLINICAL INFORMATION OBTAINED FROM A PEDIATRIC TRIAL TO SECTION 8.4 OF THE LABELING
- M-188 PROVIDES FOR DATA SUPPORTING THE SAFETY AND EFFECTIVENESS FOR THE MAINTENANCE TREATMENT OF MODERATE TO SEVERE BINGE EATING DISORDER (BED)
- M-189 LABELING DESCRIBING THE EXPECTED REDUCTION OF ABUSE OF SINGLE-ENTITY MORPHINE BY THE INTRANASAL ROUTE OF ADMINISTRATION DUE TO PHYSICOCHEMICAL PROPERTIES
- M-190 INFORMATION ADDED TO THE CLINICAL STUDIES SECTION OF THE LABELING REGARDING THE LACK OF EFFICACY OF TARCEVA IN MAINTENANCE TREATMENT OF PATIENTS WITHOUT EGFR MUTATIONS
- M-191 ADDITION OF DATA BASED ON PEDIATRIC STUDIES TO FULFILL THE POSTMARKETING

**PATENT AND EXCLUSIVITY TERMS**

ADB 39 of 117

**EXCLUSIVITY MISCELLANEOUS****REQUIREMENT 1857-2**

- M-192 PROVIDES FOR DATA EVALUATING THE NEUROPSYCHIATRIC SAFETY AND EFFICACY OF VARENICLINE FOR SMOKING CESSATION IN SUBJECTS WITH AND WITHOUT A HISTORY OF PSYCHIATRIC DISORDERS
- M-193 INFORMATION ADDED TO THE LABELING REGARDING A 15-WEEK, RANDOMIZED, DOUBLE-BLIND, PARALLEL-GROUP, PLACEBO-CONTROLLED FLEXIBLE-DOSE SAFETY AND EFFICACY STUDY OF PREGABALIN IN ADOLESCENTS (12 THROUGH 17 YEARS OLD) WITH FIBROMYALGIA
- M-194 INFORMATION ADDED TO THE LABELING REGARDING USE OF REGADENOSON ADMINISTRATION FOLLOWING AN INADEQUATE EXERCISE STRESS TEST AS COMPARED TO REGADENOSON ALONE
- M-195 REVISIONS TO THE PEDIATRIC USE SECTION OF THE LABELING REFLECTING LACK OF EFFICACY FOR IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER IN PEDIATRIC PATIENTS AGES 6-17
- M-196 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM A RANDOMIZED, PLACEBO CONTROLLED, MULTICENTER STUDY OF INTRAVENOUS ACETAMINOPHEN FOR THE TREATMENT OF ACUTE PAIN IN PEDIATRIC PATIENTS TO FULFILL THE POST-MARKETING REQUIREMENT 1704-1
- M-197 NEW CLINICAL DATA ADDED TO THE PRESCRIBING INFORMATION REGARDING CANAGLIFLOZIN ADD-ON COMBINATION THERAPY WITH METFORMIN AND A DIPEPTIDYL-PEPTIDASE-4 INHIBITOR
- M-198 PACKAGE INSERT UPDATED WITH RESULTS FROM STUDY CV181168, A MULTICENTER, RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED, PARALLEL GROUP, PHASE 3 TRIAL TO EVALUATE THE SAFETY AND EFFICACY OF SAXAGLIPTIN ADDED TO DAPAGLIFLOZIN AND METFORMIN
- M-199 INFORMATION ADDED TO LABELING REGARDING THE TREATMENT OF PATIENTS WITH ALK-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAD NOT RECEIVED PRIOR SYSTEMIC THERAPY FOR METASTATIC DISEASE.
- M-200 CLINICAL INFORMATION ADDED TO THE USE IN SPECIFIC POPULATIONS SECTION OF THE LABELING.
- M-201 REVISIONS TO THE PACKAGE INSERT BASED ON DATA FROM AN OPEN LABEL, MULTI-CENTER STUDY OF CABAZITAXEL IN PEDIATRIC PATIENTS WITH REFRACTORY SOLID TUMORS INCLUDING TUMORS OF THE CENTRAL NERVOUS SYSTEM.
- M-202 INCLUSION OF DATA FROM THE SUMMIT STUDY FOR BREO ELLIPTA (FLUTICASONE FUROATE/VILANTEROL TRIFENATATE) INHALATION POWDER IN THE PACKAGE INSERT.
- M-203 PROVIDES FOR REVISIONS TO THE PACKAGE INSERT TO REFLECT RESULTS OF TWO POSTMARKETING REQUIREMENT STUDIES ROP111662 AND ROP111569
- M-204 CLINICAL INFORMATION ADDED TO THE PACKAGE INSERT REGARDING USE OF ATORVASTATIN IN CHILDREN AND ADOLESCENTS AGES 10-17 WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- M-205 INFORMATION ADDED TO THE LABELING REGARDING RANDOMIZED, MULTICENTER, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDIES ON PATIENTS WITH SEVERE RENAL IMPAIRMENT
- M-206 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM COMPLERA TO ODEFSEY
- M-207 INFORMATION ADDED TO LABELING REGARDING 48 WEEK EFFICACY, RESISTANCE AND SAFETY DATA ON VIROLOGICALLY SUPPRESSED HIV-1 INFECTED ADULTS SWITCHING FROM ATRIPLA TO ODEFSEY
- M-208 INFORMATION ADDED TO THE LABELING TO INCLUDE RESULTS OF A RANDOMIZED, DOUBLE-BLIND, PLACEBO-CONTROLLED STUDY IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS
- M-209 INFORMATION ADDED TO THE LABELING REGARDING CABAZITAXEL AT 20 MG/M<sup>2</sup> BASED ON THE RESULTS OF THE PROSELICA STUDY
- M-210 INFORMATION ADDED TO LABELING TO SUPPORT THE USE OF SYMBICORT TO REDUCE EXACERBATIONS IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- M-211 PROVIDES FOR LABELING CHANGES REGARDING THE USE OF DAPTOMYCIN IN THE PEDIATRIC POPULATION FOR STAPHYLOCOCCUS AUREUS BACTEREMIA (SAB) BASED ON RESULTS OF A TRIAL IN PEDIATRIC PATIENTS 1 TO 17 YEARS OF AGE
- M-212 INFORMATION ADDED TO THE LABELING REGARDING THE SAFETY AND EFFICACY OF DAPAGLIFLOZIN IN PATIENTS WITH TYPE 2 DIABETES WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A BACKGROUND COMBINATION OF METFORMIN AND EXENATIDE EXTENDED RELEASE
- M-213 INFORMATION ADDED TO THE LABELING TO INCLUDE THE EFFICACY AND SAFETY OF CARIPRAZINE RELATIVE TO PLACEBO IN THE PREVENTION OF RELAPSE OF SYMPTOMS IN

**PATENT AND EXCLUSIVITY TERMS**

ADB 40 of 117

**EXCLUSIVITY MISCELLANEOUS****PATIENTS WITH SCHIZOPHRENIA**

- M-214 INFORMATION ADDED TO THE CLINICAL TRIALS SECTION OF THE LABELING REGARDING A POSTMARKETING SAFETY AND EFFICACY STUDY EVALUATING THE RISK OF SERIOUS ASTHMA-RELATED EVENTS
- M-215 INFORMATION ADDED TO THE LABELING REGARDING THE COMPARISON OF PALIPERIDONE PALMITATE COMPARED WITH ORAL ANTIPSYCHOTIC TREATMENT IN DELAYING TIME TO TREATMENT FAILURE IN ADULTS WITH SCHIZOPHRENIA WHO HAVE BEEN INCARCERATED
- M-216 UPDATE THE PRESCRIBING INFORMATION AND PATIENT LABELING WITH FINDINGS FROM STUDY RP103-08 CONDUCTED IN TREATMENT-NAIVE NEPHROPATHIC CYSTINOSIS PATIENTS TO EXPAND THE INDICATED POPULATION TO PATIENTS 1 YEAR AND OLDER
- M-217 INCORPORATION OF THE LABELING REVISIONS PROVIDED FOR IN NDA 022253/S-039 AND NDA 022255/S-022 INTO THE LACOSAMIDE INJECTION LABELING

**ORPHAN DRUG EXCLUSIVITY**

- ODE-1 TO REDUCE CHRONIC DROOLING IN PATIENTS AGED 3 - 16 WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING (E.G. CEREBRAL PALSY)
- ODE-2 FOR TREATMENT OF NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE
- ODE-3 TO TREAT INFANTILE SPASMS
- ODE-4 TREATMENT OF PATIENTS WITH SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) ASSOCIATED WITH TUBEROUS SCLEROSIS WHO REQUIRE THERAPEUTIC INTERVENTION BUT ARE NOT CANDIDATES FOR CURATIVE SURGICAL RESECTION
- ODE-5 FOR SEQUENTIAL USE FOR THE TREATMENT OF CYANIDE POISONING THAT IS JUDGED TO BE LIFE-THREATENING
- ODE-6 FOR THE MANAGEMENT OF POSTHERPETIC NEURALGIA
- ODE-7 TO REDUCE THE RISK OF PRETERM BIRTH IN WOMEN WITH SINGLETON PREGNANCY WHO HAVE A HISTORY OF SINGLETON SPONTANEOUS PRETERM BIRTH
- ODE-8 TREATMENT OF SEVERE HYPERCALCEMIA IN PATIENTS WITH PRIMARY HYPERPARATHYROIDISM WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY
- ODE-9 TREATMENT OF ASYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER IN PATIENTS WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-10 FOR USE IN COMBINATION CHEMOTHERAPY WITH 5-FLUOROURACIL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED METASTATIC COLORECTAL CANCER
- ODE-11 TREATMENT OF PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) IN PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-12 TREATMENT OF PERIPHERAL T-CELL LYMPHOMA (PTCL) IN PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-13 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA WITH THE BRAFV600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-14 TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA IN ADULTS 18 YEARS OF AGE AND OLDER
- ODE-15 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) THAT IS ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA APPROVED TEST
- ODE-16 TREATMENT OF PATIENTS WITH TRANSFUSIONAL IRON OVERLOAD DUE TO THALASSEMIA SYNDROMES WHEN CURRENT CHELATION THERAPY IS INADEQUATE
- ODE-17 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME IN PATIENTS 2 YEARS OF AGE OR OLDER
- ODE-18 ADJUNCTIVE TREATMENT OF SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME INPATIENTS 2 YEARS OF AGE OR OLDER
- ODE-19 TREATMENT OF PATIENTS WITH INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS, INCLUDING PRIMARY MYELOFIBROSIS, POST-POLYCYTHEMIA VERA MYELOFIBROSIS AND POST-ESSENTIAL THROMBOCYTHEMIA MYELOFIBROSIS
- ODE-20 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6 YEARS AND OLDER WHO HAVE A G551D MUTATION IN THE CFTR GENE.
- ODE-21 AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY

**PATENT AND EXCLUSIVITY TERMS**

ADB 41 of 117

**ORPHAN DRUG EXCLUSIVITY**

- ODE-22 FOR THE CONTROL OF HYPERGLYCEMIA SECONDARY TO HYPERCORTISOLISM IN ADULT PATIENTS WITH ENDOGENOUS CUSHING'S SYNDROME WHO HAVE TYPE 2 DIABETES MELLITUS OR GLUCOSE INTOLERANCE AND HAVE FAILED SURGERY OR ARE NOT CANDIDATES FOR SURGERY
- ODE-23 ADVANCED SOFT TISSUE SARCOMA (STS) WHO HAVE RECEIVED PRIOR CHEMOTHERAPY
- ODE-24 TREATMENT OF ADULTS WITH RENAL angiomyolipoma AND TUBEROUS SCLEROSIS COMPLEX (TSC) NOT REQUIRING IMMEDIATE SURGERY
- ODE-25 MANAGEMENT OF POSTHERPETIC NEURALGIA IN ADULTS.
- ODE-26 TREATMENT OF ENDOGENOUS ANTERIOR UVEITIS
- ODE-27 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- ODE-28 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE (PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES
- ODE-29 LOCALIZATION OF LYMPH NODES DRAINING A PRIMARY TUMOR IN PATIENTS WITH MELANOMA WHEN USED WITH A HAND-HELD GAMMA COUNTER
- ODE-30 TREATMENT OF ADULT PATIENTS WITH CHRONIC, ACCELERATED OR BLAST PHASE PHILADELPHIA CHROMOSOME-POSITIVE (PH+) CHRONIC MYELOGENOUS LEUKEMIA (CML) WITH RESISTANCE, OR INTOLERANCE TO PRIOR THERAPY
- ODE-31 TREATMENT OF CORNEAL CYSTINE CRYSTAL ACCUMULATION IN PATIENTS WITH CYSTINOSIS
- ODE-32 TREATMENT OF ADULT PATIENTS WITH CHRONIC OR ACCELERATED PHASE CHRONIC MYELOID LEUKEMIA (CML) WITH RESISTANCE AND/OR INTOLERANCE TO TWO OR MORE TYROSINE KINASE INHIBITORS (TKI)
- ODE-33 TREATMENT OF PROGRESSIVE, METASTATIC MEDULLARY THYROID CANCER (MTC)
- ODE-34 TREATMENT OF ADULT PATIENTS WITH CUSHING'S DISEASE FOR WHOM PITUITARY SURGERY IS NOT AN OPTION OR HAS NOT BEEN CURATIVE
- ODE-35 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ALL) THAT IS RESISTANT OR INTOLERANT TO PRIOR TYROSINE KINASE INHIBITOR THERAPY.
- ODE-36 ADJUNCT TO A LOW-FAT DIET AND OTHER LIPID-LOWERING TREATMENTS, INCLUDING LDL APHERESIS WHERE AVAILABLE, TO REDUCE LDL-C, TC, APOLIPOPROTEIN B, & NON-HDL-C IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- ODE-37 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME (SBS) WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- ODE-38 PART OF COMBINATION THERAPY IN ADULTS (GREATER THAN OR EQUAL TO 18 YEARS) WITH PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS (MDR-TB)
- ODE-39 TREATMENT OF CHRONIC IRON OVERLOAD IN PATIENTS 10 YRS. & OLDER WITH NON-TRANSFUSION DEPENDENT THALASSEMIA (NTDT) SYNDROMES AND WITH A LIVER IRON CONCENTRATION OF AT LEAST 5 MG OF IRON PER GRAM OF LIVER DRY WEIGHT & SERUM FERRITIN GREATER THAN 300 MCg/L.
- ODE-40 TREATMENT OF PEDIATRIC PATIENTS WITH NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME-POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA (PH+ ALL) IN COMBINATION WITH CHEMOTHERAPY, APPROVED UNDER NDA #21588/S-037
- ODE-41 ADJUNCT TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LDL-C, APOLIPOPROTEIN B (APO B), TOTAL CHOLESTEROL (TC), AND NON-HIGH DENSITY LIPOPROTEIN-CHOLESTEROL (NON-HDL-C) IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- ODE-42 USE AS A NITROGEN-BINDING ADJUNCTIVE THERAPY FOR CHRONIC MGMT OF ADULT AND PEDIATRIC PATIENTS AT LEAST 2 YRS WITH UREA CYCLE DISORDERS THAT CANNOT BE MANAGED BY DIETARY PROTEIN RESTRICTION AND/OR AMINO ACID SUPPLEMENTATION ALONE
- ODE-43 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING LENALIDOMIDE AND BORTEZOMIB AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY.
- ODE-44 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED, UNRESECTABLE OR METASTATIC GASTROINTESTINAL STROMAL TUMOR (GIST) WHO HAVE BEEN PREVIOUSLY TREATED WITH IMATINIB MESYLATE AND SUNITINIB MALATE.
- ODE-45 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS IN ADULTS AND CHILDREN AGES 6 YEARS AND OLDER.

**PATENT AND EXCLUSIVITY TERMS**

ADB 42 of 117

**ORPHAN DRUG EXCLUSIVITY**

- ODE-46 IMPROVEMENT OF NEUROLOGICAL OUTCOME BY REDUCING THE INCIDENCE AND SEVERITY OF ISCHEMIC DEFICITS IN ADULT PATIENTS WITH SUBARACHNOID HEMORRHAGE FROM RUPTURED INTRACRANIAL BERRY ANEURYSMS REGARDLESS OF THEIR POST-ICTUS NEUROLOGICAL CONDITION
- ODE-47 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA APPROVED TEST.
- ODE-48 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA APPROVED TEST
- ODE-49 TREATMENT OF MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- ODE-50 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST.
- ODE-51 TOPICAL TREATMENT OF STAGE 1A AND 1B MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN-DIRECTED THERAPY
- ODE-52 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS AS FIRST-LINE TREATMENT, IN COMBINATION WITH GEMCITABINE.
- ODE-53 TREATMENT OF ADULTS WITH PULMONARY ARTERIAL HYPERTENSION (PAH) WHO GROUP 1, TO IMPROVE EXERCISE CAPACITY, WHO FUNCTIONAL CLASS AND TO DELAY CLINICAL WORSENING.
- ODE-54 TX OF PAH TO DELAY DISEASE PROGRESSION. DISEASE PROGRESSION INCLUDED: DEATH, INITIATION OF IV OR SC PROSTANOIDS, OR CLINICAL WORSENING OF PAH (DECREASED 6-MINUTE WALK DISTANCE, WORSENED PAH SYMPTOMS AND NEED FOR ADDITIONAL PAH TREATMENT).
- ODE-55 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA (MCL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-56 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA (DCT) THAT IS REFRACTORY TO RADIOACTIVE IODINE TREATMENT.
- ODE-57 TRAMETINIB IN COMBO WITH DABRAFENIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-58 DABRAFENIB IN COMBO WITH TRAMETINIB FOR TX. OF PTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST. THIS INDICATION IS BASED ON THE DEMONSTRATION OF DURABLE RESPONSE RATE
- ODE-59 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER
- ODE-60 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-61 TREATMENT OF NEUROGENIC SYMPTOMATIC ORTHOSTATIC HYPOTENSION IN PATIENTS WITH PRIMARY AUTONOMIC FAILURE, DOPAMINE-BETA-HYDROXYLASE DEFICIENCY, AND NONDIABETIC AUTONOMIC NEUROPATHY
- ODE-62 TREATMENT OF PROLIFERATING INFANTILE HEMANGIOMA REQUIRING SYSTEMIC THERAPY.
- ODE-63 TREATMENT OF VISCELAR LEISHMANIASIS DUE TO LEISHMANIA DONOVANI; CUTANEOUS LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS, LEISHMANIA GUYANENSIS, AND LEISHMANIA PANAMENSIS; AND MUCOSAL LEISHMANIASIS DUE TO LEISHMANIA BRAZILIENSIS.
- ODE-64 SELECTIVE HEPATIC INTRA-ARTERIAL USE FOR IMAGING TUMORS IN ADULTS WITH KNOWN HEPATOCELLULAR CARCINOMA (HCC)
- ODE-65 TREATMENT OF PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS PART OF A COMBINATION REGIMEN.
- ODE-66 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB.
- ODE-67 GUIDING SENTINEL LYMPH NODE BIOPSY, USING A HAND-HELD GAMMA COUNTER IN PATIENTS WITH CLINICALLY NODE NEGATIVE SQUAMOUS CELL CARCINOMA OF THE ORAL CAVITY
- ODE-68 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA
- ODE-69 TREATMENT OF MALIGNANT HYPERTERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTERMIA IN PATIENTS AT HIGH

**PATENT AND EXCLUSIVITY TERMS**

ADB 43 of 117

**ORPHAN DRUG EXCLUSIVITY****RISK**

- ODE-70 RELAPSED CLL, IN COMBO. WITH RITUXIMAB, IN PATIENTS FOR WHOM RITUXIMAB ALONE WOULD BE CONSIDERED APPROPRIATE THERAPY DUE TO OTHER CO-MORBIDITIES; AND RELAPSED SLL IN PATIENTS WHO HAVE RECEIVED AT LEAST 2 PRIOR SYSTEMIC THERAPIES
- ODE-71 RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA (FL) IN PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-72 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-73 LONG-TERM TREATMENT OF ADULT PATIENTS WITH GAUCHER DISEASE TYPE 1 WHO ARE CYP2D6 EXTENSIVE METABOLIZERS (EMS), INTERMEDIATE METABOLIZERS (IMS), OR POOR METABOLIZERS (PMS) AS DETECTED BY AN FDA-CLEARED TEST.
- ODE-74 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY
- ODE-75 TREATMENT OF PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY.
- ODE-76 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE NOT RECEIVED AT LEAST 1 PRIOR THERAPY
- ODE-77 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- ODE-78 TREATMENT OF HYPERCALCEMIA IN ADULT PATIENTS WITH PRIMARY HYPERPARATHYROIDISM FOR WHOM PARATHYROIDECTOMY WOULD BE INDICATED ON THE BASIS OF SERUM CALCIUM LEVELS, BUT WHO ARE UNABLE TO UNDERGO PARATHYROIDECTOMY.
- ODE-79 TREATMENT OF PATIENTS WITH POLYCYTHEMIA VERA WHO HAVE HAD AN INADEQUATE RESPONSE TO OR ARE INTOLERANT OF HYDROXYUREA
- ODE-80 TREATMENT OF PEDIATRIC PATIENTS WITH TOURETTE'S
- ODE-81 TREATMENT OF PATIENTS WITH ACROMEGALY WHO HAVE HAD AN INADEQUATE RESPONSE TO SURGERY AND/OR FOR WHOM SURGERY IS NOT AN OPTION
- ODE-82 TREATMENT OF PATIENTS WITH UNRESECTABLE, WELL- OR MODERATELY-DIFFERENTIATED LOCALLY ADVANCED OR METASTATIC GASTROENTEROPANCREATIC NEUROENDOCRINE TUMORS TO IMPROVE PROGRESSION-FREE SURVIVAL
- ODE-83 USE OF AS MONOTHERAPY FOR PATIENTS WITH DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA MUTATED (AS DETECTED BY AN FDA-APPROVED TEST) ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY
- ODE-84 TREATMENT OF MOTOR FLUCUATIONS IN PATIENTS WITH ADVANCED PARKINSON'S DISEASE
- ODE-85 AS A REPLACEMENT SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY (CRRT) AND IN CASE OF DRUG POISONING WHEN CRRT IS USED TO REMOVE DIALZABLE SUBSTANCES
- ODE-86 TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA
- ODE-87 TREATMENT OF PATIENTS WITH LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, RADIOACTIVE IODINE REFRACTORY DIFFERENTIATED THYROID CANCER
- ODE-88 FOR USE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE TREATMENT)
- ODE-89 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST 2 PRIOR REGIMENS, INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT
- ODE-90 TREATMENT OF INVASIVE MUCORMYCOSIS IN PATIENTS 18 YEARS OF AGE AND OLDER
- ODE-91 TREATMENT OF BILE ACID SYNTHESIS DISORDERS DUE TO SINGLE ENZYME DEFECTS
- ODE-92 TREATMENT OF LYMPHANGIOLEIOMYOMATOSIS (LAM)
- ODE-93 TREATMENT OF CYSTIC FIBROSIS IN PATIENTS AGE 12 YEARS AND OLDER WHO ARE HOMOZYGOUS FOR F508DEL MUTATION IN THE CFTR GENE
- ODE-94 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY TRANSPLANT PATIENTS CONVERTED FROM TACROLIMUS IMMEDIATE-RELEASE FORMULATIONS IN COMBINATION WITH OTHER IMMUNOSUPPRESSANTS
- ODE-95 FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- ODE-96 TREATMENT OF PRIMARY HYPERKALEMIC PERIODIC PARALYSIS, PRIMARY HYPOKALEMIC PERIOD PARALYSIS, AND RELATED VARIANTS

**PATENT AND EXCLUSIVITY TERMS**

ADB 44 of 117

**ORPHAN DRUG EXCLUSIVITY**

- ODE-97 TO EXPAND THE INDICATION TO PEDIATRIC PATIENTS 2-6 YEARS OF AGE WITH NEPHROPATHIC CYSTINOSIS
- ODE-98 TREATMENT OF HEREDITARY OROTIC ACIDURIA
- ODE-99 FOR USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, FOR THE TREATMENT OF PATIENTS WITH METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED FOLLOWING GEMCITABINE-BASED THERAPY
- ODE-100 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA OR LEIOMYOSARCOMA WHO RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-101 FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATION, IN COMBINATION WITH VEMURAFENIB. COTELLIC IS NOT INDICATED FOR TREATMENT OF PATIENTS WITH WILD-TYPE BRAF MELANOMA
- ODE-102 FOR TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), AS DETECTED BY AN FDA-APPROVED TEST, WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY
- ODE-103 USE IN COMBINATION WITH LENALIDOMIDE AND DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-104 EMERGENCY TX OF PTS FOLLOWING A FU OR CAPECITABINE OD, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING TOXICITY AFFECTING THE CARDIAC SYSTEM OR CNS, AND/OR EARLY-ONSET, UNUSUALLY SEVERE AR W/IN 96 HRS FOLLOWING THE END OF FU OR CAPECITABINE ADMIN.
- ODE-105 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-106 FOR USE OF UPTRAVI (SELEXIPAG) TABLETS, 200, 400, 600, 800, 1000, 1200, 1400, AND 1600 MCG FOR TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH, WHO GROUP I) TO REDUCE THE RISKS OF DISEASE PROGRESSION AND HOSPITALIZATION FOR PAH
- ODE-107 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA WHO HAVE RECEIVED A PRIOR ANTHRACYCLINE-CONTAINING REGIMEN
- ODE-108 TREATMENT OF ADULT PATIENTS WITH PROGRESSIVE, WELL-DIFFERENTIATED, NON-FUNCTIONAL, NEUROENDOCRINE TUMORS (NET) OF GASTROINTESTINAL (GI) OR LUNG ORIGIN, (EXCLUDING PANCREATIC) WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC DISEASE
- ODE-109 INDICATED FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA WITHOUT 17P DELETION WHO HAVE NOT RECEIVED AT LEAST ONE PRIOR THERAPY (FIRST LINE THERAPY)
- ODE-110 FOR HIGH-DOSE CONDITIONING TREATMENT PRIOR TO HEMATOPOIETIC PROGENITOR (STEM) CELL TRANSPLANTATION IN PATIENTS WITH MULTIPLE MYELOMA
- ODE-111 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WHOSE TUMORS ARE ROS-1 POSITIVE.
- ODE-112 FOR THE TREATMENT OF ADULT AND PEDIATRIC PATIENTS WITH HEPATIC VENO-OCLUSIVE DISEASE (VOD), ALSO KNOWN AS SINUSOIDAL OBSTRUCTION SYNDROME (SOS), WITH RENAL OR PULMONARY DYSFUNCTION FOLLOWING HEMATOPOIETIC STEM CELL TRANSPLANTATION (HSCT).
- ODE-113 FOR TREATMENT OF PEDIATRIC AND ADULT PATIENTS WITH ACQUIRED METHEMOGLOBINEMIA.
- ODE-114 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- ODE-115 TREATMENT OF PATIENTS WITH METASTATIC, SQUAMOUS, NON-SMALL CELL LUNG CANCER PROGRESSING AFTER PLATINUM-BASED CHEMOTHERAPY
- ODE-116 TREATMENT OF PROGRESSIVE KERATOCONUS
- ODE-117 FOR TREATMENT OF PATIENTS WITH SMALL LYMPHOCYTIC LYMPHOMA (SLL)
- ODE-118 AN ADJUNCT TO DIET TO REDUCE LDL-C, TOTAL-C, NONHDL-C AND APOB IN CHILDREN AND ADOLESCENTS 7 TO 17 YEARS OF AGE WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA, EITHER ALONE OR WITH OTHER LIPID-LOWERING TREATMENTS (E.G., LDL APHERESIS)
- ODE-119 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC) IN COMBINATION WITH URSODEOXYCHOLIC ACID (UDCA) IN ADULTS WITH AN INADEQUATE RESPONSE TO UDCA, OR AS MONOTHERAPY IN ADULTS UNABLE TO TOLERATE UDCA
- ODE-120 FOR USE AFTER RADIOLABELING WITH GA 68, WITH POSITRON EMISSION TOMOGRAPHY (PET) FOR LOCALIZATION OF SOMATOSTATIN RECEPTOR POSITIVE NEUROENDOCRINE TUMORS (NETS) IN ADULT AND PEDIATRIC PATIENTS.

**PATENT AND EXCLUSIVITY TERMS**

ADB 45 of 117

**ORPHAN DRUG EXCLUSIVITY**

- ODE-121 TREATMENT OF CORNEAL ECTASIA FOLLOWING REFRACTIVE SURGERY
- ODE-122 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY (DMD) IN PATIENTS WHO HAVE A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- ODE-123 TREATMENT OF CYSTIC FIBROSIS (CF) IN PATIENTS AGE 6-11 YEAR OLD WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CFTR GENE
- ODE-124 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE FORMULATIONS, WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE, IN ADULTS WITH THE FOLLOWING SEIZURE TYPES: PARTIAL WITH COMPLEX SYMPTOMATOLOGY, GENERALIZED CLONIC-TONIC, AND MIXED
- ODE-125 INDICATED IN PEDIATRIC PATIENTS 10 YEARS AND OLDER FOR THE PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGES 3 AND 4 AND CKD STAGE 5 IN PATIENTS ON HEMODIALYSIS OR PERITONEAL DIALYSIS
- ODE-126 AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES
- ODE-127 TREATMENT OF SPINAL MUSCULAR ATROPHY IN PEDIATRIC AND ADULT PATIENTS
- ODE-128 TREATMENT OF PATIENTS WITH MARGINAL ZONE LYMPHOMA (MZL) WHO REQUIRE SYSTEMIC THERAPY AND HAVE RECEIVED AT LEAST ONE PRIOR ANTI-CD20-BASED THERAPY
- ODE-129 INDICATED FOR REDUCING THE RISK OF GRAFT REJECTION WHEN USED IN CONJUNCTION WITH HIGH-DOSE BUSULFAN & CYCLOPHOSPHAMIDE AS A PREPARATIVE REGIMEN FOR ALLOGENIC HEMATOPOIETIC PROGENITOR CELL TRANSPLANTATION FOR PEDS. PATIENTS WITH CLASS 3 BETA-THALASSEMIA
- ODE-130 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS 5 YEARS OF AGE AND OLDER
- ODE-131 TREATMENT OF MULTIPLE MYELOMA (MM), AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)
- ODE-132 TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY
- ODE-133 INDICATED FOR MAINTENANCE TREATMENT OF ADULT PATIENTS WITH RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE, OR PRIMARY PERITONEAL CANCER WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- ODE-134 TREATMENT OF CHOREA ASSOCIATED WITH HUNTINGTON'S DISEASE
- ODE-135 TREATMENT OF CHRONIC HCV GENOTYPE 2 OR 3 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS FOR USE IN COMBINATION WITH RIBAVIRIN
- ODE-136 TREATMENT OF PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER OR WEIGHING AT LEAST 35 KG WITH CHRONIC HEPATITIS C VIRUS GENOTYPE 1, 4, 5, OR 6 INFECTION WITHOUT CIRRHOSIS OR WITH COMPENSATED CIRRHOSIS
- ODE-137 TREATMENT OF OLIGOARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PERSISTENT OLIGOARTHRITIS, PSORIATIC JUVENILE IDIOPATHIC ARTHRITIS, ENTHESITIS-RELATED ARTHRITIS, OR UNDIFFERENTIATED ARTHRITIS) & POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS IN CHILDREN 0-16 YRS
- ODE-138 TREATMENT OF PEDIATRIC PATIENTS WITH ACUTE LYMPHOBLASTIC LEUKEMIA AS A COMPONENT OF A COMBINATION CHEMOTHERAPY MAINTENANCE REGIMEN
- ODE-139 TREATMENT OF PATIENTS WITH HEPATOCELLULAR CARCINOMA (HCC OR LIVER CANCER) WHO HAVE BEEN PREVIOUSLY TREATED WITH THE DRUG SORAFENIB.
- ODE-140 TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)
- ODE-141 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) THAT IS FLT3 MUTATION-POSITIVE AS DETECTED BY AN FDA APPROVED TEST, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION
- ODE-142 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- ODE-143 TO DECREASE THE RECURRENCE OF PNEUMOTHORAX IN ADULTS
- ODE-144 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)
- ODE-145 TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE

**PATENT AND EXCLUSIVITY TERMS**

ADB 46 of 117

**ORPHAN DRUG EXCLUSIVITY**

TUMORS ARE ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE AS DETECTED BY AN FDA-APPROVED TEST

- ODE-146 OPTICAL IMAGING AGENT INDICATED IN PATIENTS WITH GLIOMA (SUSPECTED WORLD HEALTH ORGANIZATION GRADES III OR IV ON PREOPERATIVE IMAGING) AS AN ADJUNCT FOR THE VISUALIZATION OF MALIGNANT TISSUE DURING SURGERY
- ODE-147 DABRAFENIB IN COMBINATION WITH TRAMETINIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-148 TRAMETINIB IN COMBINATION WITH DABRAFENIB, FOR THE TX. OF PTS WITH METASTATIC NON-SMALL CELL LUNG CANCER WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-149 TREATMENT OF PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER WITH CENTRAL PRECOCIOUS PUBERTY
- ODE-150 TO REDUCE THE ACUTE COMPLICATIONS OF SICKLE CELL DISEASE IN ADULT AND PEDIATRIC PATIENTS 5 YEARS OF AGE AND OLDER.
- ODE-151 TREATMENT OF ADULT PATIENTS WITH RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- ODE-152 TREATMENT OF ADULT PATIENTS WITH CHRONIC GRAFT VERSUS HOST DISEASE (CGVHD)
- ODE-153 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY WITH OR WITHOUT CONCOMITANT DOPMINERGIC MEDICATIONS
- ODE-154 FOR USE IN CHILDREN AGES 2 TO 12 YEARS OLD WITH CHAGAS DISEASE
- ODE-155 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- ODE-156 TREATMENT OF ADULTS WITH CARCINOID SYNDROME; WHEN USED, IT REDUCES THE FREQUENCY OF SHORT-ACTING SOMATOSTATTIN ANALOG RESCUE THERAPY

**PATENT USE**

- U-1 PREVENTION OF PREGNANCY
- U-2 TREATMENT OR PROPHYLAXIS OF ANGINA PECTORIS AND ARRHYTHMIA
- U-3 TREATMENT OF HYPERTENSION
- U-4 PROVIDING PREVENTION AND TREATMENT OF EMESIS AND NAUSEA IN MAMMALS
- U-5 METHOD OF PRODUCING BRONCHODILATION
- U-6 METHOD OF PRODUCING SYMPATHOMIMETIC EFFECTS
- U-7 INCREASING CARDIAC CONTRACTILITY
- U-8 ACUTE MYOCARDIAL INFARCTION
- U-9 CONTROL OF EMESIS ASSOCIATED WITH ANY CANCER CHEMOTHERAPY AGENT
- U-10 DIAGNOSTIC METHOD FOR DISTINGUISHING BETWEEN HYPOTHALAMIC MALFUNCTIONS OR LESIONS IN HUMANS
- U-11 TREATMENT OR PROPHYLAXIS OF CARDIAC DISORDERS
- U-12 METHOD OF TREATING [A] HUMAN SUFFERING FROM DEPRESSION
- U-13 A METHOD FOR TREATING ANXIETY IN A HUMAN SUBJECT IN NEED OF SUCH TREATMENT
- U-14 ADJUNCTIVE THERAPY FOR THE PREVENTION AND TREATMENT OF HYPERAMMONEMIA IN THE CHRONIC MANAGEMENT OF PATIENTS WITH UREA CYCLE ENZYMOGRAPHIES
- U-15 METHOD OF LOWERING INTRAOOCULAR PRESSURE
- U-16 USE IN LUNG SCANNING PROCEDURES
- U-17 TREATMENT OF VENTRICULAR AND SUPRAVENTRICULAR ARRHYTHMIAS
- U-18 METHOD FOR INHIBITING GASTRIC SECRETION IN MAMMALS
- U-19 TREATMENT OF INFLAMMATION
- U-20 A PROCESS FOR TREATING A PATIENT SUFFERING FROM PARKINSON'S SYNDROME AND IN NEED OF TREATMENT
- U-21 TREATMENT OF HUMANS SUFFERING UNDESIRED UROTOXIC SIDE EFFECTS CAUSED BY

**PATENT AND EXCLUSIVITY TERMS**

ADB 47 of 117

**PATENT USE**

## CYTOSTATICALLY ACTIVE ALKYLATING AGENTS

- U-22 METHOD OF COMBATTING PATHOLOGICALLY REDUCED CEREBRAL FUNCTIONS AND PERFORMANCE WEAKNESSES, CEREBRAL INSUFFICIENCY AND DISORDERS IN CEREBRAL CIRCULATION AND METABOLISM IN WARM-BLOODED ANIMALS
- U-23 METHOD FOR TREATING PROSTATIC CARCINOMA COMPRISING ADMINISTERING FLUTAMIDE
- U-24 METHOD FOR TREATING PROSTATE ADENOCARCINOMA COMPRISING ADMINISTERING AN ANTIANDROGEN INCLUDING FLUTAMIDE AND AN LHRH AGONIST
- U-25 REDUCING CHOLESTEROL IN CHOLELITHIASIS PATIENTS
- U-26 REDUCING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-27 DISSOLVING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCELAR AND RENAL ARTERIOGRAPHY, AORTOGRAPHY AND LEFT VENTRICULOGRAPHY
- U-29 CT IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY
- U-30 CEREBRAL ANGIOGRAPHY, AND VENOGRAPHY
- U-31 INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY
- U-32 PALLIATIVE TREATMENT OF PATIENTS WITH OVARIAN CARCINOMA RECURRENT AFTER PRIOR CHEMOTHERAPY, INCLUDING PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED WITH CISPLATIN
- U-33 TREATING VIRAL INFECTIONS IN A MAMMAL
- U-34 TREATING VIRAL INFECTIONS IN A WARM-BLOODED ANIMAL
- U-35 TREATING CYTOMEGALOVIRUS IN A HUMAN WITH AN INJECTABLE COMPOSITION
- U-36 METHODS OF TREATING BACTERIAL ILLNESSES
- U-37 METHOD OF TREATING GASTROINTESTINAL DISEASE
- U-38 TREATMENT OF PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA
- U-39 ANGINA PECTORIS
- U-40 METHOD OF TREATMENT OF BURNS
- U-41 METHOD OF TREATING CARDIAC ARRHYTHMIAS
- U-42 ADJUVANT TREATMENT IN COMBINATION WITH FLUOROURACIL AFTER SURGICAL RESECTION IN PATIENTS WITH DUKE'S STAGE C COLON CANCER
- U-43 MANAGEMENT OF CHRONIC PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA
- U-44 RELIEF OF NAUSEA AND VOMITING
- U-45 TREATMENT OF INFLAMMATION AND ANALGESIA
- U-46 TREATMENT OF PANIC DISORDER
- U-47 STIMULATION OF THE RELEASE OF GROWTH HORMONE
- U-48 ANALGESIA
- U-49 SYMPTOMATIC CANCER-RELATED HYPERCALCEMIA
- U-50 USE IN TREATING INFLAMMATORY DERMATOSES
- U-51 BLOOD POOL IMAGING, INCLUDING CARDIAC FIRST PASS AND GATED EQUILIBRIUM IMAGING AND FOR DETECTION OF SITES OF GASTROINTESTINAL BLEEDING
- U-52 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER SIX MONTHS OF AGE) WITH ADVANCED HIV INFECTION
- U-53 HYPERCALCEMIA OF MALIGNANCY
- U-54 REVERSAL AGENT OR ANTAGONIST OF NONDEPOLARIZING NEUROMUSCULAR BLOCKING AGENTS
- U-55 TREATMENT OF PAIN
- U-56 AID TO SMOKING CESSION
- U-57 OPHTHALMIC USE OF NORFLOXACIN
- U-58 METHOD OF TREATING INFLAMMATORY INTESTINAL DISEASES
- U-59 METHOD OF TREATING HYPERCHOLESTEROLEMIA
- U-60 NASAL ADMINISTRATION OF BUTORPHANOL
- U-61 CEREBRAL AND PERIPHERAL ARTERIOGRAPHY AND CT IMAGING OF THE HEAD

**PATENT AND EXCLUSIVITY TERMS**

ADB 48 of 117

**PATENT USE**

- U-62 CORONARY ARTERIOGRAPHY, LEFT VENTRICULOGRAPHY, CT IMAGING OF THE BODY, INTRAVENOUS EXCRETORY UROGRAPHY, INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY AND VENOGRAPHY
- U-63 ISOPRENALINE ANTAGONISM ON THE HEART RATE OR BLOOD PRESSURE
- U-64 TREATMENT OF VIRAL INFECTIONS
- U-65 METHOD OF TREATMENT OF A PATIENT INFECTED WITH HIV
- U-66 TRIPHASIC REGIMEN
- U-67 METHOD OF INDUCING ANESTHESIA IN A WARM BLOODED ANIMAL
- U-68 TREATMENT OF ACTINIC KERATOSIS
- U-69 TREATMENT OF PNEUMOCYSTIS CARINII INFECTIONS
- U-70 TREATMENT OF TRANSIENT INSOMNIA
- U-71 METHOD OF TREATMENT OF HEART FAILURE
- U-72 TREATMENT OF MIGRAINE
- U-73 METHOD OF TREATING DISEASES OR INFECTIONS CAUSED BY MYCETES
- U-74 METHOD OF PROVIDING HYPNOTIC EFFECT
- U-75 RELIEF OF OCULAR ITCHING DUE TO SEASONAL ALLERGIC CONJUNCTIVITIS
- U-76 USE TO IMAGE A SUBJECT WITH A MAGNETIC RESONANCE IMAGING SYSTEM
- U-77 TREATMENT OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS
- U-78 ULCERATIVE COLITIS
- U-79 SYMPTOMATIC TREATMENT OF PATIENTS WITH NOCTURNAL HEARTBURN DUE TO GERD
- U-80 METHOD OF TREATING OCULAR BACTERIAL INFECTIONS
- U-81 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS
- U-82 TREATMENT FOR DEMENTIA IN PATIENTS WITH ALZHEIMER'S DISEASE
- U-83 TREATMENT OF SEIZURES
- U-84 A METHOD OF BLOCKING THE UPTAKE OF MONOAMINES BY BRAIN NEURONS IN ANIMALS
- U-85 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-86 METHOD OF TREATING CERTAIN FORMS OF EPILEPSY
- U-87 METHOD FOR NONINVASIVE ADMINISTRATION OF SEDATIVES, ANALGESICS, AND ANESTHETICS
- U-88 TREATMENT OF MODERATE PLAQUE PSORIASIS
- U-89 TREATMENT OR PROPHYLAXIS OF EMESIS
- U-90 TREATMENT OF PSYCHOTIC DISORDERS
- U-91 ALTERNATIVE THERAPY TO TRIMETHOPRIM-SULFAMETHOXAZOLE FOR TREATMENT OF MODERATE-TO-SEVERE PNEUMOCYSTIS CARINII PNEUMONIA IN IMMUNOCOMPROMISED AND AIDS PATIENTS
- U-92 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN DEPENDENT DIABETES MELLITUS AND RETINOPATHY
- U-93 USE AS AN ANTIHISTAMINE/DECONGESTANT
- U-94 TREATMENT-ADULTS W/ ADVANCED HIV, INTOLERANT OF APPROVED THERAPIES, INTOLERANT OF APPROVED THERAPIES W/PROVEN BENEFIT OR HAVE EXPERIENCED CLINICAL/IMMUNOLOGICAL DETERIORATION WHILE RECEIVING..OR FOR WHOM SUCH THERAPIES-CONTRAINDED
- U-95 SHORT TERM MANAGEMENT OF MODERATE PRURITIS IN ADULTS WITH ATOPIC DERMATITIS AND LICHEN SIMPLEX CHRONICUS
- U-96 METHOD OF TREATING VARICELLA ZOSTER (SHINGLES) INFECTIONS
- U-97 A METHOD OF TREATING A PATIENT IN NEED OF MEMORY ENHANCEMENT
- U-98 A METHOD OF INDUCING REGRESSION OF LEUKEMIA CELL GROWTH IN A MAMMAL
- U-99 METHOD OF PROVIDING POTASSIUM TO A SUBJECT IN NEED OF POTASSIUM
- U-100 METHOD OF TREATING OCULAR INFLAMMATION
- U-101 ADJUNCT TO CONVENTIONAL CT OR MRI IMAGING IN THE LOCALIZATION OF STROKE IN PATIENTS IN WHOM STROKE HAS ALREADY BEEN DIAGNOSED
- U-102 METHOD OF HORMONALLY TREATING MENOPAUSAL OR POST-MENOPAUSAL DISORDERS IN WOMEN

**PATENT AND EXCLUSIVITY TERMS**

ADB 49 of 117

**PATENT USE**

- U-103 TREATMENT OF OCULAR HYPERTENSION
- U-104 TREATMENT OF AQUEOUS HUMOR FORMATION AND INTRAOCULAR PRESSURE
- U-105 EMESIS
- U-106 TREATMENT OF EPILEPSY
- U-107 TREATMENT OF HYPERTENSION AND ANGINA PECTORIS
- U-108 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER, GASTROESOPHAGEAL REFLUX DISEASE (GERD), SEVERE EROSIVE ESOPHAGITIS, POORLY RESPONSIVE SYMPTOMATIC GERD AND PATHOLOGICAL HYPERSECRETOORY CONDITIONS AND MAINTENANCE HEALING OF EROSION ESOPHAGITIS
- U-109 ADJUNCT DIET IN THE TX OF ELEVATED TOTAL CHOLESTEROL AND LDL-C LEVELS IN PTS W/PRIMARY HYPERCHOLESTEROLEMIA WHOSE RESPONSE TO DIETARY RESTRICTION OF SAT FAT AND CHOLESTEROL AND OTHER NONPHARMACOLOGICAL MEASURES HAS NOT BEEN ADEQUATE
- U-110 USE AS A RETRIEVEABLE PESSARY
- U-111 DIABETES
- U-112 CONTRACEPTION
- U-113 METHOD OF CONDUCTING RADIOLOGICAL EXAMINATION OF A PATIENT BY ADMINISTERING TO SAID PATIENT A RADIOPAQUE AMOUNT OF IOPROMIDE
- U-114 USE FOR INHIBITING BONE RESORPTION
- U-115 USE OF VASODILATORS TO EFFECT AND ENHANCE AN ERECTION (AND THUS TREAT ERECTILE DYSFUNCTION), BY INJECTION INTO THE PENIS
- U-116 METHOD OF MYOCARDIAL IMAGING
- U-117 TREATMENT OF OCULAR ALLERGIC RESPONSE IN HUMAN EYES
- U-118 METHOD OF LOWERING BLOOD SUGAR LEVEL
- U-119 TREATMENT OF NASAL HYPERSECRETION
- U-120 CONTROLLING OR PREVENTING POST-OPERATIVE INTRAOCULAR PRESSURE RISES ASSOCIATED WITH OPHTHALMIC LASER SURGICAL PROCEDURES
- U-121 METHOD OF TREATING CONDITIONS MEDIATED THROUGH HISTAMINE H<sub>2</sub>-RECEPTORS
- U-122 A THERAPEUTIC METHOD FOR CONTROLLING THROMBOSIS
- U-123 METHOD FOR CONTROLLING THROMBOSIS AND DECREASING BLOOD HYPERCOAGULATION AND HEMORRHAGING RISKS
- U-124 TREATMENT OF ACNE
- U-125 TREATMENT NEUROGENERATIVE DISEASES
- U-126 TREATMENT OF GASTRITIS
- U-127 METHOD OF PRODUCING NEUROMUSCULAR BLOCKADE
- U-128 METHOD FOR TREATMENT OF TUMORS
- U-129 METHOD TO DESTROY OR IMPAIR TARGET CELLS
- U-130 MANAGEMENT OF PATIENTS WITH MASTOCYTOSIS
- U-131 PHOTODAMAGED SKIN
- U-132 INHIBITING HIV PROTEASE
- U-133 MANAGEMENT OF OBESITY INCLUDING WEIGHT LOSS AND MAINTENANCE IN PATIENTS ON A REDUCED-CALORIE DIET
- U-134 TREATMENT OF ACNE VULGARIS
- U-135 ANTITUMOR AGENT
- U-136 PROCESS FOR WASTE NITROGEN REMOVAL
- U-137 METHOD OF TREATING BACTERIAL VAGINOSIS
- U-138 TREATMENT OF ALLERGIC RHINITIS
- U-139 TREATMENT OF ALLERGIC REACTIONS
- U-140 USE OF NORVIR TO INHIBIT HIV PROTEASE OR TO INHIBIT AN HIV INFECTION
- U-141 TREATMENT OF ULCERATIVE COLITIS

**PATENT AND EXCLUSIVITY TERMS**

ADB 50 of 117

**PATENT USE**

- U-142 METHOD OF TREATING ALLERGIC REACTIONS IN A MAMMAL BY USING THIS ACTIVE METABOLITE
- U-143 BIODEGRADABLE SUPERPARAMAGNETIC METAL OXIDES AS CONTRAST AGENTS FOR MR IMAGING
- U-144 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC MATERIALS FOR USE IN CLINICAL APPLICATIONS
- U-145 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC PARTICLES FOR USE AS NUCLEAR MAGNETIC RESONANCE IMAGING AGENTS
- U-146 METHOD OF TREATING SUSCEPTIBLE NEOPLASMS IN MAMMALS
- U-147 DETECTION OF GASTROINTESTINAL DISORDERS AND THE SUBSEQUENT BREATH COLLECTION AND MEASUREMENT OF  $^{13}\text{CO}_2$
- U-148 DEVICE FOR COLLECTING A BREATH SAMPLE
- U-149 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS, ACUTE MANIA OR MILD ANXIETY STATES
- U-150 METHOD OF USE FOR CONTROLLING HYPERGLYCEMIA BY ADMINISTRATION OF THIS SUSTAINED RELEASE DOSAGE FORM OF GLIPIZIDE
- U-151 RELIEF OF SYMPTOMS OF THE COMMON COLD
- U-152 METHOD OF TREATING ANXIETY RELATED DISORDERS INCLUDING OBSESSIVE COMPULSIVE DISORDER
- U-153 TREATMENT OF INITIAL EPISODE GENITAL HERPES
- U-154 METHOD OF TREATING ANIMALS SUFFERING FROM AN APPETITE DISORDER
- U-155 TREATMENT OF ERECTILE DYSFUNCTION
- U-156 METHOD OF PROVIDING ANESTHESIA
- U-157 TREATMENT OF A HUMAN SUFFERING FROM VITAMIN B12 DEFICIENCY
- U-158 ANGINA
- U-159 TREATMENT OF INTERSTITIAL CYSTITIS
- U-160 TREATMENT OF BACTERIAL INFECTIOUS DISEASE
- U-161 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS IN A PATIENT
- U-162 METHOD OF USE TO INHIBIT CHOLESTEROL SYNTHESIS IN A HUMAN SUFFERING FROM HYPERCHOLESTEROLEMIA
- U-163 METHOD OF USING TROGLITAZONE TO TREAT IMPAIRED GLUCOSE TOLERANCE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS
- U-164 METHOD OF USING TROGLITAZONE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS IN A DEFINED POPULATION OF PATIENTS
- U-165 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA
- U-166 TREATMENT OF H. PYLORI-ASSOCIATED DUODENAL ULCER
- U-167 METHOD FOR TREATING HIV-1 INFECTION
- U-168 METHOD OF INHIBITING LIPOXYGENASE ACTIVITY IN A MAMMAL WHICH IS THE MODE OF ACTION IN THE TREATMENT OF ASTHMA
- U-169 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS A CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING
- U-170 METHOD OF OBTAINING AN MR IMAGE USING THE COMPOSITION/DRUG PRODUCT AS A CONTRAST AGENT
- U-171 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS AN ORAL CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING OF THE GASTROINTESTINAL TRACT
- U-172 TREATMENT OF GENITAL WARTS
- U-173 ADMINISTRATION TO A HOST SUFFERING FROM GESTATIONAL DIABETES
- U-174 USE AS AN ANTIHISTAMINE AGENT
- U-175 METHOD OF TREATING MALIGNANT TUMORS
- U-176 METHOD OF TREATING A PATIENT SUFFERING FROM LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES
- U-177 FUNGICIDE
- U-178 FACILITATED ADHERENCE OF AGENTS TO SKIN

**PATENT AND EXCLUSIVITY TERMS**

ADB 51 of 117

**PATENT USE**

- U-179 ENHANCED CUTANEOUS PENETRATION OF A DERMALLY-APPLIED PHARMACOLOGICALLY ACTIVE AGENT
- U-180 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER 6 MONTHS OF AGE) WITH ADVANCED HIV INFECTION
- U-181 PRODUCING ALPHA ADRENERGIC ANTAGONISTIC ACTION IN A HOST
- U-182 USE OF SALMETEROL IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
- U-183 TREATMENT OF CONDITIONS CAUSED BY DISTURBANCE OF NEURONAL 5HT FUNCTION
- U-184 TREATING ALLERGIC EYE DISEASES IN HUMANS
- U-185 METHOD OF TREATING HYPERTENSION
- U-186 METHOD FOR TREATING GI DISORDERS CAUSED BY H. PYLORI WHICH COMPRISES ADMINISTRATION OF RANITIDINE BISMUTH CITRATE AND CLARITHROMYCIN FOR A GREATER THAN ADDITIVE EFFECT
- U-187 THERAPEUTIC TREATMENT OF CALCIFIC TUMORS
- U-188 TREATMENT OF H. PYLORI ASSOCIATED DUODENAL ULCER
- U-189 ENHANCEMENT OF THE BIOAVAILABILITY OF THE DRUG SUBSTANCE
- U-190 USE OF RITONAVIR IN COMBINATION WITH ANY REVERSE TRANSCRIPTASE INHIBITOR
- U-191 METHOD OF TREATMENT FOR CONTROLLING AND LOWERING INTRAOCULAR PRESSURE IN A HUMAN
- U-192 USE IN TREATING ALLERGIC REACTIONS
- U-193 PSORIASIS
- U-194 TREATING ANGINA PECTORIS AND HIGH BLOOD PRESSURE
- U-195 METHOD FOR THE DIAGNOSIS OF GASTROINTESTINAL DISORDERS BY UREA ISOTOPE OR NITROGEN LABELED CARBON
- U-196 TREATMENT OF METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH ESTROGEN RECEPTOR POSITIVE TUMORS
- U-197 USE IN COMBINATION WITH CERTAIN LHRH ANALOGUES FOR THE TREATMENT OF ADVANCED PROSTATE CANCER
- U-198 TREATMENT METASTATIC CARCINOMA OF OVARY AFTER 1ST LINE FAILURE OR SUBSEQUENT CHEMOTHERAPY, TREATMENT OF BREAST CANCER AFTER FAILURE OF COMBINATION CHEMOTHERAPY FOR METASTATIC DISEASE AND 2ND LINE TREATMENT OF AIDS RELATED KAPOSI'S SARCOMA
- U-199 METHOD OF TREATING INFECTIOUS UPPER GI TRACT DISORDERS CAUSED BY CAMPYLOBACTER PYLORIDIS INFECTION COMPRISING ADMINISTRATION OF A BISMUTH AGENT AND AN ANTIMICROBIAL AGENT
- U-200 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF A BISMUTH-CONTAINING AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
- U-201 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF CAMPYLOBACTER-INHIBITING ANTIMICROBIAL AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
- U-202 METHOD OF TREATING PEPTIC ULCER DISEASE CAUSED BY CAMPYLOBACTER PYLORIDIS COMPRISING ORAL ADMINISTRATION OF 50 TO 5,000MG BISMUTH DAILY FOR 3-56 DAYS
- U-203 TREATMENT OF ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-204 USE OF TAXOL IN COMBINATION WITH G-CSF FOR TREATMENT OF PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-205 METHOD FOR TREATING HEARTBURN
- U-206 METHOD OF USING FSH ALONE, WITHOUT THE PRESENCE OF EXOGENEOUS LH, IN IN VITRO FERTILIZATION
- U-207 USE AS NASAL SPRAY
- U-208 VAGINAL ADMINISTRATION USING SPECIFIED FORMULATION
- U-209 VAGINAL ADMINISTRATION OF PROGESTERONE USING SPECIFIED FORMULATION
- U-210 METHOD OF TREATING CONGESTIVE HEART FAILURE
- U-211 USE IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
- U-212 METHOD OF TREATMENT OF PARKINSON'S DISEASE
- U-213 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS AND TREATING HYPERCHOLESTEROLEMIA

**PATENT AND EXCLUSIVITY TERMS**

ADB 52 of 117

**PATENT USE**

- AND METHOD FOR TREATING HYPERLIPIDEMIA  
U-214 USE AS A BLOOD GLUCOSE-LOWERING AGENT  
U-215 TREATMENT OF EPILEPSY TWICE DAILY. TREATING A PATIENT BY ADMINISTERING CARBAMAZEPINE IN A DOSAGE FORM CAPABLE OF MAINTAINING BLOOD CONCENTRATION FROM 4-12MCG/ML OVER 12 HOURS  
U-216 TREATMENT OF ADENOCARCINOMA, INCLUDING STAGE B2-C BY ADMINISTERING AN AGONIST OF LH-RH AND FLUTAMIDE  
U-217 METHOD OF PRODUCING ANESTHESIA  
U-218 METHOD FOR LIMITING THE POTENTIAL FOR MICROBIAL GROWTH IN THE DRUG PRODUCT  
U-219 TREATMENT OF PARKINSON'S DISEASE  
U-220 METHOD OF DIAGNOSIS  
U-221 SELECTIVE VASODILATION BY CONTINUOUS ADENOSINE INFUSION  
U-222 METHOD OF TREATING PAGET'S DISEASE USING ACTONEL  
U-223 TREATMENT OF BACTERIAL CONJUNCTIVITIS CAUSED BY SUSCEPTIBLE STRAINS OF MICROORGANISMS  
U-224 CONTROLLING INTRAOCULAR PRESSURE  
U-225 METHOD FOR DELIVERY  
U-226 METHOD OF ENHANCING THE DISSOLUTION PROFILE OF A PHARMACEUTICAL FROM A SOLID DOSAGE FORM CONTAINING THE PHARMACEUTICAL AND SIMETHICONE  
U-227 NASAL ADMINISTRATION  
U-228 ASTHMA  
U-229 CARDIAC INSUFFICIENCY (CONGESTIVE HEART FAILURE)  
U-230 PREVENTION OF ACUTE CARDIAC ISCHEMIC EVENTS  
U-231 USE IN PARKINSON'S DISEASE  
U-232 METHOD OF TREATING MIGRAINE  
U-233 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE  
U-234 METHOD OF USING RIBAVIRIN TO TREAT VIRAL INFECTIONS IN MAMMALS  
U-235 METHOD OF MODULATING TH1 AND TH2 RESPONSE IN ACTIVATED T CELLS OF A HUMAN COMPRISING ADMINISTERING RIBAVIRIN TO THE T CELLS IN A DOSAGE WHICH PROMOTES THE TH1 RESPONSE AND SUPPRESSES THE TH2 RESPONSE  
U-236 TREATING MALE PATTERN BALDNESS WITH 0.05 TO 3.0MG/DAY  
U-237 METHOD OF PERFORMING NMR IMAGING WITH A PATIENT COMPRISING ADMINISTERING TO THE PATIENT AN EFFECTIVE AMOUNT OF CONTRAST AGENT DISCLOSED IN THE CLAIMS  
U-238 IMAGING A BODY TISSUE AND SUBJECTING TO NMR TOMOGRAPHY, ADMINISTERING AN AMOUNT OF PHARMACEUTICAL AGENT FOR AFFECTING THE RELAXATION TIMES OF ATOMS IN BODY TISSUES UNDERGOING NMR DIAGNOSIS, WHEREBY THE IMAGE CONTRAST IS ENHANCED....  
U-239 TREATING OR CONTROLLING OCULAR INFLAMMATION WHICH COMPRISES TOPICALLY ADMINISTERING TO AFFECTED EYE A COMPOSITION COMPRISING AN NSAID, A POLYMERIC QUATERNARY AMMONIUM COMPOUND AND BORIC ACID  
U-240 TREATMENT OF ACUTE MIGRAINE ATTACKS  
U-241 FOR SHORT-TERM TREATMENT ACTIVE DUODENAL ULCER, MAINTENANCE THERAPY FOR DUODENAL ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING OF ACTIVE ULCER, SHORT-TERM TREATMENT ACTIVE BENIGN GASTRIC ULCER & GERD, PATHOLOGICAL HYPERSECRETOORY CONDITIONS  
U-242 USE OF FOLLITROPIN ALPHA ALONE IN IN-VITRO FERTILIZATION  
U-243 TOPICAL ADMINISTRATION  
U-244 PLATELET AGGREGATION INHIBITORS  
U-245 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS  
U-246 PHOSPHATE BINDING  
U-247 TREATMENT OF RHEUMATOID ARTHRITIS  
U-248 TREATMENT OF HIV

**PATENT AND EXCLUSIVITY TERMS**

ADB 53 of 117

**PATENT USE**

- U-249 METHOD OF TREATING ALLERGIC OR NON-ALLERGIC RHINITIS IN PATIENTS BY ADMINISTERING AEROSOLIZED PARTICLES OF MOMETASONE FUROATE
- U-250 TREATMENT OF HEPATITIS B INFECTION
- U-251 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS IN THE TREATMENT OF TYPE II DIABETES
- U-252 METHOD OF TREATING A HUMAN SUBJECT HAVING GAUCHER'S DISEASE
- U-253 ORAL TRANSMUCOSAL USE
- U-254 USE OF AGGRASTAT IN COMBINATION WITH HEPARIN
- U-255 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY
- U-256 TREATMENT OF HIV INFECTION IN COMBINATION WITH ONE OR MORE ADDITIONAL HIV ANTIVIRAL AGENTS
- U-257 TREATMENT OF HIV INFECTION
- U-258 TREATMENT OF NEURODEGENERATIVE DISEASES
- U-259 TREATMENT OF ANDROGENIC ALOPECIA BY ORAL ADMINISTRATION DRUG SUBSTANCE
- U-260 REDUCTION OF INTRAOCCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA AND OCULAR HYPERTENSION WHO ARE INTOLERANT OF OTHER IOP LOWERING MEDICATIONS OR INSUFFICIENTLY RESPONSIVE TO ANOTHER IOP LOWERING MEDICATION
- U-261 TREATING BENIGN PROSTATIC HYPERPLASIA WITH A GENUS OF COMPOUNDS, INCLUDING FINASTERIDE
- U-262 TREATING BENIGN PROSTATIC HYPERTROPHY WITH FINASTERIDE
- U-263 METHOD OF TREATING A MALIGNANT CONDITION THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING LEUKEMIA OR LYMPHOMA IN A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVENOUS ADMINISTRATION OF BUSULFAN
- U-264 METHOD OF TREATING A MALIGNANT DISEASE THROUGH PARENTERAL ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN
- U-265 USE AS LAXATIVE
- U-266 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS; RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS; MANAGEMENT OF ACUTE PAIN IN ADULTS; TREATMENT OF PRIMARY DYSMENORRHEA; ACUTE TREATMENT OF MIGRAINE ATTACKS IN ADULTS
- U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMIN TO PT, 30 MIN PRIOR TO CONSUMPTION BY THE PT THE FOOD/BEVERAGE, A COMPOSITION COMPRISING 10MG FAMOTIDINE
- U-268 ACROMEGALY
- U-269 EXCESS GH-SECRETION OR GASTRO-INTESTINAL DISORDERS
- U-270 METHOD OF IMPROVING THE TIME FOR ADMINISTRATION OR THE TIME BETWEEN CHANGES OF GIVING SETS FOR THE DRUG PRODUCT
- U-271 METHOD OF TREATING TUMORS
- U-272 METHOD OF TREATING CARCINOMA
- U-273 CUTANEOUS T-CELL LYMPHOMA
- U-274 ZANAMIVIR FOR INHALATION
- U-275 METHOD OF USE OF THE DRUG SUBSTANCE
- U-276 METHOD OF USE OF LEVOBUPIVACAINE
- U-277 NEUROLOGICAL AND OTHER DISORDERS (TREATMENT OF EPILEPSY, BID ORAL DOSING)
- U-278 METHOD OF USE OF THE INDICATION OF THE DRUG PRODUCT
- U-279 METHOD OF USE OF THE APPROVED PRODUCT
- U-280 TREATING PRECIPITATED ACUTE URINARY RETENTION WITH FINASTERIDE
- U-281 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS
- U-282 METHOD OF TREATING BACTERIAL INFECTIONS
- U-283 METHOD FOR TREATING MENOPAUSAL SYMPTOMS IN A POSTMENOPAUSAL FEMALE
- U-284 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED

**PATENT AND EXCLUSIVITY TERMS**

ADB 54 of 117

**PATENT USE**

- WITH MENOPAUSE, AND VULVAR AND VAGINAL ATROPHY) AND OSTEOPOROSIS  
U-285 DEPRESSION AND SOCIAL ANXIETY DISORDER/SOCIAL PHOBIA
- U-286 DEPRESSION
- U-287 TREATMENT OR PREVENTION OF OSTEOPOROSIS
- U-288 THERAPY OF INFLUENZA
- U-289 TREATMENT OF NON-HYPERKERATOTIC ACTINIC KERATOSES OF FACE AND SCALP
- U-290 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS)
- U-291 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH CYCLOSPORIN
- U-292 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH AZATHIOPRINE
- U-293 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH A CORTICOSTEROID
- U-294 TREATMENT OF HYPERPIGMENTARY DISORDERS
- U-295 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-296 TREATING MIGRAINE PAIN AND ONE OR MORE OF A CLUSTER OF SYMPTOMS CHARACTERISTIC OF A MIGRAINE ATTACK SYMPTOMS BEING SELECTED FROM PHOTOPHOBIA, PHONOPHOBIA NAUSEA AND FUNCTIONAL DISABILITY
- U-297 PREVENTION OR TREATMENT OF REVERSIBLE VASOCONSTRICTION BY THE INHALATION OF NITRIC OXIDE WITH AN OXYGEN CONTAINING GAS
- U-298 METHOD OF COMBATING BACTERIA IN A PATIENT
- U-299 TREATMENT OF ADENOMATOUS POLYPYS
- U-300 INDICATED FOR THE REDUCTION OF ELEVATED TOTAL AND LDL CHOLESTEROL LEVELS IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA
- U-301 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS AND BIGUANIDES IN THE TREATMENT OF TYPE II DIABETES
- U-302 TO REDUCE THE RISK OF STROKE IN PATIENTS WHO HAVE HAD TRANSIENT ISCHEMIA OF THE BRAIN OR COMPLETED ISCHEMIC STROKE DUE TO THROMBOSIS
- U-303 METHOD OF USE PATENT-PRODUCT APPROVED FOR TREATMENT OF OSTEOPOROSIS, PAGET'S DISEASE, PREVENTION AND TREATMENT OF GLUCOCORTICOID INDUCED OSTEOPOROSIS
- U-304 A METHOD OF TREATMENT OF A CONDITION INVOLVING AN ANTIBODY ANTIGEN REACTION
- U-305 METHODS FOR USING THE DRUG PRODUCT
- U-306 TREATMENT OF POST-MENOPAUSAL UROGENITAL SYMPTOMS ASSOCIATED WITH ESTROGEN DEFICIENCY
- U-307 CLAIMS AN OLANZAPINE POLYMORPH USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATION OF THIS NDA
- U-308 CLAIMS A SOLID ORAL FORMULATION INCLUDING TABLETS AND GRANULES OF OLANZAPINE USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATIONS OF THIS NDA
- U-309 TREATING SJOEGREN SYNDROME
- U-310 TREATMENT OF XEROSTOMIA
- U-311 HORMONE REPLACEMENT
- U-312 PANIC DISORDER, OBSESSIVE-COMPULSIVE DISORDER, POSTTRAUMATIC STRESS DISORDER
- U-313 TREATMENT OF CONGESTIVE HEART FAILURE
- U-314 METHOD FOR TREATING HYPERPARATHYROIDISM WHICH COMPRISSES SUPPRESSING PARATHYROID ACTIVITY
- U-315 METHOD FOR ADMINISTERING DRUG TO GASTROINTESTINAL TRACT
- U-316 METHOD OF TREATING A SUBJECT SUFFERING FROM PROSTATE CANCER
- U-317 METHOD OF USING TROGLITAZONE TO TREAT PATIENTS HAVING INSULIN RESISTANCE
- U-318 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE

**PATENT AND EXCLUSIVITY TERMS**

ADB 55 of 117

**PATENT USE**

- U-319 TREATMENT OF MICROBIAL INFECTIONS
- U-320 INHIBITING OR ELIMINATING ACUTE MYELOID LEUKEMIA
- U-321 REDUCTION OF ELEVATED IPTH LEVELS IN THE MGT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS UNDERGONG CHRONIC RENAL DIALYSIS
- U-322 TREATMENT OF ALZHEIMER'S DEMENTIA
- U-323 USE AS A BILE ACID SEQUESTRANT
- U-324 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN, SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS OR ACUTE MANIA EMPLOYING OLANZAPINE
- U-325 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS, INCLUDING "BIPOLAR DISORDER NOS" EMPLOYING OLANZAPINE
- U-326 METHOD OF TREATING SCHIZOPHRENIA AND BIPOLAR DISORDER
- U-327 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED PSYCHOTIC CONDITONS EMPLOYING OLANZAPINE
- U-328 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS INCLUDING "A PSYCHOTIC CONDITION" EMPLOYING AN OLANZAPINE POLYMORPH
- U-329 USE OF AVANDIA AS MONOTHERAPY, IN COMBINATION WITH METFORMIN, AND IN COMBINATION WITH SULFONYLUREAS TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-330 TREATMENT OF NAUSEA AND VOMITING
- U-331 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-332 TREATMENT OR PREVENTION OF BRONCHOSPASM
- U-333 METHOD OF TREATING OCULAR HYPERTENSION
- U-334 TREATMENT OF EXCESSIVE FEMALE FACIAL HAIR
- U-335 USE OF PRAVASTATIN SODIUM FOR SECONDARY PREVENTION OF CORONARY EVENTS IN MEN AND WOMEN WHO HAVE HAD A MYOCARDIAL INFARCTION AND HAVE NORMAL CHOLESTEROL LEVELS
- U-336 DIAGNOSTIC RADIOIMAGING
- U-337 USE OF CARDIOLITE/MIRALUMA KIT FOR THE PREPARATION OF TC99M SESTAMIBI
- U-338 METHODS FOR TREATING DISTURBANCES OF MOOD, DISTURBANCES OF APPETITE, DEPRESSED MOOD, OR CARBOHYDRATE CRAVING ALL ASSOCIATED WITH PREMENSTRUAL SYNDROME
- U-339 PREVENTION OF CARDIO-TOXICITY CAUSED BY THE ADMINISTRATION OF DOXORUBICIN
- U-340 THE LONG TERM TREATMENT OF GROWTH FAILURE DUE TO LACK OF ADEQUATE ENDOGENOUS GROWTH HORMONE SECRETION IN CHILDREN
- U-341 METHOD FOR ENHANCING THE TREATMENT OF ... LATE LUTEAL PHASE DYSPHORIC DISORDER
- U-342 METHOD FOR TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER
- U-343 REDUCTION OF INTESTINAL GAS, CRAMPING AND ANORECTAL IRRITATION
- U-344 METHOD FOR INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH ANOTHER HIV PROTEASE INHIBITOR
- U-345 RITONAVIR AND ANOTHER HIV PROTEASE INHIBITOR FOR CONCOMITANT ADMINISTRATION FOR THE TREATMENT OF AN HIV INFECTION
- U-346 METHOD FOR INHIBITING CYTOCHROME P450 MONOOXYGENASE WITH RITONAVIR AND A METHOD FOR IMPROVING THE PHARMCOKINETICS OF A DRUG THAT IS METABOLIZED BY CYTOCHROME P450 MONOOXYGENASE BY ADMIN THE DRUG AND RITONAVIR
- U-347 METHOD OF USE IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS
- U-348 METHOD OF USE FOR INHIBITING HIV INFECTION
- U-349 METHOD OF USE WHICH IS SUBJECT OF THE APPLICATION
- U-350 PREPARATION OF A PHARMACEUTICAL COMPOSITION FOR CONCOMITANT ADMIN WITH A REVERSE TRANSCRIPTASE INHIBITOR
- U-351 INHIBITING PROTEASE WITH LOPINAVIR AND INHIBITING AN HIV INFECTION WITH LOPINAVIIR
- U-352 INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH A REVERSE TRANSCRIPTASE INHIBITOR
- U-353 PREVENTION AND TREATMENT OF OSTEOPOROSIS

**PATENT AND EXCLUSIVITY TERMS**

ADB 56 of 117

**PATENT USE**

- U-354 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-355 METHOD OF ASSISTING PERSON TO QUIT SMOKING...TRANSDERMALLY ADMIN NICOTINE VIA..PATCH ADHERED TO SKIN AT DOSING RATE APPROX SAME AS ABSORBED FROM SMOKING
- U-356 DELIVERING A MEDICINAL AEROSOL FORMULATION USING CFC-FREE PROPELLANT 134A.
- U-357 USE OF THE DRUG PRODUCT IN PHOTODYNAMIC THERAPEUTIC PROTOCOLS FOR THE TREATMENT OF AGE-RELATED MACULAR DEGENERATION AND RELATED CONDITIONS INVOLVING UNWANTED NEOVASCULATURE IN THE EYE
- U-358 DEPRESSION, OBSESSIVE COMPULSIVE DISORDER, PANIC DISORDER AND SOCIAL ANXIETY DISORDER
- U-359 METHOD OF USE OF VISICOL
- U-360 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS INCLUDING MENTAL DISORDERS EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011
- U-361 MANAGEMENT OF ANXIETY DISORDERS AND THE SHORT-TERM RELIEF OF THE SYMPTOMS OF ANXIETY
- U-362 USE OF APPROVED FORMULATIONS TO TREAT ALL APPROVED DISEASE INDICATIONS
- U-363 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS THAT RELATE TO THE USE OF A PSYCHOACTIVE SUBSTANCE EMPLOYING OLANZAPINE AS PER THE INDICATION THE SUBJECT MATTER OF SUPPLEMENT 011
- U-364 TREATING A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ANY NUMBER OF LISTED CONDITIONS INCLUDING PSYCHOSIS, EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011
- U-365 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN OUR EXTENDED, CONTROLLED RELEASE FORMULATION
- U-366 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN A DELAYED RELEASE FORMULATION
- U-367 TREATMENT OF CARDIOVASCULAR DISORDERS
- U-368 HEARTBURN
- U-369 METHOD OF CONTROLLING AND LOWERING INTRAOCULAR PRESSURE
- U-370 INTRAVAGINAL TREATMENT OF VAGINAL INFECTIONS WITH BUFFERED METRONIDAZOLE COMPOSITIONS
- U-371 APPROVAL FOR MARKETING ONLY UNDER A SPECIAL RESTRICTION PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)
- U-372 METHOD FOR ADMINISTERING A BENEFICIAL DRUG TO THE GI TRACT OF AN ANIMAL, WHICH METHOD COMPRISSES ADMITTING AN OSMOTIC DEVICE ORALLY INTO THE ANIMAL...
- U-373 GENERAL USE CLAIM SUBMITTED FOR 12 NEXIUM PATIENTS STATING "PERTINENT TO THE CAPSULE FORMULATION FOR NEXIUM AND ITS INDICATIONS FOR THE TREATMENT OF GERD AND ERADICATION OF H. PYLORI TO REDUCE THE RISK OF DUODENAL ULCER RECURRENT
- U-374 KIT ADAPTED AND DESIGNED TO PROVIDE BOTH DATA ON THE CURRENT REPRODUCTIVE STATUS OF A PATIENT AND CONTRACEPTION FOR THOSE WHO ARE NOT PREGNANT, BUT RECENTLY ENGAGED IN UNPROTECTED SEX
- U-375 METHOD OF USING RIBAVIRIN FOR TREATING A DISEASE RESPONSIVE TO RIBAVIRIN, E.G. HEPATITIS C
- U-376 TREATMENT OF INFLUENZA
- U-377 METHOD OF TREATING PT WITH CHRONIC HEPATITIS C HAVING HCV GENOTYPE 1 AND VIRAL LOAD GREATER THAN 2 MILLION COPIES/ML TO ERADICATE DETECTABLE HCV-RNA BY ADMIN COMBINATION OF RIBAVIRIN AND INTERFERON ALFA-2B FOR A LEAST 24 WEEKS
- U-378 METHOD FOR TREATING INCONTINENCE
- U-379 METHOD OF TREATINGONYCHROMYCOSIS
- U-380 COMBINATIONS OF TAXOL (PACLITAXEL) AND CISPLATIN WHICH ARE SUITABLE FOR THE TREATMENT OF OVARIAN AND NON-SMALL CELL LUNG CARCINOMAS
- U-381 TREATMENT OF HYPERPHOSPHATEMIA
- U-382 METHOD OF STABILIZING PROSTAGLANDIN
- U-383 METHOD FOR TREATING GLAUCOMA AND OCULAR HYPERTENSION

**PATENT AND EXCLUSIVITY TERMS**

ADB 57 of 117

**PATENT USE**

- U-384 TREATMENT OF CMV RETINITIS
- U-385 TREATMENT OF PEPTIC ULCERS
- U-386 TREATMENT OF PATIENTS SUFFERING FROM A LATE ASTHMATIC REACTION OR LATE PHASE ASTHMA
- U-387 TREATMENT OF PATIENTS WITH RESPIRATORY DISORDERS
- U-388 SMOKING CESSATION AID APPLIED TO THE SKIN
- U-389 SMOKING CESSATION AID APPLIED TO THE SKIN ON WAKING AND REMOVED PRIOR TO SLEEP AFTER ABOUT 16 HOURS
- U-390 METHOD OF USING THE DRUG TO TREAT NEUROIMMUNOLOGIC DISEASES (INCLUDING MULTIPLE SCLEROSIS)
- U-391 USE OF CASODEX IN COMBINATION WITH LHRH AGONISTS FOR THE TREATMENT OF PROSTATE CANCER
- U-392 TREATMENT OF PATIENTS FOR INFLAMMATION
- U-393 MANAGEMENT OF INCONTINENCE, MGT OF HORMONE REPLACEMENT THERAPY, TREATMENT OF INVOLUNTARY INCONTINENCE, MGT OVERACTIVE BLADDER AND INCREASING COMPLIANCE IN SUCH PT
- U-394 METHOD OF USE OF ALPHAGAN
- U-395 METHOD OF USE OF ALPHAGAN P
- U-396 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION
- U-397 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION WITHOUT AN INCREASE IN NAUSEA
- U-398 TREATMENT OF GENERALIZED ANXIETY DISORDER
- U-399 IN-THE-EYE USE OF CHLORINE DIOXIDE CONTAINING COMPOSITIONS
- U-400 USE OF RIBAVIRIN TO INCREASE TYPE 1 CYTOKINE RESPONSE AND SUPPRESS TYPE 2 CYTOKINE RESPONSE TO LYMPHOCYTES, INCLUDING METHODS THAT TAKE ADVANTAGE OF SUCH MODULATION TO TREAT INFECTIONS AND INFESTATIONS
- U-401 USE OF LOPINAVIR IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS FOR TREATING HIV INFECTION AND IN COMBO WITH OTHER HIV PROTEASE INHIBITORS
- U-402 TREATMENT OF ACTINIC KERATOSES
- U-403 ANTI-ALLERGIC FOR VARIOUS ALLERGIC DISEASES
- U-404 TREATMENT OF ALLERGIC CONJUNCTIVITIS
- U-405 FOR WOMEN WITH SEVERE DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS)
- U-406 METHOD OF USE OF ATOVAQUONE AND PROGUANIL
- U-407 METHOD OF TREATING OTOPATHY
- U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASENING FACTOR ANTAGONIST AND RECRUITING OOCYTES FOR IN-VITRO FERTILIZATION
- U-409 METHOD OF TREATING INFLAMMATION USING DRUG SUBSTANCE
- U-410 METHOD OF REDUCING AMOUNT OF RESPECTIVE ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (INCLUDING PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-411 METHOD OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN PREPARATION
- U-412 TREATMENT OF TYPE 2 DIABETES
- U-413 USE OF THE ACTIVE INGREDIENT FOR INHIBITING THE BIOSYNTHESIS OF CHOLESTEROL AND TREATMENT OF ATHEROSCLEROSIS
- U-414 A METHOD OF TREATING GLYCOMETABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE
- U-415 A METHOD FOR REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS
- U-416 A METHOD FOR REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING

**PATENT AND EXCLUSIVITY TERMS**

ADB 58 of 117

**PATENT USE**

- PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS  
U-417 COMBINATION USE OF AD-4833 WITH A BIGUANIDE  
U-418 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER  
U-419 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE  
U-420 METHOD OF TREATMENT OF TYPE II DIABETES  
U-421 USE FOR SEDATION  
U-422 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER AND ATTENTION DEFICIT HYPERACTIVITY DISORDER  
U-423 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER, ATTENTION DEFICIT HYPERACTIVITY DISORDER, OR AIDS RELATED DEMENTIA  
U-424 FOR ONCE DAILY, BOLUS ADMINISTRATION TO A PATIENT IN ORDER TO ENGENDER TREATMENT FOR A NERVOUS DISORDER FOR SUBSTANTIALLY AN ENTIRE DAY ON A CHRONIC BASIS  
U-425 METHOD OF REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMIN TO A DIABETIC BY ADMIN A CHEMICAL COMPOUND HAVING FORMULA (INCL PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER  
U-426 PREVENTION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION  
U-427 METHOD OF TREATING ALLERGIC REACTIONS IN MAMMALS  
U-428 METHOD OF TREATING ALLERGY IN A MAMMAL USING THIS ACTIVE METABOLITE  
U-429 METHOD OF USING DESLORATADINE TO TREAT ALLERGIC RHINITIS  
U-430 METHOD OF TREATING A DIABETIC BY ADMINISTERING AN INSULIN SENSITIZER IN COMBINATION WITH AN INSULIN SECRETION ENHANCER, AND A DRUG PRODUCT COMPRISING AN INSULIN SENSITIZER AND AN INSULIN SECRETION ENHANCER  
U-431 POSTTRAUMATIC STRESS DISORDER  
U-432 REDUCTION OF ATHEROSCLEROTIC EVENTS (MYOCARDIAL INFARCTION, STROKE, AND VASCULAR DEATH) IN PATIENTS WITH ATHEROSCLEROSIS DOCUMENTED BY RECENT STROKE, RECENT MYOCARDIAL INFARCTION OR ESTABLISHED PERIPHERAL ARTERIAL DISEASE  
U-433 USE OF LEVOCARNITINE IN PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS  
U-434 CONTROLLED SYMPTOMS OF DIARRHEA, BLOATING PRESSURE AND CRAMPS, COMMONLY REFERRED TO AS GAS  
U-435 A TITRATION DOSING REGIMEN FOR THE TREATMENT OF PAIN USING AN INITIAL DOSE OF ABOUT 25MG  
U-436 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS  
U-437 METHOD OF USE EQUAL TO PROCESS OF PREPARATION  
U-438 TREATMENT/PREVENTION OF NEURODEGENERATIVE DISEASE  
U-439 TREATMENT OF OBESITY  
U-440 METHOD FOR TRANSDERMAL ADMINISTRATION OF A DRUG THROUGH NON-SCROTAL SKIN USING A TRANSDERMAL DRUG DELIVERY DEVICE CONTAINING THE DRUG AND HAVING AN ADHESIVE SURFACE  
U-441 METHOD OF TREATING MS BY ADMINISTERING COPAXONE  
U-442 METHOD FOR DELIVERING A DRUG TO A PATIENT IN NEED OF THE DRUG, WHILE AVOIDING THE OCCURENCE OF AN ADVERSE SIDE EFFECT KNOWN OR SUSPECTED OF BEING CAUSED BY SAID DRUG  
U-443 MANAGEMENT OF MODERATE TO SEVERE PAIN WHEN A CONTINUOUS, AROUND-THE-CLOCK ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME  
U-444 TREATMENT OF MIGRAINE  
U-445 USE AS AN ANTIMYCOTIC AGENT  
U-446 TOPICAL TREATMENT OF OCULAR HYPERTENSION AND GLAUCOMA  
U-447 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT

**PATENT AND EXCLUSIVITY TERMS**

ADB 59 of 117

**PATENT USE**

- U-448 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-449 USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER WHERE THE DOSE OF LEUCOVORIN IS AT LEAST 200MG PER SQUARE METER
- U-450 INTERMEDIATE REL NICOTINIC ACID FORMULATIONS HAVING UNIQUE URINARY METAB PROFILES RESULTING FROM ABSORPTION PROFILES OF NICOTINIC ACID FROM THE INTERMEDIATE NICOTINIC ACID FORMULATIONS, SUITABLE FOR TX HYPERLIPIDEMIA FOLLOWING QD DOSING
- U-451 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER
- U-452 USE OF LANSOPRAZOLE FOR COMBATTING DISEASES CAUSED BY THE GENUS CAMPYLOBACTER (C.PYLORI=H.PYLORI)
- U-453 TREATMENT OF PLATELET ASSOCIATED ISCHEMIC DISORDERS
- U-454 METHOD OF TX A PT SUSPECTED OF HAVING HEPATITIS C BY ADMIN, IN COMBINATION, A CONJUGATE COMPRISING PEG 12000 & INTERFERON ALFA-2B IN AN AMT OF FROM 0.5MCG/KG TO 2MCG/KG, ONCE WEEKLY, AND RIBAVIRIN
- U-455 TREATMENT OF PULMONARY HYPERTENSION WITH UT-15
- U-456 METHOD OF DECREASING THE PRODUCTION OF A-BETA USING A COMPOSITION WHICH DECREASES BLOOD CHOLESTEROL IN PATIENTS AT RISK OF OR EXHIBITING SYMPTOMS OF ALZHEIMER'S DISEASE
- U-457 METHOD OF TREATING A VAGINAL FUNGAL INFECTION IN A FEMALE HUMAN
- U-458 METHOD OF USE OF IMAGENT
- U-459 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER
- U-460 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING SERTRALINE
- U-461 METHOD OF TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER (PMDD) USING SERTRALINE
- U-462 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND ADULT RHEUMATOID ARTHRITIS AND TREATMENT OF PRIMARY DYSMENORRHEA
- U-463 VENOGRAPHY
- U-464 PERIPHERAL ARTERIOGRAPHY
- U-465 CT IMAGING OF THE HEAD
- U-466 TREATMENT OF IRRITABLE BOWEL SYNDROME
- U-467 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR FOR TREATING HYPERTENSION
- U-468 METHOD OF USING FEXOFENADINE HCL IN TREATING ALLERGIC RHINITIS
- U-469 TREATMENT OF GASTROESOPHAGEAL REFLEX DISEASE (GERD) AND ERADICATION OF H.PYLORI TO REDUCE RISK OF DUODENAL ULCER RECURRENCE
- U-470 THERAPY IN CHRONIC HEPATITIS B VIRUS INFECTION
- U-471 METHOD OF TREATING A PATIENT SUFFERING FROM DIABETES MELLITUS
- U-472 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING METHYLPHENIDATE BI-MODAL RELEASE PROFILE EXTENDED-RELEASE CAPSULES
- U-473 TO REDUCE PLASMA CHOLESTEROL LEVELS IN A MAMMAL
- U-474 TO REDUCE PLASMA CHOLESTEROL LEVELS BY ADMIN EZETIMIBE IN COMBO WITH CHOLESTEROL BIOSYNTHESIS INHIB SELECTED FROM GROUP CONSISTING OF HMG COA REDUCTASE INHIBITORS INCL SIMVASTATIN
- U-475 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-476 METHOD OF TREATING ANDROGEN RESPONSIVE/MEDIATED CONDITION IN MAMMAL BY ADMIN A SAFE, EFFECTIVE AMOUNT OF DUTASTERIDE OR PHARMACEUTICALLY ACCEPTABLE DERIVATIVE THEREOF..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY
- U-477 METHOD OF INHIBITING 5 ALPHA TESTOSTERONE REDUCTASE ENZYME WITH DUTASTERIDE OR ITS DERIVATIVE AND TREATING ANDROGEN RESPONSIVE/MEDIATED DISEASE INCLUDING BENIGN PROSTATIC HYPERPLASIA

**PATENT AND EXCLUSIVITY TERMS**

ADB 60 of 117

**PATENT USE**

- U-478 METHOD OF TREATING HEPATITIS C VIRAL INFECTION BY CONTINUOUS PARENTERAL ADMIN INTERFERON ALPHA 2-10 MILLION IU WEEKLY, SUBCUTANEOUSLY, INJECTION OF POLYMER-INTERFERON ALPHA CONJUGATE-POLYMER IS PEG-INTERFERON IS ALPHA 2B
- U-479 METHOD OF USING PEG-INTRON/REBETOL COMBINATION THERAPY AND INTRON/REBETOL COMBINATION THERAPY
- U-480 CONTRAST AGENT FOR MRI
- U-481 DISUBSTITUTED ACETYLENES BEARING HETEROAROMATIC AND HETEROBICYCLIC GROUPS HAVING RETINOID-LIKE ACTIVITY
- U-482 METHOD OF IN VITRO FERTILIZATION THERAPY INCLUDING MEANS FOR INDUCING OVULATION....
- U-483 METHOD FOR THE ADMINISTRATION OF DRUGS USING THAT COMPOUND
- U-484 METHOD OF TREATING A SKIN DISEASE WITH A CORTICOSTEROID-CONTAINING PHARMACEUTICAL COMPOSITION
- U-485 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-486 EXTERNAL PREPARATION FOR APPLICATION TO THE SKIN CONTAINING LIDOCAINE-DRUG RETAINING LAYER PLACED ON SUPPORT AND COMPRIMES ADHESIVE GEL BASE 1-10% BY WEIGHT OF LIDOCAINE
- U-487 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-488 METHOD FOR REDUCING THE PAIN ASSOCIATED WITH HERPES-ZOSTER AND POST-HERPETIC NEURALGIA
- U-489 EXPECTORANT
- U-490 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-491 METHOD OF DELIVERING A DRUG TO THE LUNG
- U-492 METHOD FOR THE TREATMENT OF SKIN, SUFFERING FROM A CONDITION SELECTED FROM A GROUP CONSISTING OF NONACNE INFLAMMATORY DERMATOSES... COMPRISING APPLYING TO Affected AREA. A THERAPEUTICALLY EFFECTIVE AMT AZELAIC ACID
- U-493 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-494 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER
- U-495 PERITONEAL DIALYSIS SOLUTION
- U-496 METHOD FOR TREATING CHRONIC RENAL FAILURE
- U-497 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND RHEUMATOID ARTHRITIS
- U-498 INTRA-ARTERIAL AND INTRAVENOUS USES OF ULTRAVIST
- U-499 METHOD OF USING REBETOL CAPSULES IN COMBINATION WITH A CONJUGATE COMPRISING POLYETHYLENE GLYCOL(PEG) AND AN ALPHA INTERFERON, INCLUDING, FOR EXAMPLE, PEG-INTRON POWDER FOR INJECTION
- U-500 USE AS AN ANTIHYPERTENSIVE AGENT
- U-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN ADULTS
- U-502 PITYRIASIS VERSICOLOR
- U-503 GENERATOR MUST BE USED WITH INFUSION SYSTEM SPECIFICALLY LABELED FOR USE WITH GENERATOR
- U-504 TINEA PEDIS, TINEA CRURIS, TINEA CORPORIS
- U-505 ULTRASOUND CONTRAST AGENT
- U-506 PHARM PRODUCT CONTAINER 1ST CHAMBER IS DISPOSED AQUEOUS DILUENT SOL 2ND CHAM PHARM ACTIVE AGENT COMPRISING ACETYLCHOLINE, BUFFER IN 1ST CHAM IS SUFFICIENT TO BUFFER PH OF MIXED SOL RESULTING MIXTURE OF AQUEOUS DILUENT SOL & PHARM ACTIVE..
- U-507 ACROMEGALY IN PATIENTS W/INADEQUATE RESPONSE TO SURGERY AND/OR RADIATION THERAPY AND/OR MEDICAL THERAPIES, OR FOR WHOM THESE THERAPIES ARE NOT APPROPRIATE
- U-508 METHOD OF RELEASING 17-BETA OESTRADIOL PRECURSOR IN A SUBSTANTIALLY ZERO ORDER PATTERN FOR AT LEAST THREE WEEKS
- U-509 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY

**PATENT AND EXCLUSIVITY TERMS**

ADB 61 of 117

**PATENT USE**

- U-510 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (STAGE IA AND IB) WHO HAVE REFRACTORY OR PERSISTENT DISEASE AFTER OTHER THERAPIES OR WHO HAVE NOT TOLERATED OTHER THERAPIES
- U-511 USE OF QUINOLONE COMPOUNDS AGAINST ANAEROBIC PATHOGENIC BACTERIA
- U-512 USE OF QUINOLONE COMPOUNDS AGAINST ATYPICAL UPPER RESPIRATORY PATHOGENIC BACTERIA
- U-513 METHODS OF USE OF ANTIMICROBIAL COMPOUNDS AGAINST PATHOGENIC AMYCOPLASMA BACTERIA
- U-514 PREVENTION OF OVULATION IN A WOMAN
- U-515 TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON THE LAST THERAPY
- U-516 METHOD OF TREATING A PSYCHOTIC DISEASE
- U-517 STABLE GEL FORMULATION FOR TOPICAL TREATMENT OF SKIN CONDITIONS
- U-518 OBSESSIVE COMPULSIVE DISORDER
- U-519 POST OPERATIVE NAUSEA AND VOMITING
- U-520 PREMENOPAUSAL OSTEOPOROSIS
- U-521 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTRON A (INTERFERON ALPHA-2 B RECOMBINANT) INJECTION TO TREAT PATIENTS WITH CHRONIC HEPATITIS C
- U-522 TREATMENT OF CMV RETINITIS BY INTRAVITREAL ADMIN OF A PHOSPHOROTHIOATE OLIGONUCLEOTIDE CAPABLE OF HYBRIDIZING WITH CMV mRNA
- U-523 METHOD OF TREATING INFECTION BY CRYPTOSPORIDIUM PARVUM IN AN IMMUNOCOMPROMISED MAMMAL
- U-524 METHOD OF TREATING DIARRHEA
- U-525 METHOD OF TREATING PARASITIC INFECTIONS
- U-526 METHOD OF PROVIDING CONTROLLED RELEASE OF A TREATING AGENT USING A CONTROLLED RELEASE COMPOSITION
- U-527 METHOD OF DELIVERING AN ACTIVE INGREDIENT USING A PROGRESSIVE HYDRATION BIOADHESIVE
- U-528 PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING
- U-529 ONCE DAILY TREATMENT OF ASTHMA WITH NEBULIZED BUDESONIDE
- U-530 TREATMENT OF HERPES ZOSTER, TREATMENT OF GENITAL HERPES, TREATMENT OF COLD SORES, SUPPRESSION OF GENITAL HERPES IN IMMUNOCOMPETENT AND HIV-INFECTED INDIVIDUALS, REDUCTION OF RISK OF HETEROSEXUAL TRANSMISSION OF GENITAL HERPES
- U-531 TREATMENT OF PATIENTS WITH ESSENTIAL HYPERTENSION. MAY BE USED ALONE OR GIVEN WITH OTHER CLASSES OF ANTIHYPERTENSIVES, ESPECIALLY THIAZIDE DERIVATIVES
- U-532 TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD IN PATIENTS REQUIRING MORE THAN ONE BRONCHO DILATOR
- U-533 ERECTILE DYSFUNCTION
- U-534 HUMALOG IS AN INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS FOR THE CONTROL OF HYPERGLYCEMIA
- U-535 TREATMENT OF SOCIAL ANXIETY DISORDER
- U-536 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING
- U-537 TREATMENT OF CONDITIONS RELATED TO HYPERALDOSTERONISM SUCH AS HYPERTENSION AND CARDIAC INSUFFICIENCY, WITH EPLERENONE
- U-538 FIRST LINE TREATMENT OF SEVERE HYPERTENSION, IN PATIENTS WITH HYPERTENSION SEVERE ENOUGH THAT THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY IN THESE PATIENTS
- U-539 TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-540 TREATMENT OF FUNGAL INFECTIONS
- U-541 METHOD OF TREATMENT OF ADULTS INFECTED WITH HIV-1
- U-542 METHOD OF TREATING PATIENT WITH TYPE 2 DIABETES BY ONCE DAILY ADMINISTRATION
- U-543 TREATMENT OF SCHIZOPHRENIA
- U-544 TREATMENT OF OVERACTIVE BLADDER. TREATMENT OF URINARY INCONTINENCE.

**PATENT AND EXCLUSIVITY TERMS**

ADB 62 of 117

**PATENT USE**

- U-545 METHOD FOR THE PREVENTION AND/OR TREATMENT OF THROMBOTIC EPISODES, SUCH AS MYOCARDIAL INFARCTION, IN A HUMAN PATIENT AND METHOD FOR THE PREVENTION OF VENOUS THROMBOSIS IN A POSTOPERATIVE HUMAN PATIENT
- U-546 USE OF REPAGLINIDE IN COMBINATION WITH METFORMIN TO LOWER BLOOD GLUCOSE
- U-547 MAINTENANCE MONOTHERAPY FOR BIPOLAR DISORDER
- U-548 A METHOD OF REDUCING FLUSH IN AN INDIVIDUAL BEING TREATED FOR A LIPIDEMIC DISORDER AND EFFECTIVELY TREATING THE LIPIDEMIC DISORDER
- U-549 USE IN THE TREATMENT OF MEN WITH ADVANCED SYMPTOMATIC PROSTATE CANCER
- U-550 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA
- U-551 METHOD FOR REDUCING TOXICITY OF ALIMTA TREATED PATIENTS BY ADMINISTERING FOLIC ACID
- U-552 TREATMENT OF HYPERTENSION AND HYPERLIPIDEMIA WITH A SINGLE COMPOSITION
- U-553 MANAGEMENT OF PAIN AND DISCOMFORT ASSOCIATED WITH PERIODONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIODONTAL POCKETS
- U-554 TREATING HIV INFECTION WITH INDINAVIR SULFATE IN COMBINATION WITH ANTIRETROVIRAL AGENTS
- U-555 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
- U-556 USE AS ADJUNCT DIAGNOSTIC FOR SERUM THYROID GLLOBULIN (TG) TESTING
- U-557 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-558 INDICATED FOR THE RELIEF OF BRONCHOSPASM IN PATIENTS 2-12 YEARS OF AGE WITH ASTHMA (REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE)
- U-559 METHOD OF DECREASING OR REDUCING PARATHYROID HORMONE LEVEL; METHOD OF MODULATING PARATHYROID HORMONE SECRETION;METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF REDUCING SERUM IONIZED CALCIUM LEVEL
- U-560 METHOD OF DECREASING PARATHYROID HORMONE LEVEL;METHOD OF TREATING HYPERPARATHYROIDISM
- U-561 COSOPT IS INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION WHO ARE INSUFFICIENTLY RESPONSIVE TO BETA BLOCKERS
- U-562 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-563 MARINOL IS INDICATED FOR, INTER ALIA, ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- U-564 TREATMENT OF HIV IN CONCOMITANT THERAPY
- U-565 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS, AND CHRONIC URTICARIA
- U-566 FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-567 METHOD OF TREATING INFERTILITY
- U-568 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION
- U-569 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THEREAFTER AN OVULATORY INDUCING AMOUNT OF HCG IS ADMINISTERED
- U-570 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THE DAILY AMOUNT OF FSH IS ABOUT 5-10 IU/KG
- U-571 TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA AND BIPOLAR I MANIA
- U-572 INTENSIVE CARE UNIT SEDATION
- U-573 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-574 PROPHYLAXIS AND TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND TREATMENT OF THE NASAL SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-575 LOTEMAX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TREATMENT OF STEROID RESPONSIVE CONDITIONS OF THE PALPEBRAL BULBAR CONJUNCTIVA, CORNEA AND ANTERIOR

**PATENT AND EXCLUSIVITY TERMS**

ADB 63 of 117

**PATENT USE**

SEGMENT OF THE GLOBE.

- U-576 ALREX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TEMPORARY RELIEF OF THE SIGNS AND SYMPTOMS OF SEASONAL ALLERGIC CONJUNCTIVITIS.
- U-577 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA WITH FINASTERIDE IN COMBINATION WITH DOXAZOSIN
- U-578 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA, ACUTE EXACERBATION OF CHRONIC BRONCHITIS, AND ACUTE BACTERIAL SINUSITIS CAUSED BY SUSCEPTIBLE STRAINS OF DESIGNATED MICROORGANISMS IN PATIENTS 18 YEARS AND OLDER.
- U-579 TREATMENT OF EPILEPSY AND/OR MIGRAINE.
- U-580 TREATMENT OF DISORDERS OF THE SEROTONERGIC SYSTEM SUCH AS DEPRESSION AND ANXIETY-RELATED DISORDERS
- U-581 METHOD OF TREATING A CONDITION CAPABLE OF TREATMENT BY INHALATION, E.G. ASTHMA, COMPRISING ADMINISTRATION OF A FORMULATION CLAIMED IN US PATENT NO. 6743413
- U-582 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6253762
- U-583 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING TO A PATIENT BY INHALATION, A METERED AEROSOL DOSE OF A DRUG FORMULATION FROM THE METERED DOSE INHALER SYSTEM CLAIMED IN US 6546928
- U-584 SINGLE-DOSE ADMINISTRATION BY THE EPIDURAL ROUTE, AT THE LUMBAR LEVEL, FOR THE TREATMENT OF PAIN FOLLOWING MAJOR SURGERY
- U-585 TO PROMOTE WEIGHT GAIN AFTER WEIGHT LOSS IN CERTAIN TYPES OF PATIENTS
- U-586 AN INTERMEDIATE RELEASE NICOTINIC ACID FORMULATION SUITABLE FOR ORAL ADMINISTRATION ONCE-A-DAY AS A SINGLE DOSE FOR TREATING HYPERLIPIDEMIA WITHOUT CAUSING DRUG-INDUCED HEPATOTOXICITY OR ELEVATIONS IN URIC ACID OR GLUCOSE OR BOTH
- U-587 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR (AND OPTIONALLY A DIURETIC) FOR TREATING CONGESTIVE HEART FAILURE AND HYPERTENSION
- U-588 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER; TREATMENT OF HEARTBURN AND OTHER SYMPTOMS ASSOCIATED WITH GERD; SHORT-TERM TREATMENT OF EROSIONAL ESOPHAGITIS; MAINTENANCE OF HEALING OF EROSIONAL ESOPHAGITIS
- U-589 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN U.S. PATENT NO. 6131966
- U-590 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING TO A PATIENT BY ORAL OR NASAL INHALATION A DRUG FORMULATION BY USING THE METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6532955
- U-591 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING A DOSAGE FORM WHICH PROVIDES ONCE-DAILY ORAL ADMINISTRATION OF A PHENIDATE DRUG
- U-592 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-593 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-594 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- U-595 35 MG ORALLY ONCE A WEEK FOR PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN; 35 MG ORALLY ONCE A WEEK FOR TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-596 TREATMENT OF HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-597 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT HIGH RISK FOR FRACTURE
- U-598 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-599 METHOD FOR TREATING ALLERGIC CONJUNCTIVITIS
- U-600 A METHOD OF TREATING A PATIENT IN NEED OF OPHTHALMIC ANTIMICROBIAL THERAPY WITH LEVOFLOXACIN
- U-601 TREATMENT OF BIPOLAR DISORDER
- U-602 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS, RHEUMATOID ARTHRITIS IN ADULTS, AND/OR

**PATENT AND EXCLUSIVITY TERMS**

ADB 64 of 117

**PATENT USE**

- PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS, ACUTE PAIN IN ADULTS; PRIMARY DYSMENORRHEA; AND/OR ACUTE MIGRAINE ATTACKS IN ADULTS
- U-603 METHOD OF TREATING INFECTIONS COMPRISING ORALLY ADMINISTERING AN EFFECTIVE AMOUNT OF THE FDA APPROVED ORAL SUSPENSION
- U-604 METHOD OF LOWERING BLOOD GLUCOSE BY ONCE DAILY ADMINISTRATION
- U-605 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD); ALTHOUGH THE MECHANISM OF THE ANTIDEPRESSANT ACTION OF DULOXETINE IN HUMANS IS UNKNOWN, IT IS BELIEVED TO BE RELATED TO ITS POTENTIATION OF SERATONERGIC AND NORADRENERGIC ACTIVITY IN THE CNS
- U-606 USE OF IRINOTECAN IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER
- U-607 CANCIDAS IS INDICATED FOR EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS.
- U-608 USE OF QUINOLONE COMPOUNDS AGAINST PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-609 USE OF QUINOLONE COMPOUNDS AGAINST QUINOLONE-RESISTANT PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-610 ATROVENT HFA (IPRATROPIUM BROMIDE HFA) INHALATION AEROSOL IS INDICATED AS A BRONCHODILATOR FOR MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA.
- U-611 METHOD OF USING DESLORATADINE TO TREAT SEASONAL AND PERENNIAL ALLERGIC RHINITIS, PRURITIS, AND CHRONIC IDIOPATHIC URTICARIA IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-612 TREATMENT OF SEASONAL ALLERGY SYMPTOMS WITH NASAL CONGESTION IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-613 REDUCTION OF SERUM PHOSPHATE
- U-614 TREATMENT OF SEXUAL DYSFUNCTION
- U-615 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TOTAL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-616 MANAGEMENT OF PERSISTENT, MODERATE TO SEVERE PAIN IN PATIENTS REQUIRING CONTINUOUS, AROUND-THE-CLOCK ANALGESIA WITH A HIGH POTENCY OPIOID FOR AN EXTENDED PERIOD OF TIME GENERALLY WEEKS TO MONTHS OR LONGER
- U-617 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-618 USE OF ROSUVASTATIN CALCIUM TO REDUCE ELEVATED TOTAL-C, LDL-C, APOB, NONHDL-C OR TG LEVELS; TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA; AND TO SLOW THE PROGRESSION OF ATHEROSCLEROSIS.
- U-619 TREATMENT OF MALIGNANT NEOPLASM
- U-620 TREATMENT OF INSOMNIA
- U-621 METHOD OF TREATING CANCER
- U-622 TREATMENT OF VEGF MEDIATED OCULAR DISEASE.
- U-623 SHORT TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- U-624 REDUCTION OF RISK OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- U-625 ALLERGIC RHINITIS OR NASAL POLYPS
- U-626 CLOLAR IS INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 TO 21 YEARS OLD WITH RELAPSED OR REFRACTORY ACUTE LYMPHOBLASTIC LEUKEMIA AFTER AT LEAST TWO PRIOR REGIMENS
- U-627 TREATMENT OF PATIENTS USING EXTENDED-RELEASE CARBAMAZEPINE
- U-628 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA, AND IN COMBINATION WITH METFORMIN AND A SULFONYLUREA TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-629 METHOD OF INDUCING A HYPNOTIC OR SEDATIVE EFFECT IN A HUMAN BY ADMINISTERING ESZOPICLONE
- U-630 TREATING URINARY INCONTINENCE BY ADMINISTERING AN EXTENDED-RELEASE FORM OF DARIFENACIN
- U-631 TREATING A DISEASE OF ALTERED MOTILITY OR TONE OF SMOOTH MUSCLE BY ADMINISTERING A MUSCARINIC RECEPTOR ANTAGONIZING AMOUNT OF DARIFENACIN

**PATENT AND EXCLUSIVITY TERMS**

ADB 65 of 117

**PATENT USE**

- U-632 METHOD OF TREATMENT OF CANCER BY ADMINISTERING PARTICLES OF PACLITAXEL THAT HAVE A PROTEIN COATING
- U-633 METHOD FOR TREATMENT OF TUMORS BY ADMINISTERING PACLITAXEL AT A DOSE IN THE RANGE OF ABOUT 30MG/METER SQUARE TO ABOUT 100MG/METER SQUARE IN A PHARMACEUTICALLY ACCEPTABLE FORMULATION THAT DOES NOT CONTAIN CREMOPHOR
- U-634 METHOD FOR DELIVERY OF A BIOLOGIC (INCLUDING ANTINEOPLASTIC AGENTS) BY ADMINISTERING TO A PATIENT AN EFFECTIVE AMOUNT OF A BIOLOGIC AS A SOLID OR LIQUID WITH A POLYMERIC BIOCOMPATIBLE MATERIAL
- U-635 TREATMENT OF GERD, MAINTENANCE OF HEALING OF EROSIONAL ESOPHAGITIS AND RISK REDUCTION OF NSAID ASSOCIATED GASTRIC ULCERS
- U-636 TREATMENT OR PREVENTION OF BRONCHOSPASM OR ASTHMATIC SYMPTOMS
- U-637 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST
- U-638 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST, INCLUDING WITH INSULIN
- U-639 TREATMENT OF A MAMMAL HAVING A NEED OF OR REDUCED ABILITY TO PRODUCE INSULIN WITH AN INSULIN AND AN AMYLIN SUCH AS PRAMLINTIDE
- U-640 USE OF AN AMYLIN AGONIST TO REDUCE GASTRIC MOTILITY AND TREAT POST PRANDIAL HYPERGLYCEMIA
- U-641 USE OF AN AMYLIN AGONIST HAVING SPECIFIED BINDING ACTIVITY TO REDUCE GASTRIC MOTILITY, INCLUDING USE THROUGH PARENTERAL ADMINISTRATION
- U-642 TREATMENT AND PREVENTION OF OSTEOPOROSIS
- U-643 THE SHORT TERM TREATMENT (UP TO 10 DAYS) IN PTS HAVING GASTROESOPHAGEAL REFLUX DISEASE (GERD) AS AN ALTERNATIVE TO ORAL THERAPY IN PTS WHEN THERAPY WITH NEXIUM CAPSULES IS NOT POSSIBLE OR APPROPRIATE
- U-644 TREATMENT OF SEASONAL ALLERGIC RHINITIS
- U-645 TREATMENT OF ASTHMA
- U-646 METHOD OF TREATING OTITIS
- U-647 TREATMENT OF OSTEOPOROSIS IN POST MENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- U-648 THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN
- U-649 A METHOD FOR TREATING A TUMOR DISEASE
- U-650 TREATMENT OF ESOPHAGEAL CANDIDIASIS AND PROPHYLAXIS OF CANDIDA INFECTIONS IN HSCT PATIENTS
- U-651 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL)
- U-652 TREATMENT OF CARDIAC ARRHYTHMIA
- U-653 STIMULATING INSULIN RELEASE BY ADMINISTERING EXENATIDE
- U-654 LOWERING PLASMA GLUCAGON IN A SUBJECT IN NEED THEREOF, INCLUDING ONE WITH TYPE 2 DIABETES, BY ADMINISTERING AN EXENDIN OR ANALOG, SUCH AS EXENDIN-4
- U-655 TREATMENT OF MILD TO MODERATE ACTIVE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON AND THE MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR ASCENDING COLON FOR UP TO 3 MONTHS
- U-656 REDUCING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-657 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-658 TREATMENT OF ADVANCED HORMONE-DEPENDENT BREAST CANCER
- U-659 TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) AFTER FAILURE OF AT LEAST ONE PRIOR CHEMOTHERAPY REGIMEN
- U-660 TREATMENT OF HYPERTENSION AND TREATMENT OF HEART FAILURE
- U-661 TREATMENT OF SEIZURE DISORDER
- U-662 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-663 THE TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS
- U-664 TREATMENT OF CONDITIONS FOR WHICH AN ALDOSTERONE RECEPTOR BLOCKER IS INDICATED, SUCH AS HYPERTENSION, HEART FAILURE, AND POST-MYOCARDIAL INFARCTION

**PATENT AND EXCLUSIVITY TERMS**

ADB 66 of 117

**PATENT USE**

- U-665 METHOD OF USING THE DRUG SUBSTANCE/DRUG PRODUCT FOR ULTRASOUND IMAGING
- U-666 METHOD OF TREATING ADHD
- U-667 MANAGEMENT OF INCONTINENCE; METHOD FOR TREATING INCONTINENCE
- U-668 LEVEMIR IS A LONG-ACTING BASAL INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS
- U-669 INDICATION OF TYPE II DIABETES
- U-670 TREATMENT OF HIV-1 INFECTION BY THE CO-ADMINISTRATION OF TIPRANAVIR AND RITONAVIR.
- U-671 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 3 AND 4
- U-672 TREATMENT OF INFLAMMATION OR AN INFLAMMATION-ASSOCIATED DISORDER
- U-673 METHOD OF TREATMENT WITH ONCE-DAILY DOSES OF 625MG/5ML
- U-674 METHOD OF TREATING INSOMNIA CHARACTERIZED BY DIFFICULTY WITH SLEEP ONSET
- U-675 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA; RELIEF OF SYMPTOMS OF ALLERGIC RHINITIS
- U-676 METHOD OF TREATING ATTENTION DEFICIT DISORDER USING ORAL ADMINISTRATION OF A BI-MODAL OR PULSATILE RELEASE COMPOSITION
- U-677 A METHOD OF TREATING DISEASE AMENABLE TO TREATMENT WITH A PHENIDATE DRUG BY ONCE DAILY ORAL ADMINISTRATION OF AN EXTENDED RELEASE DOSAGE FORM
- U-678 METHOD OF TREATING ATTENTION DEFICIT DISORDER AND/OR ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-679 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-680 A METHOD OF TREATING DYSLIPIDEMIA AND DYSLIPOPROTEINEMIA USING A DOSAGE FORM THAT CAN PROVIDE AN EFFECTIVE AMOUNT OF FENOFLIBRATE TO A PATIENT IN A FASTED STATE WHICH IS AT LEAST 90% OF THE AUC AMOUNT PROVIDED BY THE DOSAGE FORM
- U-681 TREATMENT OF PRIMARY IGF-1 DEFICIENCY
- U-682 NON-BENZODIAZEPINE HYPNOTIC AGENT INDICATED FOR TREATMENT OF INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
- U-683 PREVENTION OR TREATMENT OF ISCHEMIC HEART DISEASE
- U-684 TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-685 EXPECTORANT AND COUGH SUPPRESSANT
- U-686 EXPECTORANT AND NASAL DECONGESTANT
- U-687 REDUCING FOOD INTAKE IN A SUBJECT WITH TYPE 2 DIABETES BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-688 TREATMENT OF HIV-INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS
- U-689 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-690 TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-691 USE AS A MONOTHERAPY, IN COMBINATION WITH A SULFONYLUREA, METFORMIN OR INSULIN OR IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-692 USE OF VALSARTAN TO REDUCE CARDIOVASCULAR MORTALITY IN CLINICALLY STABLE PATIENTS WITH LEFT VENTRICULAR FAILURE OR LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- U-693 THE RECOMMENDED INITIAL DOSE OF EQUETRO IS 400MG/DAY GIVEN IN DIVIDED DOSES, TWICE DAILY. THE DOSE SHOULD BE ADJUSTED IN 200MG DAILY INCREMENTS TO ACHIEVE OPTIMAL CLINICAL RESPONSE.
- U-694 LENALIDOMIDE IS AN ANALOGUE OF THALIDOMIDE. THALIDOMIDE IS A KNOWN HUMAN TERATOGEN THAT CAUSES SEVERE LIFE-THREATENING HUMAN BIRTH DEFECTS. IF LENALIDOMIDE IS TAKEN DURING PREGNANCY, IT MAY CAUSE BIRTH DEFECTS OR DEATH TO AN UNBORN BABY.
- U-695 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA AND T-CELL

**PATENT AND EXCLUSIVITY TERMS**

ADB 67 of 117

**PATENT USE**

LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS

- U-696 TREATMENT OF PATIENTS WITH T-CELL LYMPHOBLASTIC LYMPHOMA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-697 A METHOD OF USING RINFABATE RECOMBINANT (RHIGFBP-3) WITH MECASERMIN RECOMBINANT (RHIGF-1) TO PROMOTE LINEAR GROWTH IN THE TREATMENT OF PRIMARY IGF-1 DEFICIENCY
- U-698 METHOD OF USING ANTAGONIST OF ARGININE VASOPRESSIN (AVA) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH EUVOLEMIC HYponatremia
- U-699 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-700 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-701 TREATMENT OF HYPERCHOLESTEROLEMIA AND/OR HYPERTRIGLYCERIDEMIA
- U-702 TOPICAL AEROSOL HAIR REGROWTH TREATMENT
- U-703 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMOR AND RENAL CELL CARCINOMA WITH SUNITINIB
- U-704 METHOD OF ADMINISTERING INSULIN VIA INHALATION
- U-705 TREATING CHRONIC ANGINA BY ADMINISTERING AN EXTENDED RELEASE FORM OF RANOLAZINE
- U-706 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA
- U-707 ALLERGIC RHINITIS
- U-708 TREATMENT OF CHRONIC NON-INFECTIOUS UVEITIS AFFECTING THE POSTERIOR SEGMENT OF THE EYE
- U-709 METHOD OF COMBATING BACTERIA IN A PATIENT
- U-710 A METHOD OF TREATING RESPIRATORY DISORDERS, E.G., ASTHMA, WHICH COMPRISES ADMINISTRATION BY INHALATION OF AN EFFECTIVE AMOUNT OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT NO. 5658549
- U-711 ACUTE AND LONGER-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER
- U-712 A METHOD OF USING A NICOTINIC ACID FORMULATION TO REDUCE ELEVATED TC, LDL-C AND TG LEVELS, AND RAISE HDL-C LEVELS IN PATIENTS WITH HYPERLIPIDEMIA
- U-713 TREATMENT OF MILD TO MODERATE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-714 TOPICAL TREATMENT OF INTERDIGITAL TINEA PEDIS AND TINEA CORPORIS DUE TO TRICHOPHYTON RUBRUM, TRICHOPHYTON MENTAGROPHYTES OR EPIDERMOPHYTON FLOCCOSUM
- U-715 FOR CLEANSING THE BOWEL IN PREPARATION FOR COLONOSCOPY, IN ADULTS 18 YEARS OF AGE OR OLDER
- U-716 THE TREATMENT OR PREVENTION OF BRONCHOSPASM IN ADULTS AND CHILDREN 4 YEARS OF AGE AND OLDER WITH REVERSIBLE OBSTRUCTIVE AIRWAYS DISEASE AND THE PREVENTION OF EXERCISED-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-717 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT
- U-718 TREATMENT OF FUNGAL INFECTIONS
- U-719 TREATMENT OF PSYCHOSIS
- U-720 TREATMENT OF NEUROLEPTIC DISEASES
- U-721 TREATMENT OF INFLUENZA
- U-722 PROPHYLAXIS OF INFLUENZA
- U-723 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-724 METHOD OF TREATING SEIZURES
- U-725 ALLERGIC RHINITIS AND URTICARIA
- U-726 ALLERGIC RHINITIS
- U-727 FOR THE TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD)
- U-728 METHOD FOR TREATING BACTERIAL INFECTION
- U-729 TREATMENT OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER, H. PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- U-730 USE AS A NASAL SPRAY FOR TREATMENT OF THE SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

**PATENT AND EXCLUSIVITY TERMS**

ADB 68 of 117

**PATENT USE**

AND VASOMOTOR RHINITIS

- U-731 USE IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA
- U-732 ACUTE TREATMENT OF THE CUTANEOUS MANIFESTATIONS OF MODERATE TO SEVERE ERYTHEMA NODOSUM LEPROSUM (ENL)
- U-733 MAINTENANCE THERAPY FOR PREVENTION AND SUPPRESSION OF THE CUTANEOUS MANIFESTATIONS OF ENL RECURRENCE
- U-734 FIRST LINE THERAPY FOR TYPE 2 DIABETES MELLITUS
- U-735 METHOD OF TREATING CHRONIC IRON OVERLOAD
- U-736 METHOD FOR IONTOPHORETIC TRANSDERMAL DELIVERY OF FENTANYL HYDROCHLORIDE
- U-737 DISINFECTION OF PATIENT SKIN PRIOR TO AN INVASIVE PROCEDURE
- U-738 INDICATED FOR THE LONG-TERM, TWICE-DAILY MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE OR OLDER
- U-739 METHOD FOR TREATING CONSTIPATION BY OPENING CIC CHANNELS IN A MAMMALIAN SUBJECT
- U-740 FOR THE TREATMENT OF PATIENTS WITH PRIMARY BILIARY CIRRHOSIS
- U-741 COMBINATION THERAPY WITH CISPLATIN FOR THE TREATMENT OF LATE STAGE CERVICAL CANCER
- U-742 TWICE DAILY TOPICAL TREATMENT OF MODERATE TO SEVERE PLAQUE PSORIASIS.
- U-743 ONCE A DAY TOPICAL TREATMENT OF THE INFLAMMATORY LESIONS OF ROSACEA
- U-744 TREATMENT OF HIV INFECTION IN ANTIRETROVIRAL TREATMENT-EXPERIENCED ADULT PATIENTS
- U-745 TREATMENT OR PREVENTION OF EMESIS
- U-746 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT
- U-747 PREVENTION OR TREATMENT OF POST-OPERATIVE NAUSEA AND VOMITING
- U-748 A METHOD FOR THE TREATMENT OF A PROTEIN TYROSINE KINASE-ASSOCIATED DISORDER
- U-749 METHOD OF CONTRACEPTION
- U-750 TREATMENT OF HIV-1 INFECTION IN ADULTS
- U-751 ONCE DAILY DOSING OF BUDESONIDE VIA NEBULIZER FOR THE TREATMENT OF ASTHMA
- U-752 SUNSCREEN
- U-753 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES
- U-754 USE FOR THE LONG-TERM MAINTENANCE TREATMENT OF ASTHMA
- U-755 TREATMENT OF ANOREXIA, CACHEXIA, OR AN UNEXPLAINED, SIGNIFICANT WEIGHT LOSS IN PATIENTS WITH A DIAGNOSIS OF ACQUIRED IMMUNODEFICIENCY SYNDROME (AIDS)
- U-756 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- U-757 USE AS A BILE ACID SEQUESTRANT FOR LOWERING CHOLESTEROL
- U-758 TREATMENT OF SYMPTOMS OF PREMENSTRUAL DYSPHORIC DISORDER
- U-759 METHOD OF USE OF ADMINISTERING LEVOTHYROXINE
- U-760 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS AND TREATMENT OF OROPHARYNGEAL CANDIDIASIS
- U-761 TREATMENT OF SCHIZOPHRENIA INCLUDING MAINTAINING STABILITY IN PATIENTS WITH SCHIZOPHRENIA
- U-762 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-763 ADMINISTRATION OF ARIPIPRAZOLE BY INJECTION
- U-764 TREATMENT OF SCHIZOPHRENIA
- U-765 METHOD OF TREATING ALLERGIC CONJUNCTIVITIS
- U-766 TREATMENT OF SEIZURES
- U-767 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER

**PATENT AND EXCLUSIVITY TERMS**

ADB 69 of 117

**PATENT USE**

- U-768 A METHOD OF REDUCING THE CAPACITY OF EXTENDED RELEASE NICOTINIC ACID TO PROVOKE A FLUSHING REACTION BY PRETREATING AN INDIVIDUAL WITH A FLUSH INHIBITING AGENT PRIOR TO THE ADMINISTRATION OF THE EXTENDED RELEASE NICOTINIC ACID
- U-769 REVЛИMID (LENALIDOMIDE) IN COMBINATION WITH DEXAMETHASONE IS INDICATED FOR THE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-770 LONG-TERM TREATMENT OF PATHOLOGICAL HYPERSECRETOORY CONDITIONS
- U-771 METHOD FOR THE TREATMENT OF DIABETES MELLITUS, SUCH AS TYPE 1 DIABETES MELLITUS OR TYPE 2 DIABETES MELITUS, IN A HUMAN PATIENT
- U-772 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN CHILDREN 2 TO 11 YEARS AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 MONTHS TO 11 YEARS
- U-773 PATHOLOGICAL HYPERSECRETOORY CONDITIONS
- U-774 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR
- U-775 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND/OR A SULFONYLUREA
- U-776 TREATMENT OF CUTANEOUS MANIFESTATION IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL) WHO HAVE PROGRESSIVE, PERSISTENT OR RECURRENT DISEASE ON OR FOLLOWING TWO SYSTEMIC THERAPIES.
- U-777 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE
- U-778 REDUCTION OF ELEVATED INTRAOULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-779 A METHOD FOR TREATMENT OF A CANCER, WHEREIN THE CANCER IS CHRONIC MYELOGENOUS LEUKEMIA
- U-780 A METHOD FOR THE TREATMENT OF CANCER
- U-781 FOR TREATMENT OF ADULT PATIENTS WITH TYPE 2 DIABETES MELLITUS WHO ARE NAIVE TO PHARMACOLOGIC THERAPY
- U-782 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS WITH EVIDENCE OF VIRAL REPLICATION AND EITHER EVIDENCE OF PERSISTANT ELEVATIONS IN SERUM AMINOTRANSFERASES (ALT OR AST) OR HISTOLOGICALLY ACTIVE DISEASE
- U-783 DESONATE GEL IS INDICATED FOR THE TREATMENT OF MILD TO MODERATE ATOPIC DERMATITIS IN PATIENTS 3 MONTHS OF AGE AND OLDER
- U-784 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME (RLS)
- U-785 USE AS REPLACEMENT SOLUTION, HEMOFILTRATION SOLUTION OR HEMODIAFILTRATION SOLUTION IN CONTINUOUS RENAL REPLACEMENT THERAPY
- U-786 PRODUCT IS APPROVED FOR THE TOPICAL TREATMENT OF TINEA PEDIS
- U-787 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND PEDIATRIC PATIENTS SIX YEARS OF AGE OR OLDER, INCLUDING PATIENTS REQUIRING ORAL CORTICOSTEROID THERAPY FOR ASTHMA
- U-788 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING PAROXETINE
- U-789 TREATMENT OF KNOWN OR SUSPECTED CYANIDE POISONING
- U-790 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT RISK FOR FRACTURE. FORTEO CAN BE USED BY PEOPLE WHO HAVE HAD A FRACTURE RELATED TO OSTEOPOROSIS
- U-791 GLEEVEC IS ALSO INDICATED FOR THE TREATMENT OF PATIENTS WITH KIT (CD117) POSITIVE UNRESECTABLE AND/OR METASTATIC MALIGNANT GASTROINTESTINAL STROMAL TUMORS (GIST)
- U-792 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS
- U-793 FOR THE LONG TERM TREATMENT, TWICE DAILY (MORNING AND EVENING) MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-794 CLOSURE OF A CLINICALLY SIGNIFICANT PATENT DUCTUS ARTERIOSUS IN PREMATURE INFANTS WEIGHING BETWEEN 500 AND 1500G, WHO ARE NO MORE THAN 32 WEEKS GESTATIONAL AGE WHEN USUAL MEDICAL MANAGEMENT IS INEFFECTIVE
- U-795 METHOD FOR INHIBITING NOREPINEPHRINE UPTAKE

**PATENT AND EXCLUSIVITY TERMS**

ADB 70 of 117

**PATENT USE**

- U-796 METHOD OF TREATING DEPRESSION
- U-797 METHOD OF TREATING ANXIETY
- U-798 TREATMENT AND PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN BY ONCE-MONTHLY ORAL ADMINISTRATION OF IBANDRONATE SODIUM MONOHYDRATE EQUIVALENT TO 150MG OF IBANDRONIC ACID
- U-799 METHOD FOR INHIBITING SEROTONIN UPTAKE
- U-800 TREATMENT OF PATIENTS WITH ADVANCED OR METASTATIC BREAST CANCER WHOSE TUMORS OVEREXPRESS HER2 AND WHO HAVE RECEIVED PRIOR THERAPY INCLUDING ANTHRACYCLINE, A TAXANE AND TRASTUZUMAB
- U-801 METHOD OF TREATING CANCER
- U-802 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR
- U-803 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN
- U-804 TREATMENT OF ACTINIC KERATOSES BY PHOTODYNAMIC THERAPY
- U-805 TREATMENT OF IMPETIGO DUE TO STAPHYLOCOCCUS AUREUS OR STREPTOCOCCUS PYOGENES
- U-806 INTRATHECAL TREATMENT OF LYMPHOMATOUS MENINGITIS
- U-807 PREVENTION OF EXERCISE-INDUCED BRONCHOCONSTRICTION
- U-808 THE TREATMENT OF THE SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-809 TREATMENT OF CHRONIC IDIOPATHIC URTICARIA
- U-810 METHOD OF TREATMENT TO ALLEVIATE INFLAMMATION OF THE EYE
- U-811 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS AND TREATMENT OF THE UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA
- U-812 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-813 MAINTENANCE TREATMENT OF BRONCHOCONSTRICTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-814 TREATMENT OF SCHIZOPHRENIA
- U-815 TREATS COLD SORES/FEVER BLISTERS ON THE FACE OR LIPS. SHORTENS HEALING TIME AND DURATION OF SYMPTOMS: TINGLING, PAIN, BURNING AND/OR ITCHING
- U-816 DEPRESSION, PANIC DISORDER, PREMENSTRUAL DISORDERS AND SOCIAL ANXIETY DISORDER
- U-817 NASAL ADMINISTRATION OF CYANOCOBALAMIN
- U-818 TOPICAL TREATMENT OF ACNE VULGARIS
- U-819 MANAGEMENT OF FIBROMYALGIA
- U-820 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE SLEEPINESS ASSOCIATED WITH NARCOLEPSY, OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME, AND SHIFT WORK SLEEP DISORDER
- U-821 METHOD OF INHIBITING ENTHOHELIN RECEPTORS BY ADMINISTERING AMBRISENTAN TO A PATIENT TO TREAT PULMONARY ARTERIAL HYPERTENSION.
- U-822 USE IN LIPID MANAGEMENT
- U-823 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS AND FOR THE TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN CHILDREN 6 TO 11 YEARS OF AGE
- U-824 METHOD OF TREATING PATIENTS INFECTED WITH CCR5-TROPIC HIV-1
- U-825 USE FOR PREVENTION OF BREAST CANCER
- U-826 RELIEF OF MODERATE TO SEVERE PAIN
- U-827 USE FOR TREATMENT OF DIABETES, PARTICULARLY TYPE 2 DIABETES
- U-828 PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION
- U-829 TREATMENT OF EXTRAVASATION RESULTING FROM IV ANTHRACYCLINE CHEMOTHERAPY
- U-830 TREATMENT OF RELAPSED SMALL CELL LUNG CANCER

**PATENT AND EXCLUSIVITY TERMS**

ADB 71 of 117

**PATENT USE**

- U-831 METHOD OF ADMINISTERING LANREOTIDE ACETATE
- U-832 ZINGO IS INDICATED FOR THE USE ON INTACT SKIN TO PROVIDE LOCAL ANALGESIA PRIOR TO VENIPUNCTURE OR INTRAVENOUS CANNULATION.
- U-833 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.25% BY WEIGHT OF ROPIVACAINE
- U-834 INVIRASE IN COMBINATION WITH RITONAVIR AND OTHER ANTIRETROVIRAL AGENTS IS INDICATED FOR THE TREATMENT OF HIV INFECTION
- U-835 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF ATOPIC DERMATITIS IN PATIENTS ONE YEAR OF AGE OR OLDER
- U-836 A METHOD FOR THE TREATMENT OF LEUKEMIAS
- U-837 GASTROINTESTINAL LAVAGE INDICATED FOR CLEANSING OF THE COLON AS A PREPARATION FOR COLONOSCOPY IN ADULTS
- U-838 METHOD OF TREATING PAIN USING A PHARMACEUTICALLY ACCEPTABLE SALT OF ROPIVACAINE AND ADMINISTERING A COMPOSITION CONTAINING LESS THAN 0.5% BY WEIGHT OF ROPIVACAINE
- U-839 TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- U-840 TREATMENT FOR TYPE 2 DIABETES MELLITUS
- U-841 INDICATED FOR THE LONG-TERM, MAINTENANCE TREATMENT OF ASTHMA IN PATIENTS 12 YEARS OF AGE AND OLDER
- U-842 INDICATED FOR THE TREATMENT OF ATTENTION-DEFICIT/HYPERACTIVITY DISORDER (ADHD)
- U-843 METHOD FOR ADMINISTRATION OF TESTOSTERONE
- U-844 PREFEST IS INDICATED IN WOMEN WHO HAVE A UTERUS FOR THE TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE; TREATMENT OF VULVAR AND VAGINAL ATROPHY; PREVENTION OF OSTEOPOROSIS
- U-845 TREATMENT OF PATIENTS WITH CANDIDEMIA, ACUTE DISSEMINATED CANDIDIASIS, CANDIDA PERITONITIS AND ABSCESSSES
- U-846 USE FOR DELINEATION (VISUALIZATION) DURING A VITRECTOMY SURGICAL PROCEDURE
- U-847 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-848 ACUTE TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-849 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP. DOSE IS ONE DROP OF COMBIGAN IN THE AFFECTED EYE TWICE DAILY
- U-850 PREVENTION OR TREATMENT OF NAUSEA OR EMESIS INDUCED BY A CANCER CHEMOTHERAPEUTIC AGENT
- U-851 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-852 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-853 TREATMENT OR PREVENTION OF EMESIS
- U-854 PREVENTION OF CMV DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
- U-855 METHOD TO INDUCE NATRIURESIS, DIURESIS AND/OR VASODILATION
- U-856 SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN
- U-857 INHIBITION OF TRANSPLANT REJECTION
- U-858 PEDIATRIC USE AGED 1-11 YEARS, GERD AND EROSIONAL ESOPHAGITIS
- U-859 EROSIONAL ESOPHAGITIS, HYPERSECRETORY CONDITIONS INCLUDING ZOLLINGER-ELLISON SYNDROME, MAINTENANCE OF HEALING OF EROSIONAL ESOPHAGITIS AND REDUCTION OF SYMPTOMS IN PATIENTS WITH GERD
- U-860 FOR THE APPROVED USES AND CONDITIONS OF USE, INCLUDING DEPRESSION
- U-861 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER

**PATENT AND EXCLUSIVITY TERMS**

ADB 72 of 117

**PATENT USE**

- U-862 ADJUNCT TO DIET TO REDUCE ELEVATED TOTAL-C, LDL-C, NON-HDL-C, APO B, TG, AND LP(A) LEVELS AND TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA, MIXED DYSLIPIDEMIA, AND HYPERTRIGLYCERIDEMIA
- U-863 TAKING ASPIRIN OR NON-STEROIDAL ANTI-INFLAMMATORY MEDICATIONS APPROXIMATELY 30 MINUTES BEFORE DOSING CAN MINIMIZE FLUSHING, A COMMON SIDE EFFECT OF NIACIN THERAPY
- U-864 PEDIATRIC USE AGES 1-2 YEARS, GERD AND EROSIONAL ESOPHAGITIS
- U-865 TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND A HIGH RISK FOR BONE FRACTURE BY REDUCING THE RISK OF VERTEBRAL AND NONVERTEBRAL BONE FRACTURE
- U-866 THE LABEL REFERENCES THE EFFECTS OF THE ACTIVE INGREDIENT OF REVOLIMID UPON CYTOKINES
- U-867 TREATMENT OF MIGRAINE
- U-868 METHOD OF USING ANTAGONIST OF ARGinine VASOPRESSIN (AVP) V1A AND V2 RECEPTORS FOR INTRAVENOUS TREATMENT OF PATIENTS WITH HYPERVOLEMIC HYponatremia
- U-869 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART
- U-870 METHOD OF PRODUCING CORONARY VASODILATION WITHOUT PERIPHERAL VASODILATION
- U-871 METHOD OF REDUCING RISK OF MYOCARDIAL INFARCTION, STROKE AND DEATH
- U-872 TWICE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA. TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-873 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME BY OPENING CHLORIDE CHANNELS (CIC)
- U-874 METHOD OF TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME
- U-875 FIRST-LINE TREATMENT OF LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER, IN COMBINATION WITH GEMCITABINE
- U-876 TREATMENT OF MIGRAINE WITH OR WITHOUT AURA
- U-877 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PEPTIC ULCER
- U-878 A METHOD FOR BINDING A PERIPHERAL OPIOID RECEPTOR
- U-879 A METHOD OF TREATING OR PREVENTING ILEUS
- U-880 ENDOMETRIN IS A PROGESTERONE INDICATED TO SUPPORT EMBRYO IMPLANTATION AND EARLY PREGNANCY BY SUPPLEMENTATION OF CORPUS LUTEAL FUNCTION AS PART OF AN ASSISTED REPRODUCTIVE TECHNOLOGY (ART) TREATMENT PROGRAM FOR INFERTILE WOMEN
- U-881 TREATMENT OF NON-SMALL CELL LUNG CANCER
- U-882 MANAGEMENT OF FIBROMYALGIA (FM)
- U-883 TREATMENT OF GASTROINTESTINAL STROMAL TUMOR WITH SUNITINIB
- U-884 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA
- U-885 TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- U-886 ADMINISTERING DESLORATADINE TO TREAT THE SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS, SEASONAL ALLERGIC RHINITIS, OR CHRONIC IDIOPATHIC URTICARIA
- U-887 TREATMENT AND PREVENTION OF OSTEOPOROSIS
- U-888 FEMALE HORMONE REPLACEMENT THERAPY FOR POSTMENOPAUSAL WOMEN
- U-889 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE)
- U-890 REDUCTION OF SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE
- U-891 USE AS AN INTRAOCULAR IRRIGATING SOLUTION DURING SURGICAL PROCEDURES INVOLVING PERfusion OF THE EYE
- U-892 TREATMENT OF CUTANEOUS MANIFESTATIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (CTCL)
- U-893 CLEVIPREX IS A DIHYDROPYRIDINE CALCIUM CHANNEL BLOCKER INDICATED FOR THE REDUCTION OF BLOOD PRESSURE WHEN ORAL THERAPY IS NOT FEASIBLE OR NOT DESIRABLE
- U-894 TREATMENT OF COLD SORES IN PEDIATRIC PATIENTS TWELVE YEARS OF AGE AND OLDER

**PATENT AND EXCLUSIVITY TERMS**

ADB 73 of 117

**PATENT USE**

- U-895 TREATMENT OF HIV INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS
- U-896 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN TWO YEARS OF AGE AND OLDER
- U-897 METHOD OF TREATING TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES IN A ONCE-A-DAY AMOXICILLIN PRODUCT
- U-898 USE OF GLUTAMINE TOGETHER WITH GROWTH HORMONE FOR THE TREATMENT OF PATIENTS WITH SHORT BOWEL SYNDROME
- U-899 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA
- U-900 INTEGRASE INHIBITION FOR THE TREATMENT OF HIV INFECTION
- U-901 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
- U-902 USE IN THE TREATMENT OF THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA (BPH)
- U-903 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN ADULT PATIENTS
- U-904 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE
- U-905 TREATMENT OF MODERATE TO SEVERE VAGINAL DRYNESS AND PAIN WITH INTERCOURSE, SYMPTOMS OF VULVAR AND VAGINAL ATROPHY, ASSOCIATED WITH MENOPAUSE
- U-906 PROPHYLAXIS OF ORGAN REJECTION IN KIDNEY, LIVER AND HEART ALLOGENIC TRANSPLANTS; TREATMENT OF PATIENTS WITH SEVERE ACTIVE, RHEUMATOID ARTHRITIS; TREATMENT OF ADULT, NONIMMUNOCOMPROMISED PATIENTS WITH SEVERE, RECALCITRANT, PLAQUE PSORIASIS
- U-907 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS IN SUBJECTS 18 YEARS OF AGE AND OLDER
- U-908 PROPHYLAXIS OF ORGAN REJECTION IN PATIENTS RECEIVING ALLOGENEIC RENAL TRANSPLANTS
- U-909 TREATMENT OF CYSTIC FIBROSIS PATIENTS WITH PSEUDOMONAS AERUGINOSA
- U-910 TREATMENT OF METASTATIC CARCINOMA OF THE OVARY AFTER FAILURE OF INITIAL OR SUBSEQUENT CHEMOTHERAPY
- U-911 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL TREATMENT IS TEMPORARILY NOT FEASIBLE
- U-912 SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES
- U-913 TREATMENT OF OVERACTIVE BLADDER WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND FREQUENCY
- U-914 METHOD OF TREATING, AS ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER
- U-915 TREATMENT OF MUSCULOSKELETAL CONDITIONS
- U-916 TOPICAL TREATMENT OF ACNE VULGARIS IN PATIENTS 12 YEARS OR OLDER
- U-917 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS
- U-918 TO TREAT OR PREVENT INFECTIONS CAUSED BY SUSCEPTIBLE BACTERIA USING DELAYED-RELEASE TABLETS CONSISTING OF DOXYCYCLINE HYCLATE COATED PELLETS IN A TABLET
- U-919 FOR THE TREATMENT OF DERMATITIS
- U-920 STEROID-RESPONSIVE INFLAMMATORY OCULAR CONDITIONS FOR WHICH A CORTICOSTEROID IS INDICATED AND WHERE SUPERFICIAL BACTERIAL OCULAR INFECTION OR A RISK OF BACTERIAL OCULAR INFECTION EXISTS
- U-921 TREATMENT OF ACNE VULGARIS
- U-922 FOR THE TREATMENT OF FUNGAL INFECTIONS
- U-923 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION
- U-924 TREATMENT OF MILD TO MODERATE INFECTION CAUSED BY SUSCEPTIBLE STRAINS
- U-925 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA
- U-926 MGT SPECIFIC BACTERIAL INFECTIONS. TREATMENT PTS W/ COMMUNITY ACQUIRED PNEUMONIA OR BACTERIAL SINUSITIS DUE TO CONFIRMED, OR SUSPECTED B-LACTAMASE PRODUCING PATHOGENS & S. PNEUMONIAE WITH REDUCED SUSCEPTIBILITY TO PENICILLIN (MIC=2MC/ML)

**PATENT AND EXCLUSIVITY TERMS**

ADB 74 of 117

**PATENT USE**

- U-927 METHOD FOR INCREASING TEAR PRODUCTION
- U-928 TREATMENT OF BACTERIAL INFECTIOUS DISEASE
- U-929 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER TREATABLE WITH AN SSRI
- U-930 TREATMENT OF IDIOPATHIC THROMBOCYTOPENIC PURPURA (ITP)
- U-931 RELIEF OF MODERATE TO SEVERE ACUTE PAIN
- U-932 PYLERA CAPSULES, IN COMBINATION WITH OMEPRAZOLE ARE INDICATED FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI
- U-933 FOR THE TREATMENT OF PATIENTS WITH HELICOBACTER PYLORI INFECTION AND DUODENAL ULCER DISEASE TO ERADICATE H. PYLORI. THE ERADICATION OF HELICOBACTER PYLORI HAS BEEN SHOWN TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- U-934 IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELL TO THE PERIPHERAL BLOOD FOR COLLECTION AND SUBSEQUENT AUTOLOGOUS TRANSPLANTATION WITH NON-HODGKINS LYMPHOMA AND MULTIPLE MYELOMA
- U-935 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) IN PEDIATRIC PATIENTS 6 YEARS OF AGE AND OLDER
- U-936 USE IN COMBINATION WITH GRANULOCYTE-COLONY STIMULATING FACTOR (G-CSF) TO MOBILIZE HEMATOPOIETIC STEM CELLS TO PERIPHERAL BLOOD FOR COLLECTION & SUBSEQUENT AUTOLOGOUS TRANSPLANTATION IN PATIENTS WITH NON-HODGKIN'S LYMPHOMA & MULTIPLE MYELOMA
- U-937 TREATMENT OF PROSTATE CANCER
- U-938 TREATMENT OF HAIR LOSS AND HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-939 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING AND STIMULATING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-940 METHOD TO TREAT AIDS-RELATED KAPOSI'S SARCOMA
- U-941 METHOD TO TREAT OVARIAN CANCER
- U-942 METHOD TO TREAT MULTIPLE MYELOMA
- U-943 GNRH ANTAGONIST INDICATED FOR TREATMENT OF PATIENTS WITH ADVANCED PROSTATE CANCER
- U-944 TREATMENT OF PATIENTS WITH B-CELL CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-945 SEDATIVE-HYPNOTIC AGENT INDICATED FOR MONITORED ANESTHESIA CARE (MAC) SEDATION
- U-946 TREATMENT OF BREAST CANCER
- U-947 WHEN PATIENTS ARE UNABLE TO TAKE THE ORAL FORMULATIONS, PREVACID IV, FOR INJECTION IS INDICATED AS AN ALTERNATIVE FOR THE SHORT-TERM TREATMENT (UP TO 7 DAYS) OF ALL GRADES OF EROSIONAL ESOPHAGITIS
- U-948 TREATMENT OF DIABETES MELLITUS
- U-949 HEALING OF ALL GRADES OF EROSIONAL ESOPHAGITIS (EE) FOR UP TO 8 WEEKS
- U-950 MAINTAIN HEALING OF EROSIONAL ESOPHAGITIS (EE) FOR UP TO 6 MONTHS
- U-951 TREATMENT OF HEARTBURN ASSOCIATED WITH NON-EROSIVE GASTROESOPHAGEAL REFLUX DISEASE (GERD) FOR 4 WEEKS
- U-952 USE AS AN ANALGESIC
- U-953 METHOD OF TREATING OPHTHALMIC INFLAMMATION AND INFECTION
- U-954 CHRONIC MANAGEMENT OF HYPERURICEMIA IN PATIENTS WITH GOUT. NOT RECOMMENDED FOR THE TREATMENT OF ASYMPTOMATIC HYPERURICEMIA
- U-955 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-956 TREATMENT OF PATIENTS WITH H. PYLORI INFECTION AND DUODENAL ULCER DISEASE
- U-957 A METHOD OF TREATING CANCER IN A PATIENT COMPRISING ADMINISTERING IXABEPILONE OR PHARMACEUTICAL COMPOSITIONS COMPRISING IXABEPILONE
- U-958 METHOD OF TREATING PATIENT COMPRISING MIXING FIRST AND SECOND VIALS OF PRODUCT COMPRISING LYOPHILIZED IXABEPILONE TO PROVIDE AN EPOTHILONE ANALOG SOLUTION, DILUTING SOLUTION WITH A SUITABLE DILUENT TO PREPARE INTRAVENOUS FORMULATION FOR PT

**PATENT AND EXCLUSIVITY TERMS**

ADB 75 of 117

**PATENT USE**

- U-959 METHOD OF TREATING CANCER, IV ADMIN, LYOPHILIZED IXABEPILONE DILUTED, EVERY WEEK OR 3 WEEKS; LYOPHILIZED IXABEPILONE WITH SOLVENT(DEHYDRATED ETHANOL) DILUTED TO CONCENTRATION OF 0.1MG/ML TO 0.9MG/ML
- U-960 METHOD OF TREATING CANCER IN A PATIENT COMPRISING INTRAVENOUSLY ADMINISTERING TO THE PATIENT IXABEPILONE DILUTED IN A PARENTERAL DILUENT
- U-961 METHOD OF TREATING BREAST CANCER BY ADMINISTERING IXABEPILONE; A METHOD OF TREATING A CANCER RESPONSIBLE TO MICROTUBULE STABILIZATION BY ADMINISTERING IXABEPILONE
- U-962 SYMBYAX IS INDICATED FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-963 PROZAC AND OLANZAPINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-964 ZYPREXA ZYDIS AND FLUOXETINE IN COMBINATION FOR THE ACUTE TREATMENT OF TREATMENT RESISTANT DEPRESSION IN ADULTS
- U-965 USE OF IXABEPILONE IN COMBINATION WITH CAPECITABINE IN TREATMENT OF METASTASIS BREAST CANCER
- U-966 TREATMENT OF ASTHMA (MAINTENANCE AND PROPHYLACTIC THERAPY)
- U-967 A METHOD OF REVERSING SOFT-TISSUE ANESTHESIA I.E. ANESTHESIA OF THE LIP AND TONGUE, AND THE ASSOCIATED FUNCTIONAL DEFICITS RESULTING FROM AN INTRAORAL SUBMUCOSAL INJECTION OF A LOCAL ANESTHETIC
- U-968 A METHOD FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-969 TREATMENT OF MIGRAINE
- U-970 TOPICAL TREATMENT OF LICE INFESTATIONS
- U-971 INDICATED FOR THE ACUTE TREATMENT OF ADULTS WITH SCHIZOPHRENIA
- U-972 MONOTHERAPY OR AS ADJUNCTIVE THERAPY TO LITHIUM OR VALPROATE FOR THE MAINTENANCE TREATMENT OF BIPOLAR I DISORDER
- U-973 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH PIOGLITAZONE AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON PIOGLITAZONE OR METFORMIN ALONE
- U-974 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-975 TREATMENT OF PULMONARY HYPERTENSION
- U-976 IMPROVEMENT OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES
- U-977 TREATMENT OF ACUTE, UNCOMPLICATED MALARIA INFECTION DUE TO PLASMODIUM FALCIPARUM IN PATIENTS OF 5KG BODYWEIGHT AND ABOVE
- U-978 METHOD OF TREATING HYponatremia
- U-979 RELIEF OF MUSCLE SPASM
- U-980 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN
- U-981 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI-INFLAMMATORY, ANALGESIC, AND ANTIPYRETIC ACTIVITY
- U-982 A METHOD OF TREATING OSTEOPOROSIS
- U-983 METHOD OF TREATING OSTEOPOROSIS IN A POST-MENOPAUSAL WOMAN AT RISK FOR FRACTURE
- U-984 METHOD FOR THE TREATMENT OF A WOMAN WITH OSTEOPOROSIS AND AT RISK FOR BONE FRACTURE
- U-985 TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)
- U-986 TREATMENT OF PATIENTS INFECTED WITH PEDICULUS HUMANUS CAPITIS (HEAD LICE AND THEIR OVA) OF THE SCALP HAIR
- U-987 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH CHRONIC KIDNEY DISEASE ON DIALYSIS
- U-988 TREATMENT OF RHINITIS COMPRISING THE NASAL APPLICATION OF A PHARMACEUTICAL FORMULATION AS CLAIMED IN US PATENT 7541350
- U-989 FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA

**PATENT AND EXCLUSIVITY TERMS**

ADB 76 of 117

**PATENT USE**

- U-990 TREATMENT OF PROTOZOAL INFECTION
- U-991 TREATMENT OR PROPHYLAXIS OF THROMBOSIS OR EMBOLISMS
- U-992 REDUCTION OF THE RISK OF CARDIOVASCULAR HOSPITALIZATION
- U-993 METHOD OF TREATING INFERTILITY
- U-994 METHOD OF TREATMENT OF OSTEOPOROSIS WHEREIN THE OSTEOPOROSIS IS STEROID-INDUCED
- U-995 METHOD FOR TREATING TYPE II DIABETES BY ADMINISTERING SAXAGLIPTIN
- U-996 AN ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL (TC), LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES, AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA
- U-997 TREATMENT OF MAJOR DEPRESSIVE DISORDER BY DOSING AT INTERVALS OF 24 HOURS
- U-998 ADJUNCTIVE THERAPY TO DIET TO REDUCE ELEVATED TOTAL CHOLESTEROL, LOW-DENSITY LIPOPROTEIN CHOLESTEROL, APOLIPOPROTEIN B, TRIGLYCERIDES AND TO INCREASE HDL-C IN ADULT PATIENTS WITH PRIMARY HYPERLIPIDEMIA OR MIXED DYSLIPIDEMIA
- U-999 TREATMENT OF CHRONIC HEPATITIS B IN ADULT PATIENTS
- U-1000 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH HYPERLIPIDEMIAS
- U-1001 METHOD FOR DELIVERING DRUG TO LUNG OF MAMMAL, COMPRISING ADMINISTERING DRUG PRODUCT BY INHALATION. TREATING A MAMMAL HAVING A CONDITION CAPABLE OF TREATMENT BY INHALATION, COMPRISING ADMINISTERING TO THE LUNG THE DRUG PRODUCT BY INHALATION
- U-1002 METHOD OF TREATING INFLAMMATORY CONDITIONS
- U-1003 A METHOD OF MYOCARDIAL PERfusion IMAGING AND INCREASING CORONARY BLOOD FLOW
- U-1004 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA
- U-1005 METHOD OF TREATING A STAPHYLOCOCCAL INFECTION
- U-1006 NEW COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND ADOLESCENTS (12 YEARS OF AGE AND OLDER)
- U-1007 METHOD OF TREATING GOUT FLARES
- U-1008 APPLICATION OF ANTISEPTIC WITH MOISTURIZERS FOR SURGICAL AND HEALTHCARE PERSONNEL SKIN DISINFECTION
- U-1009 METHOD FOR ADMINISTRATION OF TESTOSTERONE
- U-1010 TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA DUE TO TETRA HYDROBIOPTERIN RESPONSIVE PHENYLKETONURIA. KUVAN SHOULD BE TAKEN ORALLY WITH FOOD TO INCREASE ABSORPTION
- U-1011 USE OF GRANisetron TRANSDERMAL SYSTEM TO TREAT/PREVENT CHEMOTHERAPY INDUCED NAUSEA AND VOMITING
- U-1012 METHOD FOR TREATING INSOMNIA WHILE REDUCING THE RISK OF AN ADVERSE DRUG INTERACTION
- U-1013 METHOD OF USING RIBAVIRIN IN COMBINATION WITH PEGYLATED INTERFERON ALPHA-2B TO TREAT PATIENTS WITH CHRONIC HEPATITIS C
- U-1014 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTERFERON ALPHA-2B (PEGYLATED AND NONPEGYLATED) TO TREAT PATIENTS WITH CHRONIC HEPATITIS C
- U-1015 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA
- U-1016 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-EXPERIENCED ADULT PATIENTS, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO AN NNRTI AND OTHER ANTIRETROVIRAL AGENTS
- U-1017 A METHOD OF TREATING NASAL AND NON-NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS
- U-1018 TREATMENT OF PULMONARY HYPERTENSION BY INHALATION
- U-1019 TREATMENT OF PULMONARY HYPERTENSION
- U-1020 METHOD OF USING COLCHICINE FOR THE PROPHYLAXIS OF GOUT FLARES
- U-1021 SHORT-TERM TREATMENT (4-8 WEEKS) OF ACTIVE BENIGN GASTRIC ULCER
- U-1022 FOR THE PREPARATION OF SKIN PRIOR TO SURGERY; HELPS REDUCE BACTERIA THAT CAN POTENTIALLY CAUSE SKIN INFECTION

**PATENT AND EXCLUSIVITY TERMS**

ADB 77 of 117

**PATENT USE**

- U-1023 TREATMENT OF ATROPHIC VAGINITIS DUE TO MENOPAUSE
- U-1024 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION WHO REQUIRE ADJUNCTIVE OR REPLACEMENT THERAPY DUE TO INADEQUATELY CONTROLLED IOP
- U-1025 TREATING FREQUENT HEARTBURN
- U-1026 A METHOD OF TREATING HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS.
- U-1027 REDUCTION OF ELEVATED PLASMA STEROL AND/OR STANOL LEVELS IN A MAMMAL
- U-1028 A METHOD OF DISTRIBUTING SODIUM OXYBATE UNDER CONTROL OF A CENTRAL PHARMACY
- U-1029 METHOD FOR TREATING ACUTE ELEVATIONS OF BLOOD PRESSURE IN HUMAN SUBJECT IN NEED THEREOF
- U-1030 IMPROVEMENT OF WALKING IN PATIENTS WITH MULTIPLE SCLEROSIS (MS)
- U-1031 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA
- U-1032 USE OF ROSUVASTATIN CALCIUM FOR THE PRIMARY PREVENTION OF CARDIOVASCULAR DISEASE IN INDIVIDUALS WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH INCREASED RISK FACTORS
- U-1033 TOPICAL TREATMENT OF ACNE VULGARIS
- U-1034 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER (ADHD) IN ADULTS
- U-1035 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN
- U-1036 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH INSULIN
- U-1037 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH A PPAR-GAMMA AGONIST
- U-1038 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN AND A PPAR-GAMMA AGONIST
- U-1039 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH METFORMIN
- U-1040 INHIBITION OF THROMBIN IN A PATIENT
- U-1041 TREATMENT OF DISORDERS RESPONSIVE TO GROWTH HORMONE
- U-1042 METHOD FOR STIMULATING CORONARY VASODILATION FOR PURPOSES OF IMAGING THE HEART
- U-1043 MANAGEMENT OF MODERATE TO SEVERE PAIN
- U-1044 TOPICAL TREATMENT OF SCALP PSORIASIS
- U-1045 MAINTENANCE TREATMENT IN PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHO HAVE NOT PROGRESSED ON 1ST-LINE TREATMENT WTH PLATINUM-BASED CHEMOTHERAPY
- U-1046 MAINTENANCE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NSCLC WHOSE DISEASE HAS NOT PROGRESSED AFTER FOUR CYCLES PLATINUM-BASED CHEMOTHERAPY
- U-1047 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA (SBCC)
- U-1048 WORKS THROUGH THE INDUCTION OF INTERFERON AND OTHER CYTOKINES
- U-1049 PROPHYLAXIS OF ORGAN REJECTION IN ADULT PATIENTS AT LOW-MODERATE IMMUNOLOGIC RISK RECEIVING A RENAL TRANSPLANT
- U-1050 USE OF METAXALONE FOR TREATMENT OF MUSCULOSKELETAL CONDITIONS
- U-1051 TREATMENT OF OROPHARYNGEAL CANDIDIASIS
- U-1052 RELIEF OF SIGNS AND SYMPTOMS OF ARTHRITIS AND RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER
- U-1053 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER
- U-1054 ONYCHOMYCOSIS OF THE TOENAIL CAUSED BY TRICOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES, ONCE DAILY USE FOR 12 CONSECUTIVE WEEKS
- U-1055 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO ARE ALREADY TREATED WITH A THIAZOLIDINEDIONE (TZD) AND METFORMIN OR WHO HAVE INADEQUATE GLYCEMIC CONTROL ON A TZD OR METFORMIN ALONE
- U-1056 TREATMENT OF PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE
- U-1057 TREATMENT OF INFLAMMATION AND PAIN USING A NASAL SPRAY OF KETOROLAC TROMETHAMINE

**PATENT AND EXCLUSIVITY TERMS**

ADB 78 of 117

**PATENT USE**

- U-1058 USE OF THALIDOMIDE IN COMBINATION WITH DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED MULTIPLE MYELOMA
- U-1059 ADJUNCTIVE THERAPY TO DIET TO PATIENTS WITH HYPERTRIGLYCERIDEMIA
- U-1060 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH Elevated CHOLESTEROL AND/OR LIPID LEVELS
- U-1061 ADJUNCTIVE THERAPY TO DIET IN PATIENTS WITH MIXED DYSLIPIDEMIA
- U-1062 ADMINISTRATION OF APPROVED PRODUCT FOR TREATMENT OF ALZHEIMER'S DISEASE
- U-1063 TREATMENT OF ONLY INFLAMMATORY LESIONS (PAPULES AND PUSTULES) OF ROSACEA
- U-1064 TREATMENT OF BIPOLAR DISORDER AND SCHIZOPHRENIA
- U-1065 METHOD OF TREATING ANDROGEN RESPONSIVE OR MEDICATED CONDITION IN A MAMMAL BY ADMINISTERING A SAFE & EFFECTIVE AMOUNT OF DUTASTERIDE OR A PHARMACEUTICALLY ACCEPTABLE SOLVATE THEREOF.. CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY
- U-1066 METHOD OF TREATING AN ANDROGEN RESPONSE OR MEDiated DISEASE IN A MAMMAL BY ADMINISTERING AN EFFECTIVE ANDROGEN RESPONSIVE OR MEDICATED DISEASE AMOUNT OF DUTASTERIDE..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERPLASIA
- U-1067 TREATMENT OF CANCER
- U-1068 TREATMENT OF ASTHMA
- U-1069 A METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING AN EXCLUSIVE COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION
- U-1070 A METHOD TO CONTROL ABUSE OF A SENSITIVE DRUG BY CONTROLLING WITH A COMPUTER PROCESSOR THE DISTRIBUTION OF THE SENSITIVE DRUG VIA AN EXCLUSIVITY CENTRAL PHARMACY THAT MAINTAINS A CENTRAL DATABASE
- U-1071 METHOD OF TREATING BLADDER DYSFUNCTION WITH ONCE A DAY TROSPiUM SALT FORMULATION
- U-1072 THE MANAGEMENT OF MODERATE TO SEVERE CHRONIC PAIN IN PATIENTS REQUIRING A CONTINUOUS, AROUND-THE-CLOCK OPIOID ANALGESIC FOR AN EXTENDED PERIOD OF TIME
- U-1073 USE FOR THE TREATMENT OF ASTHMA AND COPD
- U-1074 USE OF EXENATIDE MAY RESULT IN REDUCTION IN BODY WEIGHT
- U-1075 USE FOR THE TREATMENT OF ASTHMA
- U-1076 REDUCE CHRONIC SEVERE DROOLING (I.E., SIALORRHEA) IN PATIENTS WITH NEUROLOGIC CONDITIONS ASSOCIATED WITH PROBLEM DROOLING
- U-1077 PRETREATMENT OF PATIENTS WITH VITAMIN B12 AND FOLIC ACID PRIOR TO PEMETREXED DISODIUM ADMINISTRATION
- U-1078 TREATMENT OF ACNE
- U-1079 REVLIIMID (LENALIDOMIDE) IS INDICATED FOR THE TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)
- U-1080 METHOD TO TREAT PULMONARY HYPERTENSION BY ADMINISTERING AMBRISENTAN TO A PATIENT
- U-1081 LUMIGAN IS A PROSTAGLANDIN ANALOG INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-1082 USE OF A COMBINATION OF TOBRAMYCIN AND DEXAMETHASONE TO TREAT OCULAR INFLAMMATION WHERE AN INFECTION OR RISK OF INFECTION EXISTS
- U-1083 ACUTE TREATMENT OF MIGRAINE ATTACKS, WITH OR WITHOUT AURA, AND THE TREATMENT OF CLUSTER HEADACHE EPISODES
- U-1084 RELIEF OF THE INFLAMMATORY AND PRURITIC MANIFESTATIONS OF CORTICOSTEROID RESPONSIVE DERMATOSES IN PATIENTS 12 YEARS OF AGE OR OLDER
- U-1085 METHOD FOR TREATING IRRITABLE BOWEL SYNDROME AND METHOD FOR TREATING ABDOMINAL DISCOMFORT ASSOCIATED WITH IRRITABLE BOWEL SYNDROME
- U-1086 TREATMENT OF AUTOIMMUNE DISEASE
- U-1087 DETECTION OF NON-MUSCLE INVASIVE PAPILLARY CANCER OF THE BLADDER BY PHOTODYNAMIC CYSTOSCOPY
- U-1088 RELIEF OF MUSCLE SPASM
- U-1089 INHIBITION OF THROMBIN
- U-1090 LO LOESTRIN FE IS INDICATED FOR THE PREVENTION OF PREGNANCY IN WOMEN WHO ELECT TO USE ORAL CONTRACEPTIVES AS A METHOD OF CONTRACEPTION

**PATENT AND EXCLUSIVITY TERMS**

ADB 79 of 117

**PATENT USE**

- U-1091 ASSESSMENT OF BRONCHIAL HYPERRESPONSIVENESS IN PATIENTS 6 YEARS OF AGE OR OLDER WHO DO NOT HAVE CLINICALLY APPARENT ASTHMA
- U-1092 TREATMENT OF BREAST CANCER
- U-1093 TREATMENT OF PSEUDOBULBAR AFFECT
- U-1094 MANAGEMENT OF CHRONIC MUSCULOSKELETAL PAIN
- U-1095 METHOD OF TREATING OCULAR INFLAMMATION
- U-1096 TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER
- U-1097 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHEN TREATMENT WITH BOTH SAXagliptin AND METFORMIN IS APPROPRIATE
- U-1098 METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF TREATING HYPERCALCEMIA
- U-1099 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA
- U-1100 REDUCTION OF EXCESS ABDOMINAL FAT IN HIV-INFECTED PATIENTS WITH LIPODYSTROPHY
- U-1101 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY
- U-1102 METHOD OF TREATING CATAPLEXY IN PATIENTS WITH NARCOLEPSY
- U-1103 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-1104 USE OF TRAMADOL FOR THE MANAGEMENT OF MODERATE TO MODERATELY SEVERE CHRONIC PAIN
- U-1105 TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS FOUR (4) YEARS OF AGE AND OLDER
- U-1106 TREATING HYPERTRIGLYCERIDEMIAS WITH REDUCTION OF FOOD EFFECT
- U-1107 TREATING HYPERCHOLESTEROLEMIAS WITH REDUCTION OF FOOD EFFECT
- U-1108 TREATING TYPE 2 DIABETES MELLITUS WITH EXENATIDE BY STIMULATING INSULIN RELEASE
- U-1109 TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL) IN CONNECTION WITH A SPECIAL PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)
- U-1110 METHOD OF TREATING A PATIENT WITH A PRESCRIPTION DRUG USING A COMPUTER DATABASE IN A COMPUTER SYSTEM FOR DISTRIBUTION
- U-1111 NONSTEROIDAL ANTI-INFLAMMATORY DRUG INDICATED FOR RELIEF OF MILD TO MODERATE ACUTE PAIN
- U-1112 METHOD OF MR IMAGING OF A MAMMAL
- U-1113 TREATMENT AND PROPHYLAXIS OF INFLUENZA
- U-1114 TREATMENT WITH GABAPENTIN, INCLUDING TREATMENT OF NEUROPATHIC PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH POSTHERPETIC NEURALGIA
- U-1115 TREATMENT TO REDUCE THE RISK OF COPD EXACERBATIONS IN PATIENTS WITH SEVERE COPD ASSOCIATED WITH CHRONIC BRONCHITIS AND A HISTORY OF EXACERBATIONS
- U-1116 METHOD OF ADMINISTERING COLCHICINE TO FAMILIAL MEDITERRANEAN FEVER PATIENTS
- U-1117 TREATMENT OF BREAST CANCER
- U-1118 USE FOR THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-1119 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING
- U-1120 TO REDUCE GASTROINTESTINAL SIDE EFFECTS ADMINISTER WITH A MEAL; AS STARTING DOSE ADMINISTER ONCE DAILY WITH EVENING MEAL
- U-1121 METHOD OF TREATING TRAVELERS' DIARRHEA
- U-1122 TREATMENT OF SECONDARILY INFECTED TRAUMATIC SKIN LESIONS DUE TO S. AUREUS AND S. PYOGENES
- U-1123 TREATMENT OF ALCOHOL DEPENDENCE
- U-1124 PREVENTION OF RELAPSE TO OPIOID DEPENDENCE, FOLLOWING OPIOID DETOXIFICATION
- U-1125 METHOD FOR THE DETECTION OF NEUROENDOCRINE TUMORS
- U-1126 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE RECEIVED PRIOR CHEMOTHERAPY

**PATENT AND EXCLUSIVITY TERMS**

ADB 80 of 117

**PATENT USE**

CONTAINING DOCETAXEL

- U-1127 TREATMENT OF PATENT DUCTUS ARTERIOSUS
- U-1128 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION IN COMBINATION WITH PEGINFERON ALFA AND RIBAVIRIN IN ADULT PATIENTS (>=18 YEARS OF AGE) WITH COMPENSATED LIVER DISEASE
- U-1129 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITIN AGENT SUCH AS ASPIRIN
- U-1130 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1131 TREATMENT OF HYPERTRIGLYDERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITIN AGENT SUCH AS ASPIRIN
- U-1132 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1133 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1134 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1135 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A) AND INCREASE OF HDL-C
- U-1136 SECONDARY PREVENTION OF CARDIOVASCULAR EVENTS BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1137 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1138 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1139 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1140 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1141 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1142 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1143 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITIN AGENT SUCH AS ASPIRIN
- U-1144 REDUCTION IN ELEVATED TC AND LDL-C BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITIN AGENT SUCH AS ASPIRIN
- U-1145 REDUCTION IN TG BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1146 REDUCTION IN TG WITH REDUCED FLUSHING BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-1147 TREATMENT OF PRIMARY AND MIXED DYSLIPIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1148 REDUCTION IN RISK OF RECURRENT NONFATAL MYOCARDIAL INFARCTION BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION OF LDL-C, TC, TG, LP(A), AND INCREASE OF HDL-C
- U-1149 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, WITH PRETREATMENT WITH A FLUSH INHIBITING AGENT SUCH AS ASPIRIN
- U-1150 TREATMENT OF HYPERCHOLESTEROLEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, TG, LP(A), AND INCREASE OF HDL-C
- U-1151 TREATMENT OF HYPERTRIGLYCERIDEMIA BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT, THROUGH REDUCTION IN TOTAL-C, LDL-C, LP(A), AND INCREASE OF HDL-C
- U-1152 CYANOCOBALAMIN ADMINISTRATION THROUGH NASAL INFUSION

**PATENT AND EXCLUSIVITY TERMS**

ADB 81 of 117

**PATENT USE**

- U-1153 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS, IS INDICATED FOR THE TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS TYPE 1 (HIV-1) INFECTION IN ANTIRETROVIRAL TREATMENT-NAIVE ADULT PATIENTS, AS SET FORTH IN THE LABELING, INCLUDING I&U SECTION
- U-1154 TREATMENT OF PROTEIN KINASE RELATED DISORDERS, SUCH AS GASTROINTESTINAL STROMAL TUMORS, RENAL CELL CARCINOMA AND ADVANCED PANCREATIC NEUROENDOCRINE TUMORS, WITH SUNITINIB
- U-1155 USE OF THALIDOMIDE IN TREATMENT OF CUTANEOUS MANIFESTATIONS OF ERYTHEMA NODOSUM LEPROSUM (ENL)
- U-1156 TO REDUCE BLOOD PHENYLALANINE (PHE) LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA (HPA)
- U-1157 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES IN ADULTS AND CHILDREN 2 YEARS OF AGE AND OLDER AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1158 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1159 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES, SWELLING OF THE NASAL PASSAGES AND SINUS CONGESTION AND PRESSURE IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-1160 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES AND FOR THE RELIEF OF SYMPTOMS ASSOCIATED WITH HIVES (URTICARIA) IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER AND 12 YEARS OF AGE AND OLDER
- U-1161 FOR THE TREATMENT AND PROPHYLAXIS OF GOUT FLARES & THE TREATMENT OF FAMILIAL MEDITERRANEAN FEVER
- U-1162 TREATMENT OF SEBORRHEIC DERMATITIS OF THE SCALP
- U-1163 METHOD OF TREATING THROMBOSIS
- U-1164 METHOD OF TREATING AN ARGATROBAN TREATABLE CONDITION
- U-1165 USE FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1166 A METHOD FOR TREATMENT OF GOUT FLARES DURING PROPHYLAXIS
- U-1167 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT)
- U-1168 THE LONG TERM, ONCE-DAILY MAINTENANCE BRONCHODILATOR TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-1169 MANAGEMENT OF BREAKTHROUGH PAIN IN CANCER PATIENTS 18 YEARS OF AGE AND OLDER WHO ARE RECEIVING AND TOLERANT TO OPIOID THERAPY FOR THEIR UNDERLYING PERSISTENT CANCER PAIN
- U-1170 TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-1171 REDUCTION OF THE RATE OF THROMBOTIC EVENTS IN PATIENTS WITH ACUTE CORONARY SYNDROME
- U-1172 TO REDUCE ELEVATED TOTAL-C, APO B, AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE IN COMBINATION WITH A STATIN
- U-1173 TO REDUCE ELEVATED TOTAL-C, LDL-C, APO B AND NON-HDL-C IN PATIENTS WITH PRIMARY HYPERLIPIDEMIA BY ADMINISTRATION OF EZETIMIBE ALONE OR IN COMBINATION WITH A STATIN OR WITH FENOFLIBRATE
- U-1174 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION, 0.9% SODIUM CHLORIDE INJECTION, OR FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO ADMINISTRATION
- U-1175 REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
- U-1176 TREATMENT OR PREVENTION OF STROKE
- U-1177 REDUCTION OF CARDIAC TISSUE DAMAGE ASSOCIATED WITH MYOCARDIAL INFARCTION
- U-1178 RELIEF OF MODERATE TO SEVERE CHRONIC PAIN
- U-1179 TREATMENT OF A CANCER MEDIATED BY AN ANAPLASTIC LYMPHOMA KINASE (ALK)
- U-1180 TREATMENT OF THE FOLLOWING INFECTIONS: COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS AND STAPHYLOCOCCUS AUREUS BLOODSTREAM INFECTIONS (BACTEREMIA) INCLUDING THOSE WITH RIGHT-SIDED INFECTIVE ENDOCARDITIS

**PATENT AND EXCLUSIVITY TERMS**

ADB 82 of 117

**PATENT USE**

- U-1181 A METHOD OF TREATING OR PREVENTING OCULAR PAIN IN A PATIENT
- U-1182 TREATMENT OF CYCLIC HEAVY MENSTRUAL BLEEDING
- U-1183 A METHOD FOR ADMINISTERING FOLLICLE STIMULATING HORMONE (FSH) FOR OVARIAN FOLLICLE OR TESTICULAR STIMULATION IN THE HUMAN
- U-1184 TREATMENT OF ERECTILE DYSFUNCTION AND THE SIGNS AND SYMPTOMS OF BENIGN PROSTATIC HYPERPLASIA
- U-1185 TREATMENT OF OPIOID-INDUCED CONSTIPATION
- U-1186 ADMINISTRATION OF AN INHALABLE POWDER COMPRISING TIOTROPIUM VIA DEVICE
- U-1187 TREATMENT OF PATHOLOGICAL STATE BY ANTAGONIZING BRADYKININ RECEPTOR INCLUDING TREATMENT OF ACUTE ATTACKS OF HEREDITARY ANGIOEDEMA (HAE)
- U-1188 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE
- U-1189 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH METFORMIN
- U-1190 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH INSULIN
- U-1191 METHOD OF TX TYPE 2 DM IN PTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBO WITH AN AGENT ACTING ON AN ATP-DEPENDENT CHANNEL IN BETA CELLS SUCH AS A SULFONYLUREA(INCL GLIPIZIDE, GLIMEPIRIIDE & GLYBURIDE)
- U-1192 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A SULFONYLUREA (SUCH AS GLIPIZIDE, GLIMEPIRIDE AND GLYBURIDE)
- U-1193 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND SIMVASTATIN IS APPROPRIATE, IN COMBINATION WITH A PPAR-GAMMA AGONIST (SUCH AS PIOGLITAZONE AND ROSIGLITAZONE)
- U-1194 METHOD FOR TREATING INSOMNIA
- U-1195 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 5, WHICH MAY RESULT IN RENAL OSTEODYSTROPHY, WHILE AVOIDING HYPERPHOSPHATEMIA
- U-1196 RELIEF OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS AND OSTEOARTHRITIS AND TO DECREASE RISK OF DEVELOPING UPPER GASTROINTESTINAL ULCERS IN PATIENTS WHO ARE TAKING IBUPROFEN FOR THOSE INDICATIONS
- U-1197 METHOD OF TREATMENT OF CHILDREN WITH CENTRAL PRECOCIOUS PUBERTY
- U-1198 RECTIV IS A NITRATE VASODILATOR INDICATED FOR THE TREATMENT OF MODERATE TO SEVERE PAIN ASSOCIATED WITH CHRONIC ANAL FISSURE
- U-1199 TREATMENT AND PREVENTION OF POSTMENOPAUSAL OR GLUCOCORTICOID-INDUCED OSTEOPOROSIS AND TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- U-1200 REDUCING THE RISK OF STROKE AND SYSTEMIC EMBOLISM
- U-1201 FOR THE TREATMENT OF INTERMEDIATE OR HIGH-RISK MYELOFIBROSIS
- U-1202 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH IRRITABLE BOWEL SYNDROME
- U-1203 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A HUMAN CONSTIPATED PATIENT
- U-1204 TREATMENT OF UVEITIS
- U-1205 TREATMENT OF MACULAR EDEMA
- U-1206 DELIVERING AN OCULAR IMPLANT AS DESCRIBED IN THE DOSAGE AND ADMINISTRATION SECTION OF THE APPROVED LABELING OF OZURDEX
- U-1207 INFANT USE AGED 1 MONTH TO LESS THAN ONE YEAR, GERD AND EROSIONAL ESOPHAGITIS
- U-1208 TREATMENT OF HYPOTRICHOSIS OF THE EYELASHES BY INCREASING THEIR GROWTH INCLUDING LENGTH, THICKNESS AND DARKNESS
- U-1209 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN ADULT PATIENTS, AND TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER
- U-1210 USE OF REVOLIMID (LENALIDOMIDE) WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO REVOLIMID (LENALIDOMIDE)

**PATENT AND EXCLUSIVITY TERMS**

ADB 83 of 117

**PATENT USE**

- U-1211 USE OF REVIMID (LENALIDOMIDE) TO INHIBIT THE SECRETION OF PRO-INFLAMMATORY CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA
- U-1212 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA AND TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)
- U-1213 TOPICAL TREATMENT OF SEBORRHEIC DERMATITIS IN IMMUNOCOMPETENT PATIENTS 12 YEARS OF AGE AND OLDER
- U-1214 METHOD FOR RELIEVING CONSTIPATION IN A HUMAN PATIENT THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MCG+/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1215 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF TRANSFUSION-DEPENDENT ANEMIA IN MYELODYSPLASTIC SYNDROMES (MDS)
- U-1216 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1217 METHOD OF INCREASING HAIR GROWTH
- U-1218 METHOD OF STIMULATING HAIR GROWTH
- U-1219 METHOD OF INCREASING THE NUMBER OF HAIRS
- U-1220 TREATMENT OF RENAL CELL CARCINOMA
- U-1221 TO STIMULATE THE IMMUNE SYSTEM TO INDUCE T CELL PROLIFERATION
- U-1222 TO INHIBIT THE PROLIFERATIVE ACTIVITY OF NEOPLASTIC CELLS
- U-1223 METHOD FOR TREATING TYPE 2 DIABETES USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENATIDE
- U-1224 REDUCTIONS IN BODY WEIGHT ARE OBSERVED WITH EXENATIDE
- U-1225 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING PARTIAL LARGE OR SMALL BOWEL RESECTION SURGERY WITH PRIMARY ANASTOMOSIS
- U-1226 A METHOD OF PROVIDING A PREDETERMINED CONCENTRATION OF NITRIC OXIDE TO A PATIENT
- U-1227 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE
- U-1228 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS FOR WHOM TREATMENT WITH BOTH SITAGLIPTIN AND METFORMIN HCL EXTENDED RELEASE IS APPROPRIATE ALONE OR IN COMBINATION WITH INSULIN
- U-1229 TREATMENT OF MILDLY TO MODERATELY ACTIVE ULCERATIVE COLITIS IN MALE PATIENTS
- U-1230 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT
- U-1231 TREATMENT OF MODERATE-TO-SEVERE PRIMARY RESTLESS LEG SYNDROME IN ADULTS
- U-1232 USE AS ANTICOAGULANT IN PTS W/ UNSTABLE ANGINA UNDERGOING PTCA; W/ PROVISIONAL USE OF GLYCOPROTEIN IIB/IIIA INHIBITOR, AS ANTICOAGULANT IN PTS UNDERGOING PCI AND FOR PTS W/, OR AT RISK OF, HIT/HITTS UNDERGOING PCI. INTENDED FOR USE W/ASPIRIN
- U-1233 TREATMENT OF CHRONIC HEPATITIS C (CHC) GENOTYPE 1 INFECTION, ADMINISTERED WITH FOOD
- U-1234 FOR REDUCING TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES, AND TREATING HYPERTRIGLYCERIDEMIA
- U-1235 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION
- U-1236 USE OF THALOMID (THALIDOMIDE) FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1237 COMBO W/ OTHER ANTIRETROVIRALS FOR TX OF HIV-1 IN ANTIRETROVIRAL TX-EXPERIENCED PT 6 YEARS UP, WHO HAVE EVIDENCE OF VIRAL REPLICATION AND HIV-1 STRAINS RESISTANT TO NON-NUCLEOSIDE REVERSE TRANSCRIPTASE INHIBITOR AND OTHER ANTIRETROVIRALS
- U-1238 TREATMENT OF ANEMIA DUE TO CHRONIC KIDNEY DISEASE
- U-1239 MAGNETIC RESONANCE IMAGING OF THE LIVER
- U-1240 TREATMENT OF HEAVY MENSTRUAL BLEEDING IN WOMEN WITHOUT ORGANIC PATHOLOGY WHO CHOOSE TO USE AN ORAL CONTRACEPTIVE AS THEIR METHOD OF CONTRACEPTION
- U-1241 MANAGEMENT OF MODERATE TO SEVERE PAIN BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED
- U-1242 PREVENTION OF RESPIRATORY DISTRESS (RDS) IN PREMATURE INFANTS

**PATENT AND EXCLUSIVITY TERMS**

ADB 84 of 117

**PATENT USE**

- U-1243 WITH DRY HANDS, GENTLY REMOVE THE SUPRENZA (PHENTERMINE HYDROCHLORIDE ODT) TABLET FROM THE BOTTLE. IMMEDIATELY PLACE THE SUPRENZA TABLET ON TOP OF THE TONGUE WHERE IT WILL DISSOLVE, THEN SWALLOW WITH OR WITHOUT WATER
- U-1244 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH SULFONYLUREA
- U-1245 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-4 INHIBITOR IN COMBINATION WITH PIOGLITAZONE
- U-1246 SINGLE DOSE ADMINISTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA
- U-1247 MANAGEMENT OF POSTHERPETIC NEURALGIA (PHN) IN ADULTS
- U-1248 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL MEDICATION ON THE SAME KNEE
- U-1249 TREATMENT OF MALE PATIENT HAVING A DISEASE OR CONDITION RESPONSIVE TO A TERATOGENIC DRUG
- U-1250 TREATMENT OF PAIN, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY OR SPINAL CORD INJURY, POSTHERPETIC NEURALGIA, AND FIBROMYALGIA
- U-1251 A METHOD OF CONTROLLING POSTOPERATIVE OCULAR PAIN AND BURNING/STINGING IN A PATIENT
- U-1252 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE
- U-1253 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY
- U-1254 METHOD FOR CHRONIC WIEGHT MANAGEMENT BY CONTROLLING WEIGHT GAIN
- U-1255 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY
- U-1256 TREATMENT OF SEBORRHEIC DERMATITIS
- U-1257 TREATMENT OF OPHTHALMIC DISORDERS
- U-1258 VISUALIZATION DURING VITRECTOMY PROCEDURES
- U-1259 PROPHYLAXIS OF HIV-1 INFECTION
- U-1260 TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES INCLUDING BORTEZOMIB AND AN IMMUNOMODULATORY AGENT AND HAVE DEMONSTRATED DISEASE PROGRESSION ON OR WITHIN 60 DAYS OF COMPLETION OF THE LAST THERAPY
- U-1261 REDUCTION OF THE RISK OF HOSPITALIZATION FOR ATRIAL FIBRILLATION
- U-1262 USE OF QSYMIA (PHENTERMINE AND TOPIRAMATE) FOR WEIGHT MANAGEMENT, INCLUDING, BUT NOT LIMITED TO EFFECTING WEIGHT LOSS, TREATING OBESITY, AND/OR TREATING OVERWEIGHT
- U-1263 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) OR CHRONIC BRONCHITIS
- U-1264 TREATMENT OF A RESPIRATORY DISEASE
- U-1265 PATENTED METHOD OF USING REPAGLINIDE IN COMBINATION WITH METFORMIN AS INDICATED FOR IMPROVING GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-1266 METHOD OF TREATING MIDDLE-OF-THE-NIGHT INSOMNIA
- U-1267 TREATMENT OF RHEUMATOID ARTHRITIS BY DELAYED RELEASE FORMULATION OF 1MG OR 2MG OF PREDNISONE
- U-1268 TREATMENT OF PULMONARY, GASTROINTESTINAL AND/OR RHEUMATOLOGICAL DISEASES OR CONDITIONS BY USE OF DELAYED RELEASE FORMULATIONS OF 1MG OR 2MG PREDNISONE
- U-1269 TREATMENT OF RHEUMATOLOGIC, ALLERGIC, PULMONARY, GASTROINTESTINAL, DERMATOLOGIC DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 5MG PREDNISONE TABLET
- U-1270 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH INSULIN (WITH OR WITHOUT METFORMIN AND/OR PIOGLITAZONE)
- U-1271 TREATMENT OF ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME-NEGATIVE(PH-) ACUTE LYMPHOBLASTIC LEUKEMIA (ALL) IN SECOND OR GREATER RELAPSE OR WHOSE DISEASE HAS PROGRESSED FOLLOWING TWO OR MORE ANTI-LEUKEMIA THERAPIES
- U-1272 TREATMENT OF SIGNS AND SYMPTOMS OF PARKINSON'S DISEASE BY APPLICATION OF CLAIMED TRANSDERMAL SYSTEM
- U-1273 TREATMENT OF RESTLESS LEGS SYNDROME BY APPLICATION OF CLAIMED TRANSDERMAL

**PATENT AND EXCLUSIVITY TERMS**

ADB 85 of 117

**PATENT USE**

## DELIVERY SYSTEM

- U-1274 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY DUE TO CYSTIC FIBROSIS OR OTHER CONDITIONS
- U-1275 TREATMENT OF CHRONIC HEPATITIS B IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-1276 MANAGEMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-1277 METHOD OF INCREASING EYELASH GROWTH INCLUDING LENGTH, THICKNESS, DARKNESS AND/OR NUMBER OF EYELASHES BY ADMINISTERING BIMATOPROST TO AN EYELID MARGIN
- U-1278 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULTS
- U-1279 TREATMENT OF HIV INFECTION USING A COMPOSITION CONTAINING A PHARMACOKINETIC ENHANCER THAT INHIBITS CYTOCHROME P450 MONOOXYGENASE
- U-1280 USE OF A CALCIPOTRIENE CONTAINING FOAM FOR THE TREATMENT OF PSORIASIS
- U-1281 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAVE PREVIOUSLY RECEIVED DOCETAXEL
- U-1282 PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING
- U-1283 A METHOD OF TREATING CHRONIC MYELOGENOUS LEUKEMIA
- U-1284 A METHOD OF TREATING A NEOPLASM
- U-1285 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS
- U-1286 A METHOD OF REDUCING THE RISK OF PULMONARY EDEMA IN PATIENTS IN NEED OF TREATMENT WITH INHALED NITRIC OXIDE
- U-1287 METHOD OF REDUCING TG LEVELS IN PATIENT SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA
- U-1288 TREATMENT OF ERECTILE DYSFUNCTION BY ADMINISTERING A FILM-COATED TABLET
- U-1289 MANAGEMENT OF MODERATE TO SEVERE ACUTE PAIN
- U-1290 TREATMENT OF LUNG CANCER
- U-1291 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL) IN PATIENTS WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE (15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- U-1292 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED RELEASE 1, 2, OR 5 MG PREDNISONE TABLET
- U-1293 A METHOD OF LOWERING INTRAOCULAR PRESSURE IN A PATIENT WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-1294 METHOD OF TREATING GLAUCOMA IN A PATIENT
- U-1295 A METHOD OF TREATING A PATIENT WITH GLAUCOMA OR OCULAR HYPERTENSION
- U-1296 USE OF PEMETREXED WITH PRIOR AND/OR REPEATED VITAMIN B12 AND FOLIC ACID ADMINISTRATION
- U-1297 TREATMENT OF PULMONARY ARTERIAL HYPERTENSION BY INHIBITING ENDOTHELIN RECEPTORS
- U-1298 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES
- U-1299 TREATMENT OF PATIENTS WITH LEUKEMIA INCLUDING CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)
- U-1300 TREATMENT OF PATIENTS WITH TYROSINE KINASE INHIBITOR (TKI) RESISTANT OR INTOLERANT CHRONIC MYELOID/MYELOGENOUS LEUKEMIA (CML)
- U-1301 TREATMENT OF DEEP VEIN THROMBOSIS (DVT)
- U-1302 TREATMENT OF PULMONARY EMBOLISM (PE)
- U-1303 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND PULMONARY EMBOLISM
- U-1304 USE OF ONCE-A-DAY AMOXICILLIN PRODUCT TO TREAT TONSILLITIS AND/OR PHARYNGITIS SECONDARY TO STREPTOCOCCUS PYOGENES
- U-1305 TREATMENT OF HIV-1 INFECTION IN ADULT PATIENTS, AND TREATMENT OF HIV-1 INFECTION IN PEDIATRIC PATIENTS 3 YEARS OF AGE AND OLDER, CO-ADMINISTERED WITH RITONAVIR (PREZISTA/RITONAVIR) AND WITH OTHER ANTIRETROVIRAL AGENTS
- U-1306 TREATMENT OF THROMBOCYTOPENIA IN PATIENTS WITH CHRONIC HEPATITIS C TO ALLOW THE INITIATION AND MAINTENANCE OF INTERFERON-BASED THERAPY

**PATENT AND EXCLUSIVITY TERMS**

ADB 86 of 117

**PATENT USE**

- U-1307 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE ADULT PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THERAPY
- U-1308 MULTIPLE MYELOMA
- U-1309 BONE METASTASES
- U-1310 FOR THE MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
- U-1311 METHOD OF TREATING CYSTIC FIBROSIS
- U-1312 USE FOR THE TREATMENT OF HYPERGLYCEMIA
- U-1313 AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-1314 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- U-1315 THE LONG TERM TREATMENT OF PROPHYLACTIC MANAGEMENT OF OCULAR HYPERTENSION AND GLAUCOMA
- U-1316 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES
- U-1317 TREATMENT OF HYPERCHOLESTEROLEMIA, HYPERLIPIDEMIA AND HYPERLIPOPROTEINEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1318 TREATMENT OF HYPERCHOLESTEROLEMIA BY DECREASING THE AMOUNT OR ACTIVITY OF MICROSOMAL TRIGLYCERIDE TRANSFER PROTEIN IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1319 SYMPTOMATIC RELIEF OF NON-INFECTIOUS DIARRHEA
- U-1320 TREATMENT OF ADULT PATIENTS WITH SHORT BOWEL SYNDROME WHO ARE DEPENDENT ON PARENTERAL SUPPORT
- U-1321 TREATMENT OF PULMONARY MULTI-DRUG RESISTANT TUBERCULOSIS
- U-1322 METHOD OF REDUCING OCULAR HYPERTENSION
- U-1323 REDUCING THE RISK OF STROKE
- U-1324 MANAGEMENT OF CYSTIC FIBROSIS PATIENTS
- U-1325 INDUCTION OF REMISSION IN PATIENTS WITH ACTIVE, MILD TO MODERATE ULCERATIVE COLITIS
- U-1326 METHOD OF INDUCING CONTRACEPTION IN A FEMALE OF REPRODUCTIVE AGE WHO HAS NOT YET REACHED PREMENOPAUSE
- U-1327 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF, USING A FLOWABLE HYDROGEL FORMULATION
- U-1328 METHOD FOR TREATING ACUTE MIGRAINE IN ADULTS, WITH OR WITHOUT AURA, COMPRISING IONTOPHORETIC TRANSDERMAL DELIVERY OF SUMATRIPTAN OR A SALT THEREOF
- U-1329 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER
- U-1330 METHODS OF TREATING LIPID METABOLISM AND GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1331 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1332 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1333 METHODS OF LOWERING ELEVATED POST PRANDIAL BLOOD GLUCOSE LEVELS COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR
- U-1334 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-1335 METHODS OF MODIFYING GLUCOSE METABOLISM AND TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE INHIBITOR AND ONE OR MORE OTHER THERAPEUTIC AGENTS SUCH AS METFORMIN
- U-1336 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING A DIPEPTIDYL PEPTIDASE

**PATENT AND EXCLUSIVITY TERMS**

ADB 87 of 117

**PATENT USE****INHIBITOR AND METFORMIN**

- U-1337 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING ALOGLIPTIN
- U-1338 METHOD OF TREATING DIABETES COMPRISING ADMINISTERING A COMPOUND SUCH AS ALOGLIPTIN
- U-1339 METHODS OF TREATING DIABETES COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1340 METHODS OF TREATING LIPID METABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1341 METHODS OF TREATING GLYCOMETABOLISM DISORDERS COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1342 METHODS OF REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1343 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH A BIGUANIDE SUCH AS METFORMIN
- U-1344 METHODS OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT COMPRISING ADMINISTERING AN INSULIN SENSITIVITY ENHANCER SUCH AS PIOGLITAZONE IN COMBINATION WITH AN INSULIN PREPARATION
- U-1345 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH A DOSAGE UNIT COMPRISING 24MICROG+/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1346 USE OF FEBUXOSTAT FOR THE MANAGEMENT OF HYPERURICEMIA IN PATIENTS SUFFERING FROM GOUT AND, WHEN USED WITH THEOPHYLLINE WITHOUT THE NEED FOR DOSE ADJUSTMENT OF THEOPHYLLINE
- U-1347 TREATMENT OF A SKIN DISORDER
- U-1348 TREATMENT OF OSTEOARTHRITIS
- U-1349 TREATMENT OF JUVENILE RHEUMATOID ARTHRITIS
- U-1350 TREATMENT OF ANKYLOSING SPONDYLITIS
- U-1351 TREATMENT OF ACUTE PAIN
- U-1352 TREATMENT OF PRIMARY DYSMENORRHEA
- U-1353 ADJUNCTIVE THERAPY TO LIPID-LOWERING MEDICATIONS AND DIET TO REDUCE LOW DENSITY LIPOPROTEIN-CHOLESTEROL, APOLIPOPROTEIN B, TOTAL CHOLESTEROL, AND NON-HIGH DENSITY LIPOPROTEIN CHOLESTEROL IN PTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1354 INHIBITION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN HYPERSTIMULATION WITH FSH
- U-1355 MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHOD FOR TREATING A RESPIRATORY DISEASE IN A CHILD
- U-1356 TREATMENT OF NASAL SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER. TREATMENT OF NASAL SYMPTOMS ASSOCIATED W PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER
- U-1357 TREATMENT OF SYMPTOMS ASSOCIATED WITH SEASONAL AND PERENNIAL ALLERGIC RHINITIS IN ADULTS AND ADOLESCENTS 12 YEARS OF AGE AND OLDER. PATENT CLAIMS METHODS FOR TREATING A RESPIRATORY DISEASE IN A CHILD
- U-1358 TREATMENT OF BACTERIAL INFECTIONS IN THE NASAL PASSAGE OF ADULT PATIENTS AND HEALTH CARE WORKERS WITH METHICILLIN RESISTANT S. AUREUS
- U-1359 USE OF POMALIDOMIDE TO INHIBIT THE SECRETION OF PRO-INFLAMMATION CYTOKINES, INCLUDING TUMOR NECROSIS FACTOR ALPHA
- U-1360 USE OF POMALIDOMIDE FOR THE TREATMENT OF MULTIPLE MYELOMA
- U-1361 USE OF POMALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO POMALIDOMIDE
- U-1362 TREATMENT OF DISEASES OR CONDITIONS BY THE USE OF A DELAYED-RELEASE 1,2, OR 5MG PREDNISONE TABLET

**PATENT AND EXCLUSIVITY TERMS**

ADB 88 of 117

**PATENT USE**

- U-1363 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING/STINGING FOLLOWING CORNEAL SURGERY
- U-1364 MAINTENANCE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- U-1365 PROPHYLAXIS OF ALLOGRAFT REJECTION IN ADULT PATIENTS RECEIVING A LIVER TRANSPLANT
- U-1366 TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY TO ANOVULATORY INFERTILE WOMEN
- U-1367 METHOD OF ADMINISTERING FSH FOR THE TREATMENT OF INFERTILITY THROUGH INDUCTION OF OVULATION AND PREGNANCY IN ANOVULATORY INFERTILE WOMEN
- U-1368 TREATMENT OF SOLID EXCRETORY SYSTEM TUMORS; ADVANCED RENAL CELL CARCINOMA (RCC), AFTER FAILURE OF TREATMENT WITH SUNITINIB OR SORAFENIB
- U-1369 TREATMENT OF VAGINAL SYMPTOMS OF UROGENITAL ATROPHY BY ORALLY ADMINISTERING OSPEMIFENE WITH FOOD TO ENHANCE BIOAVAILABILITY OF OSPEMIFENE
- U-1370 TREATMENT OF DYSpareunia ASSOCIATED WITH MENOPAUSE
- U-1371 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH ELEVATED INTRAOCULAR PRESSURE OR GLAUCOMA
- U-1372 ADMINISTRATION WITHOUT FOOD FOR TREATMENT OF HIV-1 INFECTION
- U-1373 METHOD OF TREATING ACETAMINOPHEN OVERDOSE WITH ACETYL CYSTEINE SOLUTIONS
- U-1374 TREATMENT OF PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH+CML)
- U-1375 ADASUVE IS A TYPICAL ANTIPSYCHOTIC INDICATED FOR THE ACUTE TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA OR BIPOLAR I DISORDER IN ADULTS
- U-1376 TREATMENT OF INFLAMMATORY LESIONS OF NON-NODULAR MODERATE TO SEVERE ACNE VULGARIS
- U-1377 IMPROVE RESPIRATORY SYMPTOMS IN CYSTIC FIBROSIS IN PATIENTS WITH PSEUDOMONAS AERUGINOSA
- U-1378 TREATMENT OF A NITROGEN METABOLISM DISORDER
- U-1379 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS
- U-1380 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS WHEREIN THE PATIENT HAS CARDIOVASCULAR DISEASE
- U-1381 USE OF PRASUGREL AND ASPIRIN IN PATIENTS REQUIRING THE REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS
- U-1382 TREATMENT OF NAUSEA AND VOMITING OF PREGNANCY IN WOMEN WHO DO NOT RESPOND TO CONSERVATIVE MANAGEMENT
- U-1383 DOSAGE ADJUSTMENT OF A NITROGEN SCAVENGING DRUG IN THE TREATMENT OF A UREA CYCLE DISORDER
- U-1384 METHOD OF TREATING MULTIPLE SCLEROSIS
- U-1385 METHOD OF TREATING AN AUTOIMMUNE DISEASE SELECTED FROM AUTOIMMUNE POLYARTHRITIS AND MULTIPLE SCLEROSIS BUT NOT TREATING PSORIATIC ARTHRITIS
- U-1386 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF A PERSON IN NEED THEREOF
- U-1387 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEvere HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAILY WITH MORNING AND EVENING MEALS
- U-1388 TREATMENT OF PATIENTS WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF WITHOUT SEvere HEART FAILURE AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1389 ELLA IS A PROGESTERONE AGONIST/ANTAGONIST EMERGENCY CONTRACEPTION INDICATED FOR THE PREVENTION OF PREGNANCY FOLLOWING UNPROTECTED INTERCOURSE OR A KNOWN OR SUSPECTED CONTRACEPTIVE FAILURE. ELLA CAN BE TAKEN WITH OR WITHOUT FOOD
- U-1390 A METHOD OF INCREASING THE TESTOSTERONE BLOOD LEVEL OF AN ADULT MALE SUBJECT IN NEED THEREOF
- U-1391 METHOD FOR TREATING OPIOID-INDUCED CONSTIPATION
- U-1392 METHOD OF RELIEVING OR PREVENTING CONSTIPATION IN A HUMAN PATIENT WITH OPIOID-INDUCED CONSTIPATION

**PATENT AND EXCLUSIVITY TERMS**

ADB 89 of 117

**PATENT USE**

- U-1393 METHOD FOR RELIEVING OR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION
- U-1394 METHOD FOR RELIEVING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION THAT COMPRISES ADMINISTERING TO THE PATIENT A DOSAGE UNIT COMPRISING (I) 24MICROG+/- 10% OF A DRUG SUBSTANCE AND (II) A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1395 USE IN RELIEVING OR PREVENTING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION WITH A DOSAGE UNIT COMPRISING 24MICROG +/- 10% OF A DRUG SUBSTANCE AND A PHARMACEUTICALLY SUITABLE EXCIPIENT
- U-1396 TREATMENT OF ADVANCED HORMONE RECEPTOR POSITIVE, HER2-NEGATIVE BREAST CANCER IN COMBINATION WITH EXEMESTANE AFTER FAILURE OF TREATMENT WITH LETROZOLE OR ANASTROZOLE
- U-1397 USE AS AN ANTISEPTIC FOR THE PREPARATION OF A PATIENT'S SKIN PRIOR TO SURGERY
- U-1398 METHOD OF TREATING CHRONIC HEPATITIS C
- U-1399 MANAGEMENT OF NEPHROPATHIC CYSTINOSIS BY ADMINISTERING A TOTAL DAILY DOSE IN TWO DIVIDED DOSES
- U-1400 FOR THE TREATMENT OF PRIMARY HYPERLIPIDEMIA, MIXED HYPERLIPIDEMIA OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- U-1401 INDICATED FOR LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PTS WITH A HISTORY OF EXACERBATIONS
- U-1402 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR INDOLENT B-CELL NON-HODGKIN LYMPHOMA (NHL)
- U-1403 FIRST-LINE TREATMENT OF METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) WITH EGFR EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST
- U-1404 METHOD FOR TREATING CONSTIPATION IN A PATIENT WITH OPIOID-INDUCED CONSTIPATION BY OPENING CIC CHANNELS
- U-1405 THERAPEUTIC TREATMENT OF BONE METASTASES
- U-1406 TREATMENT OF MELANOMA
- U-1407 TREATMENT OF NEWLY DIAGNOSED PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (PH + CML)
- U-1408 TREATMENT OF PLAQUE PSORIASIS IN PATIENTS 18 YEARS OF AGE OR OLDER
- U-1409 TREATMENT OF HIV-1 BY ONCE DAILY ADMINISTRATION
- U-1410 TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES
- U-1411 THIS DRUG IS ADMINISTERED BY SUBLINGUAL ROUTE TO HUMANS FOR MAINTENANCE TREATMENT OF OPIOID DEPENDENCE
- U-1412 TREATMENT OF ATOPIC DERMATITIS
- U-1413 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH FLOLAN STERILE DILUENT FOR INJECTION PRIOR TO INFUSION
- U-1414 USE OF REVLIMID (LENALIDOMIDE) FOR THE TREATMENT OF MANTLE CELL LYMPHOMA (MCL)
- U-1415 TREATING A PATIENT HAVING A CONDITION SUSCEPTIBLE TO TREATMENT WITH METHYLPHENIDATE, SUCH AS ADHD, BY ADMINISTERING THE FORMULATION RECITED IN CLAIMS 1 OR 2
- U-1416 USE OF FENOFLIBRATE FOR REDUCING ELEVATED TOTAL CHOLESTEROL (TOTAL-C), LDL-C, APO-LIPOPROTEIN B, OR TOTAL TRIGLYCERIDES
- U-1417 USE FOR TREATMENT OF HELICOBACTER INFECTIONS
- U-1418 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAFV600E MUTATION AS DETECTED BY AN FDA APPROVED TEST
- U-1419 TREATMENT OF ACUTE CYANIDE POISONING THAT IS JUDGED TO BE LIFE THREATENING
- U-1420 METHOD OF ONCE A DAY ADMINISTRATION
- U-1421 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE
- U-1422 METHOD OF TREATING PATIENTS NEEDING AN IRON SUPPLEMENT
- U-1423 AMYVID IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET)

**PATENT AND EXCLUSIVITY TERMS**

ADB 90 of 117

**PATENT USE**

IMAGING OF THE BRAIN TO ESTIMATE BETA-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT

- U-1424 LONG-TERM, ONCE DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS
- U-1425 SUBLINGUAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE
- U-1426 USE FOR TREATMENT OF DIAPER DERMATITIS COMPLICATED BY CANDIDIASIS
- U-1427 ALKYLATING DRUG INDICATED FOR THE TOPICAL TREATMENT OF STAGE IA AND IB MYCOSIS FUNGOIDES-TYPE CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO HAVE RECEIVED PRIOR SKIN DIRECTED THERAPY
- U-1428 TOPICAL TREATMENT OF FACIAL ERYTHEMA OF ROSACEA
- U-1429 TREATMENT OF PATIENTS WITH BREAST CANCER WHOSE TUMORS OVEREXPRESS THE HER2 RECEPTOR
- U-1430 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- U-1431 METHOD OF TREATING HYPERGLYCEMIA TO IMPROVE GLYCEMIC CONTROL IN A PATIENT BY ORAL ADMIN OF ONCE A DAY OSMOTIC DOSAGE FORM OF GLIPIZIDE WITH POLYETHYLENE OXIDE, HYDROXYPROPYLMETHYLCELLULOSE, CELLULOSE ACETATE, AND SODIUM CHLORIDE
- U-1432 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX
- U-1433 IMPROVEMENTS OF GLYCEMIC CONTROL IN INDIVIDUALS WITH TYPE 2 DIABETES WHO HAVE ONE OR MORE SPECIFIED CARDIOVASCULAR RISK FACTORS
- U-1434 TREATMENT OF PANCREATIC CANCER
- U-1435 COMBINATION USE OF TOPICAL DICLOFENAC ON THE KNEE AND ADMINISTRATION OF AN ORAL NSAID.
- U-1436 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL AGENT SELECTED FROM SUNSCREEN AND INSECT REPELLANT
- U-1437 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE DILUENT FOR FLOLAN OR STERILE DILUENT FOR EPOPROSTENOL SODIUM PRIOR TO ADMINISTRATION
- U-1438 ZINGO INTRADERMAL INJECTION SYSTEM IS A DRUG DELIVERY SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION
- U-1439 METHOD OF TREATING AN AFFECTIVE DISORDER SUCH AS DEPRESSION
- U-1440 USE OF INGENOL MEBUTATE TO TREAT ACTINIC KERATOSIS
- U-1441 A METHOD OF TREATING OR REDUCING OCULAR PAIN AND BURNING/STINGING
- U-1442 SUBCUTANEOUS INJECTION OF METHOTREXATE
- U-1443 ACCELERATING THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS
- U-1444 A DOSING REGIMEN OF AGGRASTAT (TIROFIBAN HYDROCHLORIDE) (25MCG/KG FOLLOWED BY 0.15MCG/KG/MIN INFUSION) TO REDUCE THE RATE OF THROMBOTIC CORONARY EVENTS ASSOCIATED WITH ACUTE CORONARY SYNDROME (ACS) IN PATIENTS WITH NON-ST ELEVATION ACS
- U-1445 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 1% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION
- U-1446 METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING ADMINISTERING MACITENTAN IN COMBINATION WITH A COMPOUND HAVING PHOSPHODIESTERASE-5 INHIBITORY PROPERTIES
- U-1447 TREATING PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIA
- U-1448 TREATING SEVERE HYPERTRIGLYCERIDEMIA
- U-1449 METHOD OF ALLEVIATING A SKIN CONDITION
- U-1450 TREATMENT OF ALLERGIC RHINITIS SYMPTOMS
- U-1451 APPROVED INDICATIONS: APTIOM (ESLICARBAZEPINE ACETATE) IS INDICATED AS ADJUNCTIVE TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH

**PATENT AND EXCLUSIVITY TERMS**

ADB 91 of 117

**PATENT USE**

EPILEPSY. PATENT CLAIMS: IN A METHOD OF TREATING A SUBJECT AFFLICTED WITH EPILEPSY

- U-1452 METHOD FOR CHRONIC WEIGHT MANAGEMENT
- U-1453 A METHOD OF TREATING HYPOXIC RESPIRATORY FAILURE BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT
- U-1454 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS
- U-1455 TREATMENT OF PERIANAL WARTS
- U-1456 TREATMENT OF MANTLE CELL LYMPHOMA
- U-1457 A METHOD OF PURGING A NITRIC OXIDE DELIVERY SYSTEM
- U-1458 A METHOD OF REDUCING INTRAOCCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-1459 TREATMENT OF CARCINOMA OF THE THYROID
- U-1460 TREATMENT OF HERPES LABIALIS
- U-1461 A METHOD OF GENERATING AN INJECTABLE FOAM OF CONTROLLED DENSITY AND BUBBLE SIZE
- U-1462 A METHOD OF USING A SCLEROSING AGENT FOR THE TREATMENT OF INCOMPETENT GREAT SAPHENOUS VEINS, ACCESSORY SAPHENOUS VEINS AND VISIBLE VARICOSEITIES OF THE GREAT SAPHENOUS (GSV) SYSTEM ABOVE AND BELOW THE KNEE
- U-1463 A METHOD OF INTRAVENOUS INJECTION USING ULTRASOUND GUIDANCE, ADMINISTERED VIA A SINGLE CANNULA INTO THE LUMEN OF THE TARGET INCOMPETENT TRUNK VEINS OR BY DIRECT INJECTION INTO VARICOSEITIES
- U-1464 TREATMENT OF OPIOID DEPENDENCE/SUBLINGUAL OR BUCCAL APPLICATION
- U-1465 USE OF THALIDOMIDE WHILE PREVENTING THE EXPOSURE OF A FETUS OR OTHER CONTRAINDICATED INDIVIDUAL TO THALIDOMIDE
- U-1466 RELIEF OF SYMPTOMS ASSOCIATED WITH RESPIRATORY ALLERGIES ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-1467 METHOD OF TREATING HEPATITIS C
- U-1468 CONTROL OF PHOSPHOROUS LEVELS IN PATIENTS
- U-1469 USE OF PHOSYRLA FOR REDUCTION OF SERUM PHOSPHOROUS IN PATIENTS
- U-1470 FOR THE TREATMENT OF HEPATITIS C
- U-1471 A METHOD FOR TREATING CARDIOVASCULAR DISEASE COMPRISING ADMINISTERING A RECONSTITUTED LYOPHILIZED PHARMACEUTICAL COMPOSITION COMPRISING EPOPROSTENOL, ARGinine AND SODIUM HYDROXIDE.
- U-1472 INTENSIVE CARE UNIT SEDATION, INCLUDING SEDATION OF NON-INTUBATED PATIENTS PRIOR TO AND/OR DURING SURGICAL AND OTHER PROCEDURES
- U-1473 MANAGEMENT OF RISK OF DRONEDARONE/BETA-BLOCKER INTERACTION IN PATIENTS IN SINUS RYTHM WITH A HISTORY OF PAROXYSMAL OR PERSISTENT AF
- U-1474 A METHOD FOR THE TREATMENT OF A PATIENT SUFFERING FROM A DISEASE TREATABLE WITH ROTIGOTINE, COMPRISING APPLYING THE CLAIMED TRANSDERMAL DELIVERY SYSTEM (TDS) TO THE SKIN OF THE PATIENT
- U-1475 USE OF ORENITRAM FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) (WHO GROUP 1).
- U-1476 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA.
- U-1477 USE OF TOPICAL DICLOFENAC ON THE KNEE AND A SECOND TOPICAL PRESCRIPTION MEDICATION ON THE SAME KNEE
- U-1478 METHOD OF REDUCING TG LEVELS IN PATIENT ON STATIN THERAPY SUFFERING FROM SEVERE HYPERTRIGLYCERIDEMIA
- U-1479 INCREASE TEAR PRODUCTION TO TREAT PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).
- U-1480 TREATMENT OF ADVANCED RENAL CELL CARCINOMA
- U-1481 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE
- U-1482 DICLOFENAC POTASSIUM FOR RELIEF OF MILD TO MODERATE ACUTE PAIN
- U-1483 INCREASE TEAR PRODUCTION IN PATIENTS WITH KERATOCONJUNCTIVITIS SICCA (DRY EYE).

**PATENT AND EXCLUSIVITY TERMS**

ADB 92 of 117

**PATENT USE**

- U-1484 COMBINATION PRODUCT FOR THE EARLY TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) TO REDUCE THE LIKELIHOOD OF ULCERATIVE COLD SORES AND TO SHORTEN THE LESION HEALING TIME IN ADULTS AND CHILDREN (6 YEARS OF AGE AND OLDER)
- U-1485 TREATING A SUBJECT UNDERGOING ABDOMINAL SURGERY BY ADMINISTERING ALVIMOPAN TO ACCELERATE THE TIME TO UPPER AND LOWER GASTROINTESTINAL RECOVERY FOLLOWING SURGERIES THAT INCLUDE PARTIAL BOWEL RESECTION WITH PRIMARY ANASTOMOSIS
- U-1486 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER
- U-1487 METHOD OF INCREASING EYELASH GROWTH
- U-1488 USE OF TOPICAL DICLOFENAC FOR TREATING PAIN
- U-1489 USE OF TOPICAL DICLOFENAC ON A JOINT FOR TREATING OSTEOARTHRITIS
- U-1490 FOR USE IN PATIENTS HAVING SYMPTOMATIC OR PROGRESSIVE MEDULLARY THYROID CANCER, WITH UNRESECTABLE LOCALLY ADVANCED OR METASTATIC DISEASE
- U-1491 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA
- U-1492 TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- U-1493 METHOD FOR PREVENTING ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS
- U-1494 SUBLINGUAL OR BUCCAL ADMINISTRATION OF A PHARMACEUTICAL COMPOSITION COMPRISING BUPRENORPHINE AND NALOXONE
- U-1495 RISK REDUCTION OF REBLEEDING IN PTS FOLLOWING THERAPEUTIC ENDOSCOPY FOR ACUTE BLEEDING GASTRIC OR DUODENAL ULCERS IN ADULTS.
- U-1496 METHOD TO TREAT HEMANGIOMA.
- U-1497 NEURACEQ IS A RADIOACTIVE DIAGNOSTIC AGENT FOR POSITRON EMISSION TOMOGRAPHY (PET) IMAGING OF THE BRAIN TO ESTIMATE P-AMYLOID NEURITIC PLAQUE DENSITY IN ADULT PATIENTS WITH COGNITIVE IMPAIRMENT
- U-1498 METHOD OF TREATING PATIENTS WITH GASTRIC RETENTIVE DOSAGE FORM
- U-1499 MANAGEMENT OF ACUTE PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA
- U-1500 TESTOSTERONE REPLACEMENT THERAPY IN ADULT MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE; PRIMARY HYPOGONADISM (CONGENITAL OR ACQUIRED); HYPOGONADOTROPIC HYPOGONADISM (CONGENITAL OR ACQUIRED).
- U-1501 PROPHYLAXIS OF DEEP VEIN THROMBOSIS AND PULMONARY EMBOLISM
- U-1502 PROPHYLAXIS OF PULMONARY EMBOLISM
- U-1503 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLITZTIN IN COMBINATION WITH METFORMIN
- U-1504 USE OF OTEZLA (APREMILAST) FOR INHIBITING PDE4
- U-1505 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS
- U-1506 TREATMENT OF PATIENTS WITH GASTROINTESTINAL STROMAL TUMOR (GIST), INCLUDING BUT NOT LIMITED TO PATIENTS PREVIOUSLY TREATED WITH IMATINIB AND PATIENTS WITH GIST HAVING RESISTANCE TO A KIT TYROSINE KINASE INHIBITOR
- U-1507 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN
- U-1508 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING A PLURALITY OF COMPOSITE SUBUNITS AS CLAIMED
- U-1509 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING A GASTRIC ACID REDUCER
- U-1510 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT BY ORALLY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED.
- U-1511 TREATMENT OF HYPERTRIGLYCERIDEMIA
- U-1512 REDUCTION OF THROMBOTIC CARDIOVASCULAR EVENTS
- U-1513 TREATMENT OF HIV-1 INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS
- U-1514 MANAGEMENT OF BREAKTHROUGH PAIN IN PATIENTS WITH CANCER BY BUCCAL OR SUBLINGUAL ADMINISTRATION OF FENTANYL
- U-1515 METHOD OF TREATING IRRITABLE BOWEL SYNDROME WITH CONSTIPATION IN ADULT PATIENTS.
- U-1516 METHOD OF TREATING CHRONIC IDIOPATHIC CONSTIPATION IN ADULT PATIENTS.
- U-1517 TREATMENT OF BACTERIAL INFECTIONS USING A TWO-DOSE REGIMENT OF DALBAVANCIN.

**PATENT AND EXCLUSIVITY TERMS**

ADB 93 of 117

**PATENT USE**

- U-1518 MAINTAINING PUPIL SIZE BY PREVENTING INTRAOPERATIVE MIOSIS AND REDUCING POSTOPERATIVE OCULAR PAIN
- U-1519 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT WITH IRRITABLE BOWEL SYNDROME
- U-1520 METHOD FOR THE LONG TERM TREATMENT OF CHRONIC CONSTIPATION IN A HUMAN SUBJECT
- U-1521 MAINTENANCE TREATMENT OF OPIOID DEPENDENCE
- U-1522 TREATMENT OF TYPE 2 DIABETES MELLITUS IN A PATIENT, WHEREIN GLYCEMIC CONTROL (HBA1C < 7.0%) IS NOT ACHIEVABLE USING ONE OR MORE OF INSULIN, METFORMIN, PIOGLITAZONE, OR ROSIGLITAZONE
- U-1523 METHOD OF INDUCING TOPICAL ANESTHESIA IN THE EYE
- U-1524 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE
- U-1525 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS IN PATIENTS WITH NARCOLEPSY
- U-1526 THE TREATMENT OF PATIENTS WITH TRAVELERS' DIARRHEA (TD) OR THE REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) RECURRENCE
- U-1527 FOR THE TREATMENT OF OVERACTIVE BLADDER (OAB) WITH SYMPTOMS OF URGE URINARY INCONTINENCE, URGENCY, AND URINARY FREQUENCY
- U-1528 A METHOD OF LOWERING INTRAOCULAR PRESSURE
- U-1529 ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER (MDD)
- U-1530 USE OF ARIPIPRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION
- U-1531 METHOD FOR TRANSDERMAL DELIVERY OF TESTOSTERONE
- U-1532 METHOD OF TREATING EXCESSIVE DAYTIME SLEEPINESS AND/OR CATAPLEXY IN NARCOLEPSY PATIENTS WITH SODIUM OXYBATE WHEN DIVALPROEX SODIUM IS CONCOMITANTLY ADMINISTERED.
- U-1533 PULMONARY ADMINISTRATION OF PARTICLES COMPRISING A DIKETOPIPERAZINE AND INSULIN.
- U-1534 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH A DIKETOPIPERAZINE.
- U-1535 ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN COMPLEXED WITH MICROPARTICLES OF A DIKETOPIPERAZINE.
- U-1536 ADMINISTRATION OF A COMPOSITION COMPRISING A DIKETOPIPERAZINE AND INSULIN.
- U-1537 TREATMENT OF A PATIENT HAVING DIABETES MELLITUS WITH A PRANDIAL RAPID ACTING INSULIN.
- U-1538 ADMINISTRATION OF FDKP MICROPARTICLES COMPRISING INSULIN.
- U-1539 PULMONARY ADMINISTRATION OF AN INSULIN COMPOSITION COMPRISING FDKP AT THE BEGINNING OF A MEAL TO A PATIENT ALSO BEING TREATED WITH A LONG-ACTING INSULIN.
- U-1540 BUTRANS IS A PARTIAL OPIOID AGONIST PRODUCT INDICATED FOR THE MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.
- U-1541 TREATMENT OF PATIENTS WITH TUBEROUS SCLEROSIS COMPLEX (TSC) WHO HAVE SUBEPENDYMAL GIANT CELL ASTROCYTOMA (SEGA) THAT REQUIRES THERAPEUTIC INTERVENTION BUT CANNOT BE CURATIVELY RESECTED.
- U-1542 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA AND/OR NON-HODGKINS LYMPHOMA
- U-1543 TREATMENT OF A PATIENT BY ADMINISTERING THE FORMULATION RECITED IN CLAIM 1 OR CLAIM 23
- U-1544 TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY PERIPHERAL T-CELL LYMPHOMA (PTCL).
- U-1545 A METHOD OF TRANSDERMALLY DELIVERING TESTOSTERONE
- U-1546 FOR USE IN THE TREATMENT OF MALIGNANT HYPERTERMIA IN CONJUNCTION WITH APPROPRIATE SUPPORTIVE MEASURES AND FOR THE PREVENTION OF MALIGNANT HYPERTERMIA IN PATIENTS AT HIGH RISK.
- U-1547 TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), CHRONIC BRONCHITIS OR EMPHYSEMA
- U-1548 FOR THE LONG-TERM, ONCE-DAILY MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH COPD, INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA, ALSO TO REDUCE EXACERBATIONS OF COPD IN PATIENTS WITH A HISTORY OF EXACERBATIONS

**PATENT AND EXCLUSIVITY TERMS**

ADB 94 of 117

**PATENT USE**

- U-1549 FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA
- U-1550 METHOD OF TREATING METASTATIC PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS.
- U-1551 METHOD OF TREATING PAPILLARY RENAL CELL CARCINOMA WITH TEMSIROLIMUS, IN THE ABSENCE OF INTERFERON ALPHA.
- U-1552 FOR HEALING OF ALL GRADES OF EROSIVE ESOPHAGITIS (EE)
- U-1553 TO MAINTAIN HEALING OF EE AND RELIEF OF HEARTBURN
- U-1554 FOR THE TREATMENT OF HEARTBURN ASSOCIATED WITH SYMPTOMATIC NON-EROSIVE GASTROESOPHAGEAL DISEASE (GERD)
- U-1555 MANAGEMENT OF MODERATE TO SEVERE PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.
- U-1556 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE.
- U-1557 A METHOD OF TESTOSTERONE REPLACEMENT THERAPY COMPRISING THE STEP OF NASALLY ADMINISTERING TO A PATIENT IN NEED OF SUCH TREATMENT AN EFFECTIVE AMOUNT OF TESTOSTERONE GEL FORMULATION.
- U-1558 FOR THE TREATMENT OF PATIENTS WITH RELAPSED FOLLICULAR B-CELL NON-HODGKIN LYMPHOMA OR [RELAPSED] SMALL LYMPHOCYTIC LYMPHOMA
- U-1559 INDICATED FOR THE ONCE-DAILY MAINTENANCE TREATMENT OF ASTHMA AS PROPHYLACTIC THERAPY IN PATIENTS AGED 12 YEARS OF AGE AND OLDER
- U-1560 A METHOD OF DISRUPTING LEUKOCYTE FUNCTION, INCLUDING AS AN INHIBITOR OF PI3KDELTA KINASE
- U-1561 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIATIC ARTHRITIS
- U-1562 TREATMENT OF PATIENTS WITH HEPATIC ENCEPHALOPATHY (HE)
- U-1563 A METHOD OF TRANSDERMAL ADMINISTRATION OF A PHYSIOLOGICALLY ACTIVE AGENT TO A SUBJECT.
- U-1564 A METHOD OF TREATING GAUCHER'S DISEASE
- U-1565 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE
- U-1566 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER
- U-1567 METHOD OF TREATING, AS INITIAL LOADING DOSE FOR MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL ONSET-SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS OR OLDER
- U-1568 METHOD OF TREATING, AS MONOTHERAPY OR ADJUNCTIVE THERAPY, PARTIAL-ONSET SEIZURES IN A PATIENT WITH EPILEPSY AGED 17 YEARS AND OLDER WHEN ORAL ADMINISTRATION IS TEMPORARILY NOT FEASIBLE
- U-1569 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS
- U-1570 TREATMENT OF BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS USING A SINGLE DOSE
- U-1571 TREATMENT OF GAUCHER DISEASE TYPE 1
- U-1572 TREATMENT OF HUMAN IMMUNODEFICIENCY VIRUS (HIV) INFECTION.
- U-1573 USE OF RUXOLITINIB (JAKAFI) FOR INHIBITING JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2.
- U-1574 A METHOD OF CATALYZING THE HYDROLYSIS OF GLUCOCEREBROSIDE TO GLUCOSE AND CERAMIDE.
- U-1575 PATIENTS WITH SEVERE APLASTIC ANEMIA WHO HAVE HAD AN INSUFFICIENT RESPONSE TO IMMUNOSUPPRESSIVE THERAPY
- U-1576 TREATMENT OF LEUKEMIA
- U-1577 CONTROL OF SERUM PHOSPHOROUS LEVELS
- U-1578 TREATMENT OF ACUTE OTITIS MEDIA
- U-1579 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER
- U-1580 USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER WHO HAD RECEIVED PRIOR DOCETAXEL

**PATENT AND EXCLUSIVITY TERMS**

ADB 95 of 117

**PATENT USE**

## CHEMOTHERAPY

- U-1581 IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA.
- U-1582 TREATMENT OF UNRESECTABLE OR METASTATIC MELANOMA
- U-1583 FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY
- U-1584 USE OF NALTREXONE AND BUPROPION IN A LAYERED FORMULATION FOR CHRONIC WEIGHT MANAGEMENT FOR AFFECTING WEIGHT LOSS
- U-1585 USE OF NALTREXONE AND BUPROPION BASED ON AN ESCALATING DOSE SCHEDULE
- U-1586 FOR EFFECT ON BLOOD GLUCOSE PARAMETERS IN PATIENTS WITH INSULIN RESISTANCE
- U-1587 SINGLE-DOSE INFILTRATION INTO THE SURGICAL SITE TO PRODUCE POSTSURGICAL ANALGESIA.
- U-1588 THE TREATMENT OF PATIENTS WITH METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (CRPC).
- U-1589 METHOD OF USE FOR REDUCING BLOOD PHENYLALANINE LEVELS IN A HUMAN SUFFERING FROM HYPERPHENYLALANINEMIA
- U-1590 KUVAN IS INDICATED TO REDUCE BLOOD PHENYLALANINE LEVELS IN PATIENTS WITH HYPERPHENYLALANINEMIA
- U-1591 TREATMENT OF ASTHMA IN PATIENTS AGED 12 YEARS AND OLDER
- U-1592 TO REDUCE SERUM PHOSPHATE IN PATIENTS WITH END STAGE RENAL DISEASE
- U-1593 MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA, AND REDUCTION OF EXACERBATIONS IN COPD PATIENTS.
- U-1594 DILATION OF THE PUPIL
- U-1595 USE OF OTEZLA (APREMILAST) FOR THE TREATMENT OF PSORIASIS
- U-1596 LAMICTAL IS AN ANTIEPILEPTIC DRUG (AED) INDICATED FOR: EPILEPSY-ADJUNCTIVE THERAPY IN PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE: (1.1) PARTIAL SEIZURES PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
- U-1597 TREATMENT OF DIABETIC MACULAR EDEMA
- U-1598 METHOD OF ADMINISTRATION OF CONTROLLED RELEASE OXYMORPHONE
- U-1599 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANTI INFLAMMATORY, ANALGESIC, AND ANTIPIRETIC ACTIVITY
- U-1600 DOSAGE MODIFICATION FOLLOWING Elevated LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- U-1601 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- U-1602 METHOD OF ADMINISTERING PIRFENIDONE CAPSULES TO TREAT A FIBROTIC CONDITION
- U-1603 METHOD FOR ADMINISTERING PIRFENDONE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-1604 METHOD FOR ADMINISTERING PIRFENDONE TO REDUCE DRUG INTERACTIONS WITH A STRONG INHIBITOR OF CYP1A2
- U-1605 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING OR AVOIDING ANOTHER STRONG CYP1A2 INDUCER
- U-1606 METHOD FOR ADMINISTERING PIRFENIDONE WHILE AVOIDING OR DISCONTINUING CONCOMITANT USE OF A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME INVOLVED IN PIRFENIDONE METABOLISM
- U-1607 METHOD OF ADMINISTERING A DOSAGE FORM THAT INCLUDES A GRANULATE FORMULATION OF PIRFENIDONE TO TREAT A FIBROTIC CONDITION
- U-1608 DOSE ESCALATION OVER 14 DAYS FOR TREATMENT OF A FIBROSIS CONDITION
- U-1609 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING Elevated LIVER ENZYMES IN TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS
- U-1610 CONTINUED DOSING OR DOSAGE MODIFICATION FOLLOWING ELVATED LIVER ENZYMES IN USE OF PIRFENIDONE
- U-1611 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY DISCONTINUING SMOKING OR BY DISCONTINUING A STRONG CYP1A2 INDUCER
- U-1612 METHOD FOR ADMINISTERING PIRFENIDONE TO AVOID REDUCED EFFICACY BY AVOIDING SMOKING OR BY AVOIDING ANOTHER STRONG CYP1A2 INDUCER

**PATENT AND EXCLUSIVITY TERMS**

ADB 96 of 117

**PATENT USE**

- U-1613 DOSAGE MODIFICATION IN TREATMENT WITH PIRFENIDONE TO REDUCE DRUG INTERACTIONS WITH CIPROFLOXACIN
- U-1614 USE OF TOPICAL DICLOFENAC SODIUM FOR TREATING PAIN
- U-1615 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL
- U-1616 NASAL ADMINISTRATION OF A TESTOSTERONE GEL TO A PATIENT TO TREAT THE PATIENT FOR A CONDITION ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-1617 METHOD OF TREATING MEDULLARY THYROID CANCER
- U-1618 A METHOD OF TREATING A PATIENT SUFFERING FROM A PAIN ASSOCIATED SLEEP DISTURBANCE COMPRISING ADMINISTERING A LIQUID COMPOSITION FORMULATED INSIDE A SOFT GEL CAPSULE, AS CLAIMED, TO THE PATIENT
- U-1619 TREATMENT OF IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1620 METHOD OF TREATMENT OF IRON-RELATED CONDITIONS WITH AT LEAST 0.6 GRAMS OF ELEMENTAL IRON VIA AN IRON CARBOHYDRATE COMPLEX, WITH A SUBSTANTIALLY NON-IMMUNOGENIC CARBOHYDRATE COMPONENT, IN ABOUT 15 MINUTES OR LESS.
- U-1621 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A COMPLEXING AGENT.
- U-1622 FOR THE TREATMENT OF POLYCYTHEMIA VERA
- U-1623 USE OF EXENATIDE MAY RESULT IN REDUCTION IN APPETITE.
- U-1624 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA, ADVANCED RENAL CELL CARCINOMA, OR DIFFERENTIATED THYROID CARCINOMA.
- U-1625 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE POOR METABOLIZERS OF CYP2D6
- U-1626 A METHOD OF TREATING OR PREVENTING OCULAR PAIN AND BURNING
- U-1627 TREATMENT OF ACUTE UNCOMPLICATED INFLUENZA IN ADULTS
- U-1628 METHOD OF TREATING DISORDERS WITH AN ETIOLOGY COMPRISING OR ASSOCIATED WITH EXCESS GH-SECRETION
- U-1629 METHOD OF TREATING ACROMEGALY
- U-1630 TREATMENT IN COMBINATION WITH A CORTICOID SUCH AS PREDNISONE OF PROSTATE CANCER PREVIOUSLY TREATED WITH DOCETAXEL
- U-1631 TREATMENT OF INFLAMMATORY LESIONS OF ROSACEA.
- U-1632 TREATMENT OF SCHIZOPHRENIA, WITH EFFICACY IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA
- U-1633 USE OF ARIPIPRAZOLE IN EXTENDED RELEASE INJECTABLE SUSPENSION IN TREATING ACUTE EPISODES OF SCHIZOPHRENIA
- U-1634 TREATMENT OF BRCA MUTATED OVARIAN CANCER USING PARP INHIBITOR
- U-1635 USE OF RITONAVIR AS A POTENT CYP3A INHIBITOR TO INCREASE PLASMA DRUG CONCENTRATION OF PARITAPREVIR AND OVERALL DRUG EXPOSURE FOR TREATMENT OF HCV INFECTION
- U-1636 USE OF DASABUVIR TO INHIBIT VIRAL REPLICATION FOR THE TREATMENT OF HCV INFECTION.
- U-1637 TREATMENT OF HCV INFECTION USING PARITAPREVIR, OMBITASVIR, RITONAVIR, AND DASABUVIR WITH RIBAVIRIN.
- U-1638 TREATMENT OF HCV INFECTION USING PARITAPREVIR
- U-1639 USE OF NALTREXONE AND BUPROPION IN EXTENDED-RELEASE FORM FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY
- U-1640 TREATMENT OF MODERATE TO SEVERE CHRONIC PAIN BY ADMINISTERING AN INTACT COMPOSITION AS CLAIMED
- U-1641 MEMANTINE HCL/DONEPEZIL HCL COMBINATION FOR THE TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE
- U-1642 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND FOR WHOM METFORMIN THERAPY IS INAPPROPRIATE BY ADMINISTERING LINAGLITPIN
- U-1643 TREATING CUSHING'S SYNDROME
- U-1644 TREATMENT OF OVERACTIVE BLADDER BY APPLICATION OF OXYBUTYNIN CHLORIDE GEL TO SKIN

**PATENT AND EXCLUSIVITY TERMS**

ADB 97 of 117

**PATENT USE**

- U-1645 TREATMENT OF PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1646 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1647 TREATMENT OF PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1648 TREATMENT OF PATIENTS WITH PARKINSON'S DISEASE, POST-ENCEPHALITIC PARKINSONISM, AND PARKINSONISM THAT MAY FOLLOW CARBON MONOXIDE INTOXICATION OR MANGANESE INTOXICATION
- U-1649 TREATMENT OF POST-ENCEPHALITIC PARKINSONISM
- U-1650 TREATMENT OF WALDENSTROM'S MACROGLOBULINEMIA
- U-1651 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPITIN IN COMBINATION WITH EMPAGLIFLOZIN
- U-1652 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPITIN IN COMBINATION WITH EMPAGLIFLOZIN AND METFORMIN
- U-1653 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPITIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT METFORMIN)
- U-1654 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPITIN IN COMBINATION WITH EMPAGLIFLOZIN (WITH OR WITHOUT INSULIN OR A SULFONYLUREA)
- U-1655 A METHOD TO ACCELERATE THE TIME TO GASTROINTESTINAL RECOVERY BY ADMINISTERING ABOUT 12 MG OF ALVIMOPAN TO THE PATIENT FROM ABOUT 30 TO 60 MINUTES PRIOR TO SURGERY
- U-1656 METHOD OF IRON ADMINISTRATION TO TREAT PATIENTS IN NEED OF IRON REPLACEMENT
- U-1657 METHOD FOR PROVIDING POST COITAL CONTRACEPTION TO A WOMAN BY ADMINISTERING ABOUT 30 MG OF ULIPRISTAL ACETATE WITHIN ABOUT 120 HOURS AFTER INTERCOURSE, WHEREIN THE WOMAN IS OVERWEIGHT HAVING A BMI OF 25 TO 29.99
- U-1658 TREATMENT OF ER-POSITIVE, HER2-NEGATIVE ADVANCED BREAST CANCER IN COMBINATION WITH LETROZOLE AS INITIAL ENDOCRINE-BASED THERAPY FOR METASTATIC DISEASE IN POSTMENOPAUSAL WOMEN
- U-1659 MANAGEMENT OF PAIN
- U-1660 TREATMENT OF HIV-1 INFECTION IN ADULTS WITH NO DARUNAVIR RESISTANCE-ASSOCIATED SUBSTITUTIONS
- U-1661 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS ALSO TAKING LOW DOSE ASPIRIN
- U-1662 A METHOD OF TREATING OCULAR PAIN
- U-1663 TREATMENT OF HIV-1 INFECTION
- U-1664 TREATMENT OF BACTERIAL VAGINOSIS WITH METRONIDAZOLE GEL
- U-1665 METHOD OF TREATING ATTENTION DEFICIT HYPERACTIVITY DISORDER BY ADMINISTERING THE COMPOSITION OF CLAIM 1
- U-1666 PALLIATIVE TREATMENT OF PROSTATE CANCER
- U-1667 TREATMENT OF ALLERGIC RHINITIS, INCLUDING SEASONAL ALLERGIC RHINITIS
- U-1668 METHOD OF TREATING DEPRESSION OR MAJOR DEPRESSIVE DISORDER
- U-1669 TREATMENT OF MULTIPLE MYELOMA, IN COMBINATION WITH BORTEZOMIB AND DEXAMETHASONE
- U-1670 NATROBA TOPICAL SUSPENSION IS A PEDICULICIDE INDICATED FOR THE TOPICAL TREATMENT OF HEAD LICE INFESTATION IN PATIENTS SIX (6) MONTHS OF AGE AND OLDER.
- U-1671 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH CONJUNCTIVITIS
- U-1672 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTION
- U-1673 TREATMENT OF COMPLICATED URINARY TRACT INFECTION, INCLUDING PYELONEPHRITIS
- U-1674 DOSAGE MODIFICATION TO REDUCE RISKS ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE
- U-1675 USE OF TROKENDI XR FOR THE TREATMENT OF EPILEPSY
- U-1676 METHODS FOR TREATING BACTERIAL INFECTIONS
- U-1677 TREATMENT OF IDIOPATHIC PULMONARY FIBROSIS (IPF)

**PATENT AND EXCLUSIVITY TERMS**

ADB 98 of 117

**PATENT USE**

- U-1678 FOR THE TREATMENT OF PATIENTS WITH CLL, FL, OR SLL
- U-1679 TREATMENT OF ACUTE OTITIS EXTERNA
- U-1680 TREATMENT OF OCULAR ITCHING ASSOCIATED WITH ALLERGIC CONJUNCTIVITIS
- U-1681 TREATMENT OF PATIENTS WITH PROGRESSIVE NEUROENDOCRINE TUMORS OF PANCREATIC ORIGIN (PNET) THAT ARE UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC
- U-1682 TREATMENT OF BACTERIAL VAGINOSIS
- U-1683 TREATMENT FOR CHRONIC LYMPHOCYTIC LEUKEMIA WITH 17P DELETION
- U-1684 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA
- U-1685 DOSAGE MODIFICATION TO REDUCE THE RISK ASSOCIATED WITH QT PROLONGATION NOT INDUCED BY OTHER DRUGS DURING TREATMENT WITH ILOPERIDONE
- U-1686 A METHOD TO REDUCE WITHDRAWAL SYMPTOMS, INCLUDING NICOTINE CRAVING, ASSOCIATED WITH SMOKING CESSATION
- U-1687 TREATMENT OF HCV INFECTION USING OMBITASVIR
- U-1688 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD
- U-1689 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD
- U-1690 METHOD FOR REDUCTION OF SUBMENTAL FAT
- U-1691 INDICATED FOR THE ONCE-DAILY INHALED TREATMENT FOR ASTHMA IN ADULTS AGED 18 YEARS AND OLDER
- U-1692 METHOD FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN AN INDIVIDUAL WHO DOES NOT HAVE SEVERE RENAL IMPAIRMENT OR ESRD
- U-1693 METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS
- U-1694 A METHOD FOR TREATING HEART FAILURE IN A HUMAN USING A CRYSTALLINE FORM OF IVABRADINE HYDROCHLORIDE
- U-1695 METHOD FOR TREATING THYROID CARCINOMA INCLUDING DIFFERENTIATED THYROID CANCER
- U-1696 TREATMENT OF UNRESECTABLE HEPATOCELLULAR CARCINOMA
- U-1697 PULMONARY ADMINISTRATION OF A COMPOSITION COMPRISING INSULIN BOUND TO A DIKETOPIPERAZINE.
- U-1698 PROPHYLAXIS OF INVASIVE ASPERGILLUS AND CANDIDA INFECTIONS
- U-1699 A METHOD FOR TREATING ACUTE LYMPHOBLASTIC LEUKEMIA
- U-1700 A METHOD FOR TREATING PHILADELPHIA CHROMOSOME POSITIVE ACUTE LYMPHOBLASTIC LEUKEMIA
- U-1701 A METHOD FOR TREATING LEUKEMIA RESULTING FROM A MUTATION IN THE BCR-ABL KINASE DOMAIN
- U-1702 TREATMENT OF COPD
- U-1703 TREATMENT OF RESPIRATORY COMPLAINTS
- U-1704 USE FOR TREATMENT IN PATIENTS WITH DIABETES
- U-1705 USE FOR TREATMENT IN PATIENTS WITH HYPERGLYCEMIA
- U-1706 TREATMENT OF TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGogue WHEREIN THE COMBINED THERAPEUTIC EFFECT IS GREATER THAN THE ADDITIVE EFFECT OF ADMINISTERING EACH AGENT ALONE
- U-1707 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS AND SYMPTOMS THEREOF.
- U-1708 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) IN ADULTS.
- U-1709 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE).
- U-1710 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH FLUVOXAMINE
- U-1711 FOR THE TREATMENT OF PATIENTS WITH CLL, FL OR SLL
- U-1712 MEKENIST IN COMBINATION WITH DABRAFENIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA

**PATENT AND EXCLUSIVITY TERMS**

ADB 99 of 117

**PATENT USE**

- U-1713 TAFINLAR IN COMBINATION WITH TRAMETINIB FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA
- U-1714 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 6 YEARS AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1715 P2Y12 PLATELET INHIBITOR FOR USE AS ADJUNCT TO PERCUTANEOUS CORONARY INTERVENTION TO REDUCE RISK OF VARIOUS DISEASES/CONDITIONS IN PATIENTS NOT TREATED WITH A P2Y12 PLATELET INHIBITOR AND NOT GIVEN A GLYCOPROTEIN IIB/IIIA INHIBITOR
- U-1716 TREATMENT OF COUGH AND SYMPTOMS ASSOCIATED WITH UPPER RESPIRATORY ALLERGIES OR A COMMON COLD WITH CODEINE PHOSPHATE AND CHLORPHENIRAMINE MALEATE ORALLY ADMINISTERED EXTENDED RELEASE TABLETS
- U-1717 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO ARE HOMOZYGOUS FOR THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE
- U-1718 METHOD OF TREATING CYSTIC FIBROSIS IN PATIENTS WHO HAVE THE F508DEL MUTATION IN THE CYSTIC FIBROSIS TRANSMEMBRANE CONDUCTANCE REGULATOR (CFTR) GENE.
- U-1719 ACUTE TREATMENT OF MIGRAINE
- U-1720 METHOD OF PROVIDING A THERAPEUTICALLY EFFECTIVE AND STABLE MEDIAN BLOOD PLASMA LEVEL OF LEVODOPA
- U-1721 USE OF RUXOLITINIB (JAKAFI) FOR BLOCKING SIGNAL TRANSDUCTION OF JANUS ASSOCIATED KINASES (JAKS) JAK1 AND/OR JAK2
- U-1722 TREATMENT OF BASAL CELL CARCINOMA
- U-1723 TREATMENT OF HEART FAILURE
- U-1724 METHOD OF INHIBITING HEPATITIS C VIRUS
- U-1725 METHOD OF INHIBITING HEPATITIS C VIRUS WITH DAKLINZA AND AT LEAST ONE ADDITIONAL COMPOUND HAVING ANTI-HCV ACTIVITY
- U-1726 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH CORONARY HEART DISEASE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1727 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA
- U-1728 REDUCTION IN RISK OF HOSPITALIZATION IN PATIENTS WITH STABLE NYHA CLASS III HEART FAILURE AND A HISTORY OF PAROXYSMAL OR PERSISTENT AF AND WITH ONE OR MORE RISK FACTORS BY ADMINISTRATION TWICE A DAY WITH MORNING AND EVENING MEALS
- U-1729 REDUCE THE RISK OF RECURRENT DEEP VEIN THROMBOSIS (DVT)
- U-1730 REDUCE THE RISK OF RECURRENT PULMONARY EMBOLISM
- U-1731 TEMPORARY RELIEF OF MINOR ACHES AND PAINS
- U-1732 TEMPORARY REDUCTION OF FEVER
- U-1733 TREATMENT/PREVENTION OF CARDIOVASCULAR DISEASE
- U-1734 USE OF FLIBANSERIN OR A PHARMACEUTICALLY ACCEPTABLE ACID ADDITION SALT THEREOF TO TREAT HYPOACTIVE SEXUAL DESIRE DISORDER (HSDD)
- U-1735 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER WITH INTRAVENOUS IBUPROFEN SUCH THAT MEAN ARTERIAL BLOOD PRESSURE DOES NOT INCREASE THE DOSAGE INTERVAL
- U-1736 TREATMENT OF THROMBOCYTOPENIA IN ADULT AND PEDIATRIC PATIENTS 1 YEAR AND OLDER WITH CHRONIC IMMUNE (IDIOPATHIC) THROMBOCYTOPENIA (ITP)
- U-1737 METHOD OF TREATING SCHIZOPHRENIA BY ADMINISTERING ILOPERIDONE TO A PATIENT BY REDUCING THE DOSE IN PATIENTS WHO ARE BEING TREATED WITH FLUOXETINE
- U-1738 TREATMENT OF IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)
- U-1739 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND -THE-CLOCK, LONG-TERM OPIOID TREATMENT, INCLUDING NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-1740 IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS FOR THE TREATMENT OF HIV-1 INFECTION IN TREATMENT-NAIVE PATIENTS WITH HIV-1 RNA LESS THAN OR EQUAL TO 100,000 AT THE START OF THEARPY
- U-1741 PREVENTION OF DELAYED NAUSEA AND VOMITING ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY

**PATENT AND EXCLUSIVITY TERMS**

ADB 100 of 117

**PATENT USE**

- U-1742 ROLAPITANT IS APPROVED FOR THE PREVENTION OF DELAYED NAUSEA AND VOMITING (I.E., EMESIS) ASSOCIATED WITH EMETOGENIC CANCER CHEMOTHERAPY
- U-1743 FOR THE PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY
- U-1744 PREVENTION OF POST-OPERATIVE NAUSEA AND VOMITING
- U-1745 FOR THE TREATMENT OF PATIENTS WITH WALDENSTROM'S MACROGLOBULINEMIA
- U-1746 MONOTHERAPY OR ADJUNCTIVE THERAPY FOR TREATMENT OF PARTIAL-ONSET SEIZURES AND APPROVED IN PATIENTS WITH EPILEPSY
- U-1747 FOR CLAIMS 1-3, 6-13, 16-24 AND 26-32: METHOD OF TREATING ADHD
- U-1748 FOR CLAIMS 1-4, 6-14, 16-24 AND 26-32: METHOD OF TREATING ADHD IN CHILDREN 6 YEARS OF AGE AND OLDER AND ADOLESCENTS
- U-1749 ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPO极 I DISORDER
- U-1750 TREATMENT OF SCHIZOPHRENIA AND/OR ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPO极 I DISORDER WITH CARIPRAZINE
- U-1751 TREATMENT OF PATIENTS WITH METASTATIC COLORECTAL CANCER WHO HAVE BEEN PREVIOUSLY TREATED WITH FLUOROPYRIMIDINE-, OXALIPLATIN- AND IRINOTECAN-BASED CHEMOTHERAPY, AN ANTI-VEGF BIOLOGICAL THERAPY, AND IF RAS WILD-TYPE, AN ANTI-EGFR THERAPY
- U-1752 PROPHYLAXIS OF ORGAN REJECTION
- U-1753 TREATMENT OF HCV INFECTION USING DASABUVIR
- U-1754 FOR THE TREATMENT OF PULMONARY HYPERTENSION (PAH) IN COMBINATION WITH Tadalafil
- U-1755 FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHOSE TUMORS HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS
- U-1756 METHODS OF TREATING PAIN, INFLAMMATION AND/OR FEVER IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF
- U-1757 INHIBITION ON PI3K KINASE
- U-1758 METHOD OF TREATING ALLERGIC REACTION VIA INJECTION
- U-1759 METHOD OF REVERSING THE ANTICOAGULANT EFFECT OF DABIGATRAN USING IDARUCIZUMAB
- U-1760 RISK-REDUCTION OF NSAID GASTRIC ULCER IN PATIENTS REQUIRING CHRONIC NSAID TREATMENT
- U-1761 PLAQUE PSORIASIS
- U-1762 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1763 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1764 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1765 USE OF BELVIQ (LORCASERIN HYDROCHLORIDE) FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1766 TREATMENT OF HYPERKALEMIA
- U-1767 USE OF CALCIPOTRIENE FOAM FOR THE TOPICAL TREATMENT OF PLAQUE PSORIASIS IN PATIENTS AGED 18 YEARS AND OLDER
- U-1768 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN
- U-1769 TREATMENT OF PAIN BY TRANSMUCOSAL DELIVERY OF BUPRENORPHINE
- U-1770 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN NEGATIVE SYMPTOMS AND/OR COGNITIVE DYSFUNCTION OF SCHIZOPHRENIA
- U-1771 ADMINISTRATION OF REMODULIN DILUTED FOR INTRAVENOUS INFUSION WITH STERILE WATER FOR INJECTION OR 0.9% SODIUM CHLORIDE INJECTION PRIOR TO ADMINISTRATION
- U-1772 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH EMPAGLIFLOZIN
- U-1773 LONG - TERM MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)

**PATENT AND EXCLUSIVITY TERMS**

ADB 101 of 117

**PATENT USE**

- U-1774 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE
- U-1775 USE OF A LOTION CONTAINING HALOBETASOL PROPIONATE FOR THE TREATMENT OF CORTICOSTEROID-RESPONSIVE DERMATOSES INCLUDING PSORIASIS
- U-1776 METHOD OF USING COBIMETINIB FOR THE TREATMENT OF MELANOMA
- U-1777 TREATMENT OF PATIENTS WITH METASTATIC EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC), WHO HAVE PROGRESSED ON OR AFTER EGFR TKI THERAPY
- U-1778 METHOD FOR TREATING MULTIPLE MYELOMA
- U-1779 METHOD FOR TREATING MULTIPLE MYELOMA WITH ONE OR MORE OTHER THERAPEUTIC AGENTS
- U-1780 METHOD FOR TREATING CANCER, INCLUDING MULTIPLE MYELOMA
- U-1781 RISK-REDUCTION OF NSAID-ASSOCIATED GASTRIC ULCER IN PATIENTS REQUIRING NSAID TREATMENT
- U-1782 FOR HEAD LICE INFESTATIONS
- U-1783 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM AS CLAIMED
- U-1784 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM TRIHYDRATE AS CLAIMED
- U-1785 METHOD OF TREATING FREQUENT HEARTBURN BY ADMINISTERING AN ESOMEPRAZOLE MAGNESIUM FORMULATION AS CLAIMED
- U-1786 TREATMENT OF PATIENTS WITH RELAPSING FORMS OF MULTIPLE SCLEROSIS WHILE MANAGING THE RISK OF TERIFLUONOMIDE AND ROSUVASTATIN INTERACTION BY LIMITING THE ROSUVASTATIN DOSE TO NO MORE THAN 10MG AND/OR ADMINISTERING ABOUT HALF THE NORMAL DOSE
- U-1787 TREATMENT OF EXOCRINE PANCREATIC INSUFFICIENCY
- U-1788 TREATMENT OF PATIENT HAVING DIABETES MELLITUS VIA ORAL INHALATION OF FDNP MICROPARTICLES COMPRISING INSULIN
- U-1789 METHOD OF ADMINISTERING AN ETHANOL-FREE TAXANE LIQUID NANODISPERSION FORMULATION TO A SUBJECT COMBINING THE FORMULATION WITH AN AQUEOUS MEDIUM TO PROVIDE AN ETHANOL-FREE TAXANE DILUTED SOLUTION
- U-1790 FOR USE IN TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) AND/OR NON-HODGKIN'S LYMPHOMA
- U-1791 EMERGENCY TREATMENT OF ADULT & PEDIATRIC PATIENTS FOLLOWING FLUOROURACIL OR CAPECITABINE OVERDOSE, OR WHO EXHIBIT EARLY-ONSET, SEVERE OR LIFE-THREATENING CARDIAC OR CNS TOXICITY OR UNUSUALLY SEVERE ADVERSE REACTIONS WITHIN 96 HOURS
- U-1792 TREATMENT OF OTIC INFECTION OR INFLAMMATION
- U-1793 TREATMENT OF PEDIATRIC PATIENTS WITH OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT
- U-1794 REVERSAL OF DRUG-INDUCED NEUROMUSCULAR BLOCK
- U-1795 REVERSAL OF NEUROMUSCULAR BLOCKAGE INDUCED BY ROCURONIUM BROMIDE OR VECURONIUM BROMIDE
- U-1796 TOPICAL TREATMENT OF INFLAMMATORY PAPULES AND PUSTULES OF MILD TO MODERATE ROSACEA
- U-1797 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING SELEXIPAG
- U-1798 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING SELEXIPAG IN COMBINATION WITH THE ENDOTHELIN RECEPTOR ANTAGONIST MACITENTAN
- U-1799 METHOD OF INCREASING GROWTH OF HAIR INCLUDING EYELASHES
- U-1800 A METHOD OF TREATING OCULAR PAIN AND/OR ENHANCING OCULAR COMFORT
- U-1801 REDUCTION OF SERUM URIC ACID LEVELS
- U-1802 TREATMENT OF GOUT
- U-1803 TREATMENT OF HYPERURICEMIA
- U-1804 ACHIEVING A THERAPEUTIC BENEFIT IN A SUBJECT WITH GOUT
- U-1805 USE OF DEXLANSOPRAZOLE IN PATIENTS TAKING CLOPIDOGREL WITHOUT MEANINGFUL CYP2C19 INTERACTIONS

**PATENT AND EXCLUSIVITY TERMS**

ADB 102 of 117

**PATENT USE**

- U-1806 COADMINISTERING WITH ALLOPURINOL TO REDUCE SERUM URIC ACID (SUA) BELOW 4 MG/DL; BELOW 6MG/DL IN PATIENTS HAVING URIC ACID DEPOSITS; AND/OR BELOW 6MG/DL WITH SUA INTRADAY CHANGE MORE THAN 50% AND/OR ADVERSE EVENT RATE LESS THAN 15%
- U-1807 TREATMENT OF PEDIATRIC PATIENTS 8 TO 17 YEARS OF AGE WITH HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HEFH)
- U-1808 USE OF NALTREXONE AND BUPROPION FOR CHRONIC WEIGHT MANAGEMENT FOR TREATING OVERWEIGHT OR OBESITY IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER
- U-1809 METHOD OF DRUG DELIVERY VIA THE NASAL CAVITY
- U-1810 TREATMENT OF PAIN IN PATIENTS WITH HEPATIC IMPAIRMENT
- U-1811 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATIONS AFTER CONFIRMING THE PRESENCE OF BRAF V600E MUTATION
- U-1812 TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC LIPOSARCOMA
- U-1813 TREATMENT OF PATIENTS INFECTED WITH HEPATITIS C VIRUS
- U-1814 METHOD OF TREATING GLAUCOMA OR ELEVATED INTRAOCULAR PRESSURE
- U-1815 TREATMENT OF PARTIAL-ONSET SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
- U-1816 TREATMENT OF A UREA CYCLE DISORDER
- U-1817 PEDIATRIC USE AGES 1 MONTH TO 2 YEARS, GERD AND EROSIONAL ESOPHAGITIS
- U-1818 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH LETROZOLE AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN, OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-1819 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE
- U-1820 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION BY ADMINISTERING A PHARMACEUTICAL COMPOSITION COMPRISING MACITENTAN AND A POLYSORBATE, WHEREIN THE POLYSORBATE REPRESENTS 0.1 TO 3% OF THE WEIGHT OF SAID PHARMACEUTICAL COMPOSITION
- U-1821 METHOD FOR CONTRACEPTION TO A WOMAN COMPRISING ADMINISTERING TO THE WOMAN 30MG OF ULIPRISTAL ACETATE MORE THAN 72 HOURS AND UP TO 120 HOURS AFTER AN UNPROTECTED INTERCOURSE
- U-1822 TREATMENT OF SCHIZOPHRENIA OR BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA AND/OR BIPOLAR DISORDER
- U-1823 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY COMPENSATING LONG-TERM SENSITIVITY DRIFT OF ELECTROCHEMICAL GAS SENSORS USED IN SYSTEMS FOR DELIVERING THERAPEUTIC NITRIC OXIDE TO A PATIENT
- U-1824 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY VERIFYING GAS INFORMATION OF NITRIC OXIDE PRIOR TO DELIVERY TO PATIENT
- U-1825 METHOD OF USING VISMODEGIB TO TREAT CANCER IN A MAMMAL
- U-1826 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCOCLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-1827 A METHOD OF PROVIDING A SUBJECT WITH THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET ACCORDING TO CLAIM 1
- U-1828 INCREASING MEAN ARTERIAL BLOOD PRESSURE IN ADULT PATIENTS WITH HYPOTENSION ASSOCIATED WITH SEPTIC SHOCK
- U-1829 EMERGENCY TREATMENT OF ALLERGIC REACTIONS (TYPE I), INCLUDING ANAPHYLAXIS
- U-1830 INDUCTION AND MAINTENANCE OF MYDRIASIS DURING INTRAOCULAR SURGERY
- U-1831 METHOD OF TREATING PULMONARY ARTERIAL HYPERTENSION COMPRISING ADMINISTERING A CRYSTALLINE FORM OF SELEXIPAG
- U-1832 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
- U-1833 REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-1834 TREATMENT OF POSTOPERATIVE INFLAMMATION AND PREVENTION OF OCULAR PAIN IN

**PATENT AND EXCLUSIVITY TERMS**

ADB 103 of 117

**PATENT USE**

## PATIENTS UNDERGOING CATARACT SURGERY

- U-1835 TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) WITH 17P DELETION, AS DETECTED BY AN FDA APPROVED TEST, WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-1836 TREATMENT OF HEREDITARY TYROSINEMIA TYPE 1 (HT-1) IN COMBINATION WITH DIETARY RESTRICTION OF TYROSINE AND PHENYLALANINE
- U-1837 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN ALONE OR IN COMBINATION WITH INSULIN, METFORMIN, A THIAZOLIDINEDIONE, GLYBURIDE OR METFORMIN PLUS A SULFONYLUREA
- U-1838 METHOD FOR TREATING TYPE II DIABETES MELLITUS BY ADMINISTERING SAXAGLIPTIN IN COMBINATION WITH METFORMIN
- U-1839 COMPOSITION AND METHOD FOR PROVIDING A REDUCTION IN SIDE EFFECTS FOR HUMAN PATIENTS IN NEED OF ACETYL CYSTEINE THERAPY
- U-1840 TREATMENT OF HCV INFECTION USING PARITAPREVIR, OMBITASVIR, RITONAVIR, AND DASABUVIR, WITHOUT RIBAVIRIN
- U-1841 USE IN THE LONG-TERM, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD)
- U-1842 METHOD OF TREATING EPILEPSY
- U-1843 TREATMENT OF PSYCHOSIS
- U-1844 TREATMENT OF PARKINSON'S DISEASE PSYCHOSIS
- U-1845 TREATMENT OF PSYCHOSIS OR A SYMPTOM THEREOF
- U-1846 TREATMENT OF A NEURODEGENERATIVE DISEASE OR A SYMPTOM THEREOF
- U-1847 METHOD OF TREATING A BACTERIAL INFECTION
- U-1848 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN
- U-1849 METHOD OF TREATING PULMONARY HYPERTENSION BY ADMINISTERING TREPROSTINIL OR A SALT THEREOF BY INHALATION USING A DEVICE
- U-1850 METHOD OF ADMINISTERING LEVETIRACETAM
- U-1851 A DOSING REGIMEN FOR THE TREATMENT OF HYPERCHOLESTEROLEMIA AND HYPERLIPIDEMIA IN PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA USING AT LEAST THREE STEP-WISE INCREASING DOSES
- U-1852 METHOD OF TREATING TYPE 2 DIABETES
- U-1853 METHOD OF TREATING TYPE 2 DIABETES MELLITUS BY ADMINISTERING A DIPEPTIDYL PEPTIDASE-IV INHIBITOR IN COMBINATION WITH METFORMIN AND, OPTIONALLY, A SULFONYLUREA
- U-1854 TREATMENT OF PRIMARY BILIARY CHOLANGITIS (PBC)
- U-1855 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS
- U-1856 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT HOMOZYGOUS FOR THE UGT1A1\*28 ALLELE
- U-1857 TO INCREASE BLOOD PRESSURE IN ADULTS WITH VASODILATORY SHOCK (E.G., POST-CARDIOTOMY OR SEPSIS) WHO REMAIN HYPOTENSIVE DESPITE FLUIDS AND CATECHOLAMINES
- U-1858 TREATMENT OF PLAQUE PSORIASIS
- U-1859 TREATMENT OF SCHIZOPHRENIA, ACUTE TREATMENT OF MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER, ADJUNCTIVE TREATMENT OF MAJOR DEPRESSIVE DISORDER, AND TREATMENT OF IRRITABILITY ASSOCIATED WITH AUTISTIC DISORDER
- U-1860 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1861 USE OF AN INHALER TO ADMINISTER DRY POWDER MEDICAMENT
- U-1862 TREATMENT OF POST-MYOCARDIAL INFARCTION
- U-1863 TREATMENT OF STROKE
- U-1864 TREATMENT OF MYOCARDIAL INFARCTION
- U-1865 TREATMENT OF THROMBOTIC STROKE
- U-1866 TREATMENT OF STABLE AND UNSTABLE ANGINA

**PATENT AND EXCLUSIVITY TERMS**

ADB 104 of 117

**PATENT USE**

- U-1867 METHOD OF INHIBITING PLATELET AGGREGATION
- U-1868 TREATMENT OF ARTERIAL THROMBOTIC COMPLICATIONS SELECTED FROM THE GROUP CONSISTING OF UNSTABLE ANGINA, THROMBOTIC OR EMBOLIC STROKE, TRANSIENT ISCHAEMIC ATTACKS, PERIPHERAL VASCULAR DISEASE AND MYOCARDIAL INFARCTION
- U-1869 TREATMENT OF AN ARTERIAL THROMBOTIC COMPLICATION IN A PATIENT WITH CORONARY ARTERY, CEREBROVASCULAR OR PERIPHERAL VASCULAR DISEASE
- U-1870 ZINGO IS A POWDER INTRADERMAL SYSTEM THAT IS CAPABLE OF DELIVERING FINE DRY POWDERED LIDOCAINE HYDROCHLORIDE MONOHYDRATE FOR LOCAL ANESTHETIC ACTION
- U-1871 TREATMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH STAGE 3 OR 4 CHRONIC KIDNEY DISEASE USING CONTROLLED RELEASE, ORAL 25-HYDROXYVITAMIN D
- U-1872 USE OF SUSTAINED RELEASE 25-HYDROXYVITAMIN D IN TREATING PATIENTS HAVING 25-HYDROXYVITAMIN D INSUFFICIENCY OR DEFICIENCY
- U-1873 ADMINISTRATION OF 25-HYDROXYVITAMIN D3 BY CONTROLLED RELEASE
- U-1874 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING OMEPRAZOLE ACCORDING TO CLAIMS 1-8
- U-1875 TREATMENT OF FREQUENT HEARTBURN BY ADMINISTERING S-OMEPRAZOLE TRIHYDRATE ACCORDING TO CLAIMS 1-3
- U-1876 METHOD OF ANESTHETIZING AT LEAST A PORTION OF THE MAXILLARY DENTAL ARCH
- U-1877 METHOD OF TREATING PULMONARY HYPERTENSION BY ORALLY ADMINISTERING A FORMULATION OF A PHARMACEUTICALLY ACCEPTABLE SALT OF TREPROSTINIL
- U-1878 FOR OPIOID DEPENDENCE
- U-1879 METHOD OF DIAGNOSING TUMORS USING POSITRON EMISSION TOMOGRAPHY
- U-1880 TREATMENT OF SIGNS AND SYMPTOMS OF DRY EYE DISEASE (DED)
- U-1881 IMPROVEMENT IN GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR
- U-1882 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS, REDUCTION IN FEVER THROUGH ANALGESIC AND ANTIPYRETIC ACTIVITY
- U-1883 TREATMENT OF GASTROINTESTINAL STROMAL TUMORS (GIST)
- U-1884 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1885 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY DECREASING FOOD INTAKE IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1886 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY INDUCING SATIETY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1887 USE OF LORCASERIN HYDROCHLORIDE FOR CHRONIC WEIGHT MANAGEMENT BY TREATING OBESITY IN PATIENTS ON A REDUCED-CALORIE DIET AND WHO HAVE ACHIEVED A GREATER THAN OR EQUAL TO 5% WEIGHT LOSS BY WEEK 12 OF TREATMENT
- U-1888 USE OF CONTROLLED RELEASE 25-HYDROXYVITAMIN D IN TREATING SECONDARY HYPERPARATHYROIDISM IN PATIENTS HAVING CHRONIC KIDNEY DISEASE
- U-1889 TREATMENT OF HCV INFECTION USING DASABUVIR/OMBITASVIR/PARITAPREVIR/RITONAVIR FIXED DOSE COMBINATION
- U-1890 OTC USE: ALLERGY SYMPTOM RELIEVER; TEMPORARILY RELIEVES THESE SYMPTOMS DUE TO HAY FEVER OR OTHER UPPER RESPIRATORY ALLERGIES; NASAL CONGESTION, RUNNY NOSE, SNEEZING, ITCHY NOSE, AND (ITCHY WATER EYES (AGES 12 AND UP))
- U-1891 TREATMENT OR PREVENTION OF NAUSEA AND VOMITING
- U-1892 METHOD OF TREATING LEFT VENTRICULAR DYSFUNCTION
- U-1893 METHOD OF TREATING MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER IN PEDIATRIC PATIENTS
- U-1894 COMBINATION TREATMENT WITH A GLITAZONE FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS
- U-1895 METHOD OF TREATING PROSTATE CANCER
- U-1896 SUPPLEMENT FOR VITAMIN B12 DEFICIENCIES

**PATENT AND EXCLUSIVITY TERMS**

ADB 105 of 117

**PATENT USE**

- U-1897 METHOD OF TREATING ACS USING ANGIOPLASTY WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)
- U-1898 METHOD OF INHIBITING PLATELET AGGREGATION WITH AGGRASTAT (TIROFIBAN HYDROCHLORIDE)
- U-1899 TREATMENT OF PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN
- U-1900 TREATMENT OF THE SIGNS SYMPTOMS OF DRY EYE DISEASE (DED)
- U-1901 TREATMENT OF SCHIZOAFFECTIVE DISORDER AS A MONOTHERAPY AND AS AN ADJUNCT TO MOOD STABILIZERS OR ANTIDEPRESSANTS
- U-1902 TREATMENT OR SECONDARY PREVENTION OF CARDIOVASCULAR DISEASE, CARDIOVASCULAR EVENTS, OR CEREBROVASCULAR EVENTS AND RISK-REDUCTION OF ASPIRIN-ASSOCIATED GASTRIC ULCERS
- U-1903 USE OF NALOXONE HYDROCHLORIDE FOR EMERGENCY TREATMENT OF KNOWN OR SUSPECTED OPIOID OVERDOSE, AS MANIFESTED BY RESPIRATORY AND/OR CENTRAL NERVOUS SYSTEM DEPRESSION.
- U-1904 (I) TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY; (II) RESTORING/INCREASING FUNCTIONAL DYSTROPHIN PROTEIN; OR (III) INDUCING SKIPPING; EACH OF (I)-(III) IN PATIENTS HAVING A CONFIRMED MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-1905 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, THE PATIENT HAVING A R117H MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE
- U-1906 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS, SUCH AS A PATIENT HAVING A G551D MUTATION IN CFTR, USING N-(5-HYDROXY-2,4-DI-TERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE
- U-1907 USE OF A DELIVERY DEVICE TO ADMINISTER A DOSE OF NALOXONE
- U-1908 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND FORM I LUMACAFTOR
- U-1909 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING IVACAFTOR AND LUMACAFTOR
- U-1910 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING THE DOSAGE UNIT OF CLAIM 1 OF U.S. PATENT NO. 8,716,338
- U-1911 METHOD OF TREATING A PATIENT HAVING CYSTIC FIBROSIS USING IVACAFTOR AND LUMACAFTOR
- U-1912 METHOD OF TREATING CYSTIC FIBROSIS IN A PATIENT, THE PATIENT HAVING THE F508DEL MUTATION IN CFTR, USING A DOSAGE UNIT AS DEFINED IN CLAIM 1 OF U.S. PATENT NO. 9,192,606
- U-1913 TREATMENT OF PEDIATRIC PATIENTS WITH BILATERAL OTITIS MEDIA WITH EFFUSION UNDERGOING TYMPANOSTOMY TUBE PLACEMENT
- U-1914 IN COMBINATION WITH RITUXIMAB, FOR THE TREATMENT OF PATIENTS WITH RELAPSED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-1915 METHOD OF TREATING TYPE 2 DIABETES MELLITUS IN PATIENTS WITH SEVERE CHRONIC RENAL IMPAIRMENT AND WHO ARE INELIGIBLE FOR METFORMIN THERAPY BY ADMINISTERING LINAGLIPTIN
- U-1916 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH CHEMOTHERAPY (CINV)
- U-1917 TREATMENT OF EXOCRINE PANCREATIC CANCER THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN
- U-1918 TREATMENT OF DUCHENNE MUSCULAR DYSTROPHY IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-1919 RESTORING AN mRNA READING FRAME TO INDUCE DYSTROPHIN PROTEIN PRODUCTION IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-1920 USE OF EXTENDED RELEASE ORAL 25-HYDROXYVITAMIN D3 IN TREATING SECONDARY HYPERPARATHYROIDISM IN ADULT PATIENTS HAVING CHRONIC KIDNEY DISEASE STAGE 3 OR STAGE 4
- U-1921 MANAGEMENT OF PAIN SEVERE ENOUGH TO REQUIRE DAILY, AROUND-THE-CLOCK, LONG-TERM OPIOID TREATMENT AND FOR WHICH ALTERNATIVE TREATMENT OPTIONS ARE INADEQUATE BY PROVIDING AN ABUSE-DETERRENT ORAL CONTROLLED RELEASE COMBINATION DRUG PRODUCT
- U-1922 INTRAVAGINAL PRASTERONE (DEHYDROEPIANDROSTERONE) AT A DAILY DOSE OF 6.5MG FOR THE TREATMENT OF DYSPAREUNIA, A SYMPTOM OF VULVAR AND VAGINAL ATROPHY, DUE TO

**PATENT AND EXCLUSIVITY TERMS**

ADB 106 of 117

**PATENT USE**

## MENOPAUSE

- U-1923 IMPROVEMENT IN GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS INADEQUATELY CONTROLLED BY BASAL INSULIN OR LIXISENATIDE BY USE OF A PEN INJECTOR WITH A THREADED DRIVE SLEEVE
- U-1924 KYPROLIS IS INDICATED IN COMBINATION WITH LENALIDOMIDE PLUS DEXAMETHASONE FOR THE TREATMENT OF PATIENTS WITH RELAPSED OR REFRACTORY MULTIPLE MYELOMA WHO HAVE RECEIVED ONE TO THREE LINES OF THERAPY
- U-1925 USE OF AN AUTO INJECTOR TO ADMINISTER NALOXONE HCL
- U-1926 METHOD OF TREATING, REDUCING THE INCIDENCE OF, OR PREVENTING AN ISCHEMIC EVENT IN A PATIENT UNDERGOING PCI BY ADMINISTERING INTRAVENOUSLY 30 UG/KG BOLUS BEFORE PCI AND CONTINUOUS INFUSION OF 4 UG/KG/MIN FOR AT LEAST 2 HOURS OR THE DURATION OF THE PCI
- U-1927 TREATMENT OF PATIENTS WITH ANAPLASTIC LYMPHOMA KINASE (ALK)-POSITIVE METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WHO HAVE PROGRESSED ON OR ARE INTOLERANT TO CRIZOTINIB
- U-1928 RUBRACA IS INDICATED AS MONOTHERAPY FOR THE TREATMENT OF PATIENTS WITH DELETERIOUS BRCA MUTATION (GERMLINE AND/OR SOMATIC) ASSOCIATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH TWO OR MORE CHEMOTHERAPIES.
- U-1929 TREATMENT OF DIABETES MELLITUS WITH AN INHALED INSULIN TO IMPROVE GLYCEMIC CONTROL USING A DRY POWDER INHALATION SYSTEM COMPRISING AN INHALER, A CARTRIDGE AND A DRY POWDER MEDICAMENT COMPRISING INSULIN IN A SINGLE INHALATION
- U-1930 METHOD OF AEROSOLIZING/DEAGGLOMERATING AN INSULIN DRY POWDER FOR USE IN TREATING DIABETES MELLITUS VIA ORAL INHALATION USING AN INHALER WITH A CARTRIDGE CONTAINING THE INSULIN DRY POWDER.
- U-1931 PROPHYLAXIS OR TREATMENT OF VENOUS AND ARTERIAL THROMBOTIC DISEASE
- U-1932 METHOD OF TREATING MILD TO MODERATE ATOPIC DERMATITIS.
- U-1933 TREATMENT OF POSTOPERATIVE INFLAMMATION AND REDUCTION OF OCULAR PAIN IN PATIENTS WHO HAVE UNDERGONE CATARACT SURGERY
- U-1934 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH A STRONG CYP1A2 INHIBITOR
- U-1935 REDUCTION OF THE RATE OF CARDIOVASCULAR DEATH, MYOCARDIAL INFARCTION, AND STROKE IN PATIENTS WITH A HISTORY OF MYOCARDIAL INFARCTION
- U-1936 TREATMENT OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1937 TREATMENT OF MYOCARDIAL INFARCTION IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1938 TREATMENT OF STROKE IN PATIENTS WITH ACUTE CORONARY SYNDROME OR A HISTORY OF MYOCARDIAL INFARCTION
- U-1939 ADMINISTRATION ONCE DAILY WITHIN TWO HOURS AFTER WAKING IN THE MORNING FOR IMPROVEMENT OF GLYCEMIC CONTROL IN A TYPE 2 DIABETES PATIENT
- U-1940 IMPROVEMENT IN THE APPEARANCE OF MODERATE TO SEVERE CONVEXITY OR FULLNESS ASSOCIATED WITH SUBMENTAL FAT IN ADULTS BY MEANS OF REDUCING SUBMENTAL FAT VOLUME AS DESCRIBED IN THE APPROVED LABELING
- U-1941 TREATMENT OF INFANTILE-ONSET SPINAL MUSCULAR ATROPHY
- U-1942 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INCREASING EXON-7 INCLUSION IN SMN2 mRNA
- U-1943 TREATMENT OF SPINAL MUSCULAR ATROPHY
- U-1944 TREATMENT OF SPINAL MUSCULAR ATROPHY BY INHIBITING AN SMN2 PRE-MRNA INTRONIC SPLICING SILENCER SITE
- U-1945 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN.
- U-1946 TREATMENT OF SMALL LYMPHOCYTIC LYMPHOMA
- U-1947 TREATMENT OF MARGINAL ZONE LYMPHOMA
- U-1948 A METHOD FOR TREATING CHRONIC MYELOID LEUKEMIA
- U-1949 FOR USE IN THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)
- U-1950 TREATMENT OF PATIENTS WITH ADVANCED (METASTATIC) NON-SMALL CELL LUNG CANCER WHOSE DISEASE PROGRESSED DURING OR AFTER PLATINUM-BASED CHEMOTHERAPY

**PATENT AND EXCLUSIVITY TERMS**

ADB 107 of 117

**PATENT USE**

- U-1951 TREATMENT OF ONYCHOMYCOSIS OF A TOENAIL
- U-1952 FOR USE IN THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA
- U-1953 REDUCE THE RISK OF STROKE IN PATIENTS WITH NONVALVULAR ATRIAL FIBRILLATION WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1954 TREATMENT OF DEEP VEIN THROMBOSIS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1955 TREATMENT OF PULMONARY EMBOLISM WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1956 FOLLOWING INITIAL 6 MONTHS TREATMENT FOR DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE), REDUCTION IN THE RISK OF RECURRENCE OF DVT AND OF PE WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1957 PROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM IN PATIENTS UNDERGOING KNEE OR HIP REPLACEMENT SURGERY, WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST FIVE CONSECUTIVE DAYS
- U-1958 FOR THE TREATMENT OF GENOTYPE 1, 2, 3 OR 4 CHRONIC HEPATITIS C VIRUS (HCV) INFECTION AS A COMPONENT OF A COMBINATION ANTIVIRAL TREATMENT REGIMEN WITH RIBAVIRIN
- U-1959 TOPICAL TREATMENT OF PERSISTENT FACIAL ERYTHEMA ASSOCIATED WITH ROSACEA IN ADULTS WITH 1% OXYMETAZOLINE HYDROCHLORIDE CREAM
- U-1960 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPIINE FOR TREATMENT OF SCHIZOPHRENIA IN ADULTS
- U-1961 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPIINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES (AGES 10 TO ADULT)
- U-1962 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPIINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: MAINTENANCE MONOTHERAPY IN ADULTS
- U-1963 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPIINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: AS ADJUNCTIVE TREATMENT TO LITHIUM OR VALPROATE IN ADULTS
- U-1964 ELEVATION OF INTRACELLULAR CGMP RESULTING IN INCREASED INTESTINAL FLUID AND ACCELERATED TRANSIT
- U-1965 FOR THE TREATMENT OF PULMONARY ARTERIAL HYPERTENSION (PAH) IN COMBINATION WITH Tadalafil, WHEREIN THE WEIGHT RATIO OF AMbrisentan TO Tadalafil IS ABOUT 1:2 TO ABOUT 1:3
- U-1966 USE OF THE ATYPICAL ANTIPSYCHOTIC ASENAPIINE FOR TREATMENT OF MANIC OR MIXED EPISODES OF BIPOLAR I DISORDER: ACUTE MONOTHERAPY OF MANIC OR MIXED EPISODES IN PEDIATRIC PATIENTS AGE 10-17
- U-1967 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WITH INSUFFICIENT GLYCEMIC CONTROL DESPITE THERAPY WITH ONE OR MORE CONVENTIONAL ANTIHYPERGLYCEMIC AGENTS BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN
- U-1968 METHOD OF TREATING TYPE 2 DIABETES IN PATIENTS WHO HAVE NOT BEEN PREVIOUSLY TREATED WITH AN ANTIHYPERGLYCEMIC AGENT BY ADMINISTERING LINAGLIPTIN IN COMBINATION WITH METFORMIN
- U-1969 TOPICAL TREATMENT OF ONYCHOMYCOSIS OF THE TOENAIL(S) DUE TO TRICHOPHYTON RUBRUM AND TRICHOPHYTON MENTAGROPHYTES
- U-1970 TREATMENT OF ONYCHOMYCHOSIS OF A TOENAIL CAUSED BY TRICHOPHYTON RUBRUM OR TRICHOPHYTON MENTAGROPHYTES
- U-1971 FOR THE TREATMENT OF PATIENTS WITH CHRONIC LYMPHOCYTIC LEUKEMIA
- U-1972 FOR THE TREATMENT OF PATIENTS WITH INDOLENT B-CELL NON-HODGKIN LYMPHOMA
- U-1973 METHOD OF TREATING CYSTIC FIBROSIS USING N-(5-HYDROXY-2,4-DITERT-BUTYL-PHENYL)-4-OXO-1H-QUINOLINE-3-CARBOXAMIDE AND 3-(6-(1-2,2-DIFLUOROBENZO[D][1,3]DIOXOL-5-YL) CYCLOPROPANECARBOXAMIDO)-3-METHYL PYRIDIN-2-YL) BENZOIC ACID
- U-1974 TREATMENT OF HALLUCINATIONS AND DELUSIONS ASSOCIATED WITH PARKINSON'S DISEASE PSYCHOSIS
- U-1975 METHOD OF INCREASING EYELASH GROWTH WITH BIMATOPROST
- U-1976 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO HAVE INADEQUATE CONTROL WITH DAPAGLIFLOZIN

**PATENT AND EXCLUSIVITY TERMS**

ADB 108 of 117

**PATENT USE**

- U-1977 METHOD FOR TREATING TYPE 2 DIABETES MELLITUS (T2DM) IN PATIENTS WHO ARE ALREADY TREATED WITH DAPAGLIFLOZIN AND SAXagliptin
- U-1978 TREATMENT OF ADVANCED PROSTATE CANCER WITH A REDUCED LIKELIHOOD OF CAUSING A GONADOTROPHIN RELEASING HORMONE AGONIST SIDE-EFFECT
- U-1979 THE TREATMENT OF CARCINOID SYNDROME DIARRHEA IN COMBINATION WITH SOMATOSTATIN ANALOG (SSA) THERAPY IN ADULTS INADEQUATELY CONTROLLED BY SSA THERAPY
- U-1980 A METHOD OF TREATING NOCTURIA DUE TO NOCTURNAL POLYURIA IN ADULTS
- U-1981 IN COMBINATION WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE-BASED THERAPY FOR TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER
- U-1982 USE OF REVIMID (LENALIDOMIDE) FOR TREATMENT OF PATIENTS WITH TRANSFUSION-DEPENDENT ANEMIA DUE TO LOW-OR INTERMEDIATE-1-RISK MYELODYSPLASTIC SYNDROMES ASSOCIATED WITH A DELETION 5Q ABNORMALITY WITH OR WITHOUT ADDITIONAL CYTOGENETIC ABNORMALITIES
- U-1983 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MANTLE CELL LYMPHOMA WHOSE DISEASE HAS RELAPSED OR PROGRESSED AFTER TWO PRIOR THERAPIES, ONE OF WHICH INCLUDED BORTEZOMIB
- U-1984 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE
- U-1985 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, AS MAINTENANCE FOLLOWING AUTOLOGOUS HEMATOPOIETIC STEM CELL TRANSPLANTATION (AUTO-HSCT)
- U-1986 USE OF REVIMID (LENALIDOMIDE) FOR THE TREATMENT OF PATIENTS WITH MULTIPLE MYELOMA, IN COMBINATION WITH DEXAMETHASONE, WHEREIN THOSE PATIENTS HAVE NOT RECEIVED PREVIOUS TREATMENT FOR MULTIPLE MYELOMA
- U-1987 METHOD OF CONTROLLING GLYCEMIA IN DIABETICS BY ADMINISTERING AN INITIAL DOSE OF INSULIN-FDKP WITH A MEAL; DETERMINING BLOOD GLUCOSE LEVEL 1-2 HRS AFTER AND ADMINISTERING A SUPPLEMENTAL DOSE OF INSULIN-FDKP IF POSTPRANDIAL GLUCOSE LEVEL IS >140 MG/DL
- U-1988 METHOD TO TREAT INFANTILE HEMANGIOMA
- U-1989 INTRAVITREAL TREATMENT OF MACULAR EDEMA FOLLOWING BRANCH RETINAL VEIN OCCLUSION (BRVO) OR CENTRAL RETINAL VEIN OCCLUSION (CRVO)
- U-1990 INTRAVITREAL TREATMENT OF DIABETIC MACULAR EDEMA
- U-1991 REDUCTION OF MORTALITY IN ACUTE MYCARDIAL INFARCTION
- U-1992 USE OF TROKENDI XR FOR PROPHYLACTIC TREATMENT OF MIGRAINE
- U-1993 ADJUNCTIVE TREATMENT TO LEVODOPA/CARBIDOPA IN PATIENTS WITH PARKINSON'S DISEASE EXPERIENCING "OFF" EPISODES
- U-1994 REDUCTION IN RISK OF OVERT HEPATIC ENCEPHALOPATHY (HE) IN ADULTS
- U-1995 TREATMENT OF TARDIVE DYSKINESIA
- U-1996 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-1997 IMPROVEMENT OF GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS IN COMBINATION WITH METFORMIN AND/OR A PPAR-GAMMA AGONIST AND/OR SULFONYLUREA AND/OR INSULIN
- U-1998 TREATING HR-POS., HER2-NEG. ADVANCED OR METASTATIC BREAST CANCER WITH PALBOCICLIB IN COMBO WITH AN AROMATASE INHIBITOR AS INITIAL ENDOCRINE BASED THERAPY IN POSTMENOPAUSAL WOMEN OR FULVESTRANT IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-1999 CHRONIC IDIOPATHIC CONSTIPATION
- U-2000 MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS
- U-2001 USE FOR THE TREATMENT OF ASTHMA IN PATIENTS 6 YEARS OF AGE AND OLDER
- U-2002 USE FOR MAINTENANCE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2003 A METHOD OF POSITIONING AN INTRAUTERINE SYSTEM BY HOLDING AN INSERTER HANDLE WITH ONE HAND, ADVANCING THE INSERTER THROUGH THE CERVIX AND INTO THE UTERUS, AND RETRACTING A SLIDER ON THE HANDLE TO RELEASE THE INTRAUTERINE SYSTEM
- U-2004 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY

**PATENT AND EXCLUSIVITY TERMS**

ADB 109 of 117

**PATENT USE**

- U-2005 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH GENERALIZED TONIC-CLONIC SEIZURES
- U-2006 REPLACEMENT THERAPY FOR ORAL CARBAMAZEPINE IN ADULTS WITH MIXED SEIZURE PATTERNS THAT INCLUDE PARTIAL SEIZURES WITH COMPLEX SYMPTOMATOLOGY, GENERALIZED TONIC-CLONIC SEIZURES, OR OTHER PARTIAL OR GENERALIZED SEIZURES
- U-2007 TREATMENT OF ADULT PATIENTS WITH NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML) WHO ARE FLT3 MUTATION-POSITIVE, IN COMBINATION WITH STANDARD CYTARABINE AND DAUNORUBICIN INDUCTION AND CYTARABINE CONSOLIDATION CHEMOTHERAPY
- U-2008 TREATMENT OF ADULT PATIENTS WITH AGGRESSIVE SYSTEMIC MASTOCYTOSIS (ASM), SYSTEMIC MASTOCYTOSIS WITH ASSOCIATED HEMATOLOGICAL NEOPLASM (SM-AHN), OR MAST CELL LEUKEMIA (MCL)
- U-2009 METHOD OF TREATING POSTMENOPAUSAL WOMEN WITH OSTEOPOROSIS AT HIGH RISK FOR FRACTURE.
- U-2010 ACUTE TREATMENT OF MIGRAINE BY DELIVERING A POWDERED SUBSTANCE COMPRISING SUMATRIPTAN VIA A BREATH-POWERED DELIVERY DEVICE
- U-2011 TREATMENT OF MIGRAINE VIA DELIVERY OF SUMATRIPTAN VIA THE NASAL CAVITY
- U-2012 A METHOD FOR TREATING OVARIAN CANCER BY ADMINISTERING RUCAPARIB, WHEREIN THE CANCER IS ASSOCIATED WITH A DELETERIOUS BRCA MUTATION
- U-2013 TREATMENT OF AMYOTROPHIC LATERAL SCLEROSIS (ALS)
- U-2014 A METHOD OF TREATING SECONDARY HYPERPARATHYROIDISM (SHPT)
- U-2015 SODIUM THIOSULFATE INJECTION IS INDICATED FOR SEQUENTIAL USE WITH SODIUM NITRITE FOR THE TREATMENT OF ACUTE CYANIDE POISONING
- U-2016 TREATMENT FOR ONYCHOMYCOSIS THAT IS TINEA UNGUIUM
- U-2017 TREATMENT OF OPIOID DEPENDENCE
- U-2018 MANAGEMENT OF MILD TO MODERATE PAIN, MANAGEMENT OF MODERATE TO SEVERE PAIN AS AN ADJUNCT TO OPIOID ANALGESICS IN A CRITICALLY ILL PATIENT WITH INTRAVENOUS IBUPROFEN IN NEED THEREOF
- U-2019 METHOD OF DELIVERING TO A PATIENT WITH DIABETES MELLITUS IN A SINGLE INHALATION, GREATER THAN 75% OF A DRY POWDER DOSE COMPRISING INSULIN AND FUMARYL DIKETOPIPERAZINE USING A HIGH RESISTANCE TO FLOW DRY POWDER INHALER.
- U-2020 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- U-2021 METHOD OF ADMINISTERING LEVETIRACETAM UNDER FASTED CONDITIONS
- U-2022 METHOD OF ADMINISTERING LEVETIRACETAM UNDER FED CONDITIONS
- U-2023 A METHOD OF INCREASING THE BIOAVAILABILITY OF GUAIFENESIN IN A SOLUTION CONTAINING 54% TO 66% BY WEIGHT OF PROPYLENE GLYCOL AND GLYCEROL, WHEREIN THE METHOD INCREASES THE CMAX BY AT LEAST 1.5 AND/OR INCREASES THE AUC (0-INF) BY AT LEAST 1.4
- U-2024 METHOD FOR TRANSDERMALLY DELIVERING A DRUG TO A USER IN NEED THEREOF
- U-2025 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-2026 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.
- U-2027 TAFINLAR(R) IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST.
- U-2028 TREATMENT OF ACUTE BACTERIAL SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY DESIGNATED SUSCEPTIBLE BACTERIA IN ADULTS
- U-2029 PREVENTING CONDITION CHARACTERIZED BY UNDESIRED THROMBOSIS
- U-2030 PROPHYLAXIS OF VENOUS THROMBOSIS
- U-2031 TAFINLAR IS INDICATED AS A SINGLE AGENT FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- U-2032 TAFINLAR IS INDICATED, IN COMBINATION WITH TRAMETINIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST.

**PATENT AND EXCLUSIVITY TERMS**

ADB 110 of 117

**PATENT USE**

- U-2033 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS
- U-2034 INHIBITING COAGULATION
- U-2035 PROPHYLAXIS OF VENOUS THROMBOEMBOLISM
- U-2036 A METHOD OF TREATING PULMONARY HYPERTENSION COMPRISING PARENTERALLY ADMINISTERING A FORMULATION COMPRISING A) 0.1 TO 5% W/V OF TREPROSTINIL OR A PHARMACEUTICALLY ACCEPTABLE SALT THEREOF AND B) A CITRATE BUFFER
- U-2037 MEKINIST IS INDICATED, AS A SINGLE AGENT OR IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE OR METASTATIC MELANOMA WITH BRAF V600E OR V600K MUTATIONS, AS DETECTED BY AN FDA-APPROVED TEST
- U-2038 MEKINIST IS INDICATED, IN COMBINATION WITH DABRAFENIB, FOR THE TREATMENT OF PATIENTS WITH METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH BRAF V600E MUTATION AS DETECTED BY AN FDA-APPROVED TEST
- U-2039 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1, 2, 3, 4, 5, OR 6 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING AN NS5A INHIBITOR
- U-2040 TREATMENT OF ADULT PATIENTS WITH CHRONIC HCV INFECTION WHO HAVE GENOTYPE 1A OR 3 INFECTION AND HAVE PREVIOUSLY BEEN TREATED WITH AN HCV REGIMEN CONTAINING SOFOSBUVIR WITHOUT AN NS5A INHIBITOR
- U-2041 TREATMENT OF PARTIAL-ONSET SEIZURES
- U-2042 DISCONTINUING ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2043 EXTENDED ADJUVANT TREATMENT OF ADULT PATIENTS WITH EARLY STAGE HER2-OVEREXPRESSED/AMPLIFIED BREAST CANCER, TO FOLLOW ADJUVANT TRASTUZUMAB BASE THERAPY
- U-2044 DOSE REDUCTION OF PIRFENIDONE BY ABOUT ONE HALF DURING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TWICE DAILY (1500 MG/DAY) TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER
- U-2045 ADMINISTRATION OF PIRFENIDONE AND AVOIDING CONCURRENT ADMINISTRATION OF CIPROFLOXACIN AT A DOSE OF 750 MG TO REDUCE DRUG INTERACTIONS IN TREATMENT OF A FIBROTIC, INFLAMMATORY, OR AUTOIMMUNE DISORDER
- U-2046 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6
- U-2047 ADMINISTERING PIRFENIDONE CONCURRENTLY WITH FLUVOXAMINE, THE PIRFENIDONE AT A DOSE OF ABOUT 801 MG/DAY TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2048 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2049 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INHIBITOR TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE AND THEN ADMINISTERING PIRFENIDONE
- U-2050 ADMINISTERING PIRFENIDONE WHILE AVOIDING CO-ADMINISTRATION OF FLUVOXAMINE TO AVOID DRUG INTERACTIONS WITH PIRFENIDONE
- U-2051 DISCONTINUING SMOKING TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2052 DISCONTINUING ADMINISTRATION OF A STRONG CYP1A2 INDUCER TO AVOID REDUCED PIRFENIDONE EFFICACY AND THEN ADMINISTERING PIRFENIDONE
- U-2053 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2, INCLUDING CIGARETTE SMOKE, TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2054 ADMINISTERING PIRFENIDONE WHILE AVOIDING CONCOMITANT ADMINISTRATION OF A STRONG INDUCER OF CYP1A2 TO AVOID REDUCED PIRFENIDONE EFFICACY
- U-2055 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 LIVER ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2056 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY, FOLLOWING BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2057 DOSING 2403 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF

**PATENT AND EXCLUSIVITY TERMS**

ADB 111 of 117

**PATENT USE**

- U-2058 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400MG/DAY DOSE, FOLLOWED BY ADMINISTERING 2403MG/DAY IN TREATMENT OF IPF
- U-2059 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE IN TREATMENT OF IPF
- U-2060 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER AST AND/OR ALT AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN AT LEAST 1600MG/DAY IN TREATMENT OF IPF
- U-2061 DOSING OF AT LEAST 1600 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2062 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY DOSE IN TREATMENT OF IPF
- U-2063 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING AT LEAST 1600 MG/DAY IN TREATMENT OF IPF
- U-2064 DOSING AT LEAST 1602 MG/DAY FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2065 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION
- U-2066 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE, FOLLOWED BY FULL DAILY DOSE
- U-2067 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2068 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, FOLLOWED BY FULL DAILY DOSE
- U-2069 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING A SUB-1600 MG/DAY DOSE, FOLLOWED BY ADMINISTERING AT LEAST 1602 MG/DAY
- U-2070 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN BIOMARKER ALT OR AST AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS OF LIVER FUNCTION ARE WITHIN NORMAL LIMITS, THEN SUB-1600 MG/DAY, THEN AT LEAST 1602 MG/DAY
- U-2071 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2072 FULL DAILY DOSING FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2073 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY ADMINISTERING FULL DAILY DOSE IN TREATMENT OF IPF
- U-2074 DOSING 1602 MG/DAY PIRFENIDONE FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION IN TREATMENT OF IPF
- U-2075 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2076 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING 801 MG/DAY FOLLOWED BY ADMINISTERING 1602 MG/DAY IN TREATMENT OF IPF
- U-2077 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY ADMINISTERING SUB-2400 MG/DAY DOSE THEN FULL DAY DAILY DOSE IN TREATMENT OF IPF

**PATENT AND EXCLUSIVITY TERMS**

ADB 112 of 117

**PATENT USE**

- U-2078 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN LIMITS, THEN SUB-2400MG/DAY DOSE, THEN FULL DAILY DOSE IN TREATMENT OF IPF
- U-2079 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF FIBROSIS AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2080 PIRFENIDONE DOSE ESCALATION REGIMEN FOR TREATMENT OF IPF AS 801 MG/DAY FOR DAYS 1-7 OF THE REGIMEN, 1602 MG/DAY FOR DAYS 8-14 OF THE REGIMEN, AND 2403 MG/DAY FOR AT LEAST DAY 15 OF THE REGIMEN
- U-2081 DISCONTINUING USE OF A CYP1A2 INHIBITOR THAT IS A MODERATE TO STRONG INHIBITOR OF BOTH CYP1A2 AND ANOTHER CYP ENZYME SELECTED FROM CYP2C9, CYP2C19, AND CYP2D6 AND THEN ADMINISTERING PIRFENIDONE
- U-2082 MODIFYING PIRFENIDONE ADMINISTRATION FROM A DOSE OF ABOUT 2400 MG/DAY DOWNWARD BY ABOUT 1600 MG/DAY WHILE CO-ADMINISTERING FLUVOXAMINE TO REDUCE DRUG INTERACTIONS WITH FLUVOXAMINE
- U-2083 DOSAGE MODIFICATION FOLLOWING GRADE 2 ABNORMALITY IN LIVER FUNCTION BIOMARKER AFTER PIRFENIDONE ADMINISTRATION, BY DISCONTINUING PIRFENIDONE UNTIL BIOMARKERS ARE WITHIN NORMAL LIMITS, FOLLOWED BY 801 MG/DAY, DOSE, THEN 1602 MG/DAY IN TREATMENT OF IPF
- U-2084 TREATMENT OF SEVERE CHRONIC PAIN VIA INTRATHECAL INFUSION OF ZICONOTIDE IN PATIENTS ALSO RECEIVING MORPHINE
- U-2085 TREATMENT OF NON-24-HOUR SLEEP-WAKE DISORDER BY AVOIDING THE USE OF TASIMELTEON IN COMBINATION WITH RIFAMPIN
- U-2086 A METHOD FOR ADMINISTERING ESTRADIOL COMPRISING A MONOLITHIC TRANSDERMAL DRUG DELIVERY SYSTEM CONSISTING OF (I) A BACKING LAYER AND (II) A SINGLE ADHESIVE POLYMER MATRIX LAYER AS CLAIMED IN US PATENT NO. 9730900
- U-2087 TREATMENT OF RELAPSED OR REFRACTORY ACUTE MYELOID LEUKEMIA (AML) WITH AN ISOCITRATE DEHYDROGENASE-2 (IDH2) MUTATION
- U-2088 TREATMENT OF PARTIAL-ONSET SEIZURES WITH OR WITHOUT SECONDARILY GENERALIZED SEIZURES IN PATIENTS WITH EPILEPSY
- U-2089 TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES AS ADJUNCTIVE THERAPY IN PATIENTS WITH EPILEPSY
- U-2090 FOR THE TREATMENT OF ADULTS WITH NEWLY-DIAGNOSED THERAPY-RELATED ACUTE MYELOID LEUKEMIA (T-AML) OR AML WITH MYELODYSPLASIA-RELATED CHANGES (AML-MRC)
- U-2091 TREATMENT OF METASTATIC ADENOCARCINOMA OF THE PANCREAS THAT HAS PROGRESSED ON GEMCITABINE-BASED THERAPY, IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN, IN A PATIENT NOT HOMOZYGOUS FOR THE UGT1A1\*28 ALLELE
- U-2092 METHOD FOR CONFIRMING DOSE DELIVERY
- U-2093 TREATMENT OF TYPE II SPINAL MUSCULAR ATROPHY
- U-2094 TREATMENT OF TYPE III SPINAL MUSCULAR ATROPHY
- U-2095 MITOSOL IS AN ANTIMETABOLITE INDICATED AS AN ADJUNCT TO AB EXTERNO GLAUCOMA SURGERY. IT IS INTENDED FOR TOPICAL APPLICATION TO THE SITE OF GLAUCOMA FILTRATION SURGERY
- U-2096 SOTYLIZE IS INDICATED FOR THE MAINTENANCE OF NORMAL SINUS RHYTHM [DELAY IN TIME TO RECURRANCE OF ATRIAL FIBRILLATION/ATRIAL FLUTTER (AFIB/AFL)] IN PATIENTS WITH SYMPTOMATIC AFIB/AFL WHO ARE CURRENTLY IN SINUS RHYTHM
- U-2097 TREATMENT OF DMD IN PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2098 INCREASING PRODUCTION OF FUNCTIONAL DYSTROPHIN PROTEIN IN DMD PATIENTS HAVING A MUTATION OF THE DMD GENE THAT IS AMENABLE TO EXON 51 SKIPPING
- U-2099 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF AIRFLOW OBSTRUCTION IN PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING BRONCHITIS AND/OR EMPHYSEMA
- U-2100 INDICATED FOR THE ONCE-DAILY TREATMENT OF ASTHMA IN PATIENTS 18 YEARS AND OLDER
- U-2101 MAINTENANCE TREATMENT OF RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY

**PATENT AND EXCLUSIVITY TERMS**

ADB 113 of 117

**PATENT USE**

- U-2102 TREATMENT OF DELETERIOUS OR SUSPECTED DELETERIOUS GERMLINE BRCA-MUTATED ADVANCED OVARIAN CANCER WHO HAVE BEEN TREATED WITH THREE OR MORE PRIOR LINES OF CHEMOTHERAPY BASED ON AN FDA-APPROVED COMPANION DIAGNOSTIC FOR LYNPARZA
- U-2103 MAINTENANCE TREATMENT OF BRCA-MUTATED RECURRENT EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER, WHO ARE IN A COMPLETE OR PARTIAL RESPONSE TO PLATINUM-BASED CHEMOTHERAPY
- U-2104 TREATMENT OF HYPERURICEMIA ASSOCIATED WITH GOUT IN PATIENTS WHO HAVE NOT ACHIEVED TARGET SERUM URIC ACID LEVELS WITH A MEDICALLY APPROPRIATE DAILY DOSE OF ALLOPURINOL ALONE
- U-2105 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING IMMEDIATE RELEASE LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2106 TREATMENT OF DYSKINESIA IN PATIENTS WITH PARKINSON'S DISEASE RECEIVING LEVODOPA-BASED THERAPY, WITH OR WITHOUT CONCOMITANT DOPAMINERGIC MEDICATIONS
- U-2107 TREATMENT OF LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, DIFFERENTIATED THYROID CARCINOMA REFRACTORY TO RADIOACTIVE IODINE TREATMENT
- U-2108 TREATMENT OF HORMONE RECEPTOR POSITIVE HER2-NEGATIVE ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN NOT PREVIOUSLY TREATED WITH ENDOCRINE THERAPY
- U-2109 CARSPIR IS INDICATED FOR TREATMENT OF NYHA CLASS III-IV HEART FAILURE AND REDUCED EJECTION FRACTION TO INCREASE SURVIVAL, MANAGE EDEMA, AND TO REDUCE THE NEED FOR HOSPITALIZATION FOR HEART FAILURE
- U-2110 METHOD FOR CHRONIC WEIGHT MANAGEMENT IN PATIENTS WITH MODERATE RENAL IMPAIRMENT WHO ARE OBESE, OR OVERWEIGHT AND HAVE AT LEAST ONE WEIGHT RELATED COMORBID CONDITION
- U-2111 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 1-5 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 1-5
- U-2112 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 6 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 6
- U-2113 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 7 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 7
- U-2114 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 9 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 9
- U-2115 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 10 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 10
- U-2116 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 12 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 12
- U-2117 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 14-15 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 14-15
- U-2118 ADJUNCT TO DIET AND EXERCISE TO TREAT GLUCOSE INTOLERANCE IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIMS 16-18 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIMS 16-18
- U-2119 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING BROMOCRIPTINE MESYLATE AND A FIRST-PHASE INSULIN SECRETAGOGUE AS RECITED IN CLAIM 19 AND WHEREIN THE EFFECTS ARE AS RECITED IN CLAIM 19
- U-2120 TREATMENT OF PATIENTS 18 YEARS OF AGE AND OLDER WITH COMPLICATED URINARY TRACT

**PATENT AND EXCLUSIVITY TERMS**

ADB 114 of 117

**PATENT USE**

## INFECTIONS CAUSED BY SUSCEPTIBLE MICROORGANISMS

- U-2121 TREATMENT OF PARTIAL-ONSET SEIZURES IN A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ABSENCE SEIZURES
- U-2122 USE FOR REDUCING EXACERBATIONS OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2123 TREATMENT OF PARTIAL-ONSET SEIZURES IN PATIENTS WITH EPILEPSY WHO HAVE BEEN PREVIOUSLY TREATED WITH OXCARBAZEPINE
- U-2124 TREATMENT OF ADULT PATIENTS WITH RELAPSED FOLLICULAR LYMPHOMA WHO HAVE RECEIVED AT LEAST TWO PRIOR SYSTEMIC THERAPIES
- U-2125 THE TREATMENT OF AN INFLAMMATORY DISORDER OF THE RESPIRATORY TRACT BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR AGONIST
- U-2126 USE OF FLUTICASONE FUROATE FOR THE TREATMENT OF AN INFLAMMATORY OR ALLERGIC CONDITIONS, INCLUDING CHRONIC OBSTRUCTIVE PULMONARY DISEASE
- U-2127 INDICATED FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF PATIENTS WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND/OR EMPHYSEMA
- U-2128 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHELINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLINIMUM, VIA INHALATION
- U-2129 METHOD OF INHIBITING THE BINDING OF ACETYLCHOLINE TO AN ACETYLCHOLINE RECEPTOR IN THE RESPIRATORY TRACT OF A HUMAN, COMPRISING CONTACTING THE RECEPTOR WITH AN EFFECTIVE AMOUNT OF UMECLINIMUM, VIA TOPICAL APPLICATION
- U-2130 TREATMENT OF PARTIAL ONSET SEIZURES IN PATIENTS WITH EPILEPSY AGED 16 YEARS AND OLDER WITH EPILEPSY
- U-2131 REDUCTION OF ELEVATED INTRAOCCULAR PRESSURE (IOP) IN PATIENTS WITH GLAUCOMA OR OCULAR HYPERTENSION, WITH COMPARABLE EFFICACY, AND A REDUCTION IN SPECIFIED ADVERSE EVENTS, COMPARED TO BRIMONIDINE 0.2% TID
- U-2132 IN COMBINATION WITH FULVESTRANT FOR THE TREATMENT OF WOMEN WITH HORMONE RECEPTOR (HR)-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY
- U-2133 METHOD OF DELIVERING FLUTICASONE PROPIONATE TO A NASAL AIRWAY
- U-2134 THE TREATMENT OF CHRONIC OBSTRUCTIVE PULMONARY DISEASE BY ONCE-PER-DAY ADMINISTRATION OF A PHARMACEUTICAL FORMULATION COMPRISING FLUTICASONE FUROATE AND A LONG-ACTING BETA2 ADRENORECEPTOR
- U-2135 AS MONOTHERAPY FOR THE TREATMENT OF ADULT PATIENTS WITH HR-POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER WITH DISEASE PROGRESSION FOLLOWING ENDOCRINE THERAPY AND PRIOR CHEMOTHERAPY IN THE METASTATIC SETTING
- U-2136 TREATMENT OF NEUROPATHIC PAIN ASSOCIATED WITH DIABETIC PERIPHERAL NEUROPATHY
- U-2137 TREATMENT OF POSTHERPETIC NEURALGIA
- U-2138 TOPICAL TREATMENT OF ACTINIC KERATOSIS OF THE FACE OR SCALP USING MORE THAN ONE TREATMENT COURSE OF INGENOL MEBUTATE
- U-2139 TREATMENT OF TYPE 2 DIABETES MELLITUS IN COMBINATION WITH EXENATIDE
- U-2140 METHOD OF TREATING PARTIAL ONSET SEIZURES IN PATIENTS 4 YEARS OF AGE AND OLDER
- U-2141 TREATMENT OF CHRONIC HEPATITIS C VIRUS (HCV) GENOTYPE 1, 2, 3, 4, 5, OR 6
- U-2142 REDUCTION IN THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS (DVT) AND/OR PULMONARY EMBOLISM (PE) IN PATIENTS AT CONTINUED RISK FOR RECURRENT DVT AND/OR AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS
- U-2143 AFTER COMPLETION OF INITIAL TREATMENT LASTING AT LEAST 6 MONTHS, TO REDUCE THE RISK OF RECURRENCE OF DEEP VEIN THROMBOSIS AND/OR PULMONARY EMBOLISM IN CERTAIN PATIENTS WITH ONCE DAILY, RAPID-RELEASE TABLET ADMINISTERED FOR AT LEAST 5 CONSECUTIVE DAYS
- U-2144 REDUCTION OF INTRAOCCULAR PRESSURE (IOP) IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- U-2145 TREATMENT OF ADULT PATIENTS WITH MANTLE CELL LYMPHOMA WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY
- U-2146 IMPROVEMENT IN GLYCEMIC CONTROL IN DIABETES MELLITUS PATIENTS BY USE OF A PEN INJECTOR WITH A ROTATING DRIVE SLEEVE

**PATENT AND EXCLUSIVITY TERMS**

ADB 115 of 117

**PATENT USE**

- U-2147 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ORALLY ADMINISTERING 20MG OF TASIMELTEON ONCE DAILY BEFORE BEDTIME
- U-2148 A METHOD OF PROVIDING NITRIC OXIDE THERAPY TO A PATIENT BY MEASURING AND DISPLAYING AN INDICATION OF THE CALCULATED DELIVERY CONCENTRATION OF NITRIC OXIDE AS COMPARED TO THE DESIRED DELIVERY CONCENTRATION OF NITRIC OXIDE
- U-2149 TREATMENT OF NON-24 HOUR SLEEP-WAKE DISORDER BY ADMINISTERING TASIMELTEON
- U-2150 TREATMENT OF CHRONIC GRAFT-VERSUS-HOST DISEASE
- U-2151 METHOD OF TREATING PAIN OR INFLAMMATION WITH AN INJECTABLE CONTROLLED OR SUSTAINED RELEASE FORMULATION OF TRIAMCINOLONE ACETONIDE
- U-2152 TREATMENT OF PAIN ASSOCIATED WITH IRRITABLE BOWEL SYNDROME WITH DIARRHEA (IBS-D) WITH VIBERZI (ELUXADOLINE)
- U-2153 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2154 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2155 REDUCING BODY WEIGHT IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2156 REDUCING HBA1C IN A HUMAN IN NEED THEREOF USING A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2157 TREATING TYPE 2 DIABETES MELLITUS BY STIMULATING INSULIN RELEASE
- U-2158 DECREASING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY USING A SUSTAINED-RELEASE COMPOSITION
- U-2159 TREATMENT OF CHRONIC LYMPHOCYTIC LEUKEMIA/SMALL LYMPHOCYTIC LYMPHOMA
- U-2160 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 5 MG OF MELOXICAM
- U-2161 TREATMENT OF NAUSEA AND VOMITING, INCLUDING THE PREVENTION OF ACUTE AND DELAYED NAUSEA AND VOMITING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF HIGHLY OR MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- U-2162 FOR CLEANSING THE LARGE INTESTINE AS A PREPARATION FOR COLONOSCOPY
- U-2163 TREATMENT OF HR-POSITIVE, HUMAN EPIDERMAL GROWTH FACTOR RECEPTOR 2 (HER-2)-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH PALBOCOCLIB OR ABEMACICLIB IN WOMEN WITH DISEASE PROGRESSION AFTER ENDOCRINE THERAPY
- U-2164 ZELBORAF IS INDICATED FOR THE TREATMENT OF PATIENTS WITH ERDHEIM-CHESTER DISEASE WITH BRAF V600 MUTATION
- U-2165 MANAGEMENT OF OSTEOARTHRITIS PAIN BY ADMINISTERING 10 MG OF MELOXICAM
- U-2166 TREATMENT OF MAJOR DEPRESSIVE EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- U-2167 METHOD OF USING A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER
- U-2168 METHOD OF USING A LOGIC CIRCUIT TO STABILIZE BATTERY VOLTAGE SUPPLIED TO A SENSOR EMBEDDED WITH A TABLET AND THAT COMMUNICATES INFORMATION VIA A SIGNAL THROUGH THE BODY OF A PATIENT TO A RECEIVER
- U-2169 METHOD OF USING A RECEIVER TO IDENTIFY A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT
- U-2170 METHOD OF USING A RECEIVER TO RECEIVE A SIGNAL FROM A TABLET EMBEDDED WITH A SENSOR THAT COMMUNICATES INFORMATION THROUGH THE BODY OF A PATIENT
- U-2171 ADJUVANT TREATMENT OF ADULT PATIENTS AT HIGH RISK OF RECURRENT RCC FOLLOWING NEPHRECTOMY
- U-2172 METHOD TO TREAT SEVERE ALLERGIC EMERGENCIES IN PATIENTS WEIGHING 7.5 TO 15 KG (16.5 TO 33 LBS)
- U-2173 TREATING OPIOID DEPENDENCE BY ADMINISTERING BUPRENORPHINE
- U-2174 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH
- U-2175 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE ONCE MONTHLY
- U-2176 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE
- U-2177 TREATING OPIOID ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE
- U-2178 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE COMPOSITION WITH 28 DAY DOSE DURATION

**PATENT AND EXCLUSIVITY TERMS**

ADB 116 of 117

**PATENT USE**

- U-2179 IN SITU FORMATION OF SOLID BUPRENORPHINE COMPOSITION
- U-2180 TREATING ADDICTION WITH 100 MG OR 300 MG DOSE OF BUPRENORPHINE
- U-2181 TREATING OPIOID DEPENDENCY BY SUBCUTANEOUSLY ADMINISTERING BUPRENORPHINE
- U-2182 IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS
- U-2183 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1 AND 13
- U-2184 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 1, 13, AND 14
- U-2185 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15 AND 27
- U-2186 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 15, 27, AND 28
- U-2187 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29 AND 39
- U-2188 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 29, 39, AND 40
- U-2189 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41 AND 52
- U-2190 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 41, 52, AND 53
- U-2191 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54 AND 64
- U-2192 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 54, 64, AND 65
- U-2193 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66 AND 75
- U-2194 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ADMINISTERING A DOSAGE FORM COMPRISING MICRONIZED BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 66, 75, AND 76
- U-2195 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77 AND 87
- U-2196 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 77, 87, AND 88
- U-2197 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89 AND 99
- U-2198 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES BY ORALLY ADMINISTERING A DOSAGE FORM COMPRISING BROMOCRIPTINE AND ONE OR MORE EXCIPIENTS AS RECITED IN CLAIMS 89, 99, AND 100
- U-2199 TREATMENT OF SCHIZOPHRENIA WITH IMPROVEMENT IN ATTENTION FUNCTION IN SCHIZOPHRENIA
- U-2200 COMBINATION TREATMENT WITH INSULIN GLARGINE WITH OR WITHOUT METFORMIN FOR IMPROVEMENT OF GLYCEMIC CONTROL IN TYPE 2 DIABETES MELLITUS PATIENTS
- U-2201 TREATMENT OF BIPOLAR DEPRESSION WITH IMPROVEMENT IN ATTENTION FUNCTION IN BIPOLAR DISORDER
- U-2202 OZEMPIC IS INDICATED AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC

**PATENT AND EXCLUSIVITY TERMS**

ADB 117 of 117

**PATENT USE**

## CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS

- U-2203 A METHOD OF PROVIDING A SUBJECT WITH A THERAPEUTICALLY EFFECTIVE AMOUNT OF RACEMIC METHYLPHENIDATE BY ORALLY ADMINISTERING TO SAID SUBJECT A SINGLE METHYLPHENIDATE EXTENDED RELEASE CHEWABLE TABLET AS CLAIMED
- U-2204 TREATING PATIENTS WITH ACUTE PROMYELOCYTIC LEUKEMIA (APL) WHO ARE REFRACTORY TO, OR HAVE RELAPSED FROM, RETINOID AND ANTHRACYCLINE CHEMOTHERAPY, AND WHOSE APL IS CHARACTERIZED BY THE PRESENCE OF THE T(15;17) TRANSLOCATION OR PML/RAR-ALPHA GENE EXPRESSION
- U-2205 TREATMENT OF SEBORRHEIC KERATOSES THAT ARE RAISED
- U-2206 TREATING OPIOID DEPENDENCY BY ADMINISTERING BUPRENORPHINE
- U-2207 TREATING ADDICTION BY SUBCUTANEOUS INJECTION OF BUPRENORPHINE
- U-2208 TREATING ADDICTION BY ONCE PER MONTH ADMINISTRATION OF BUPRENORPHINE
- U-2209 TREATING OPIOID ADDICTION BY ADMINISTERING BUPRENORPHINE ONCE PER MONTH
- U-2210 TREATING OPIOID ADDICTION BY 100 MG OR 300 MG DOSE BUPRENORPHINE
- U-2211 TREATING OPIOID ADDICTION BY ADMINISTRATION OF BUPRENORPHINE
- U-2212 REDUCING FASTING PLASMA GLUCOSE IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2213 REDUCING HBA1C IN A HUMAN IN NEED THEREOF IN COMBINATION WITH A SUSTAINED-RELEASE COMPOSITION CONTAINING EXENDIN-4
- U-2214 AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES
- U-2215 ERTUGLIFLOZIN IN COMBINATION WITH SITAGLIPTIN AND IN FURTHER COMBINATION WITH METFORMIN AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2216 ERTUGLIFLOZIN AND SITAGLIPTIN IN COMBINATION AS AN ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN ADULTS WITH TYPE 2 DIABETES MELLITUS
- U-2217 TREATING HIGH OUTPUT SHOCK WITH ANGIOTENSIN II BY INCREASING MEAN ARTERIAL PRESSURE IN PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2218 MAINTAINING MEAN ARTERIAL PRESSURE OF ABOUT 65 MMHG OR HIGHER WITH ANGIOTENSIN II IN SHOCK PATIENTS TREATED WITH CATECHOLAMINES AND REDUCING CATECHOLAMINE USE
- U-2219 TREATMENT OF CHRONIC SMALL LYMPHOCYTIC LEUKEMIA
- U-2220 A METHOD FOR THE DIAGNOSIS OF ADULT GROWTH HORMONE DEFICIENCY BY MEASURING THE LEVEL OF GROWTH HORMONE AFTER ORAL ADMINISTRATION OF MACIMORELIN
- U-2221 TREATING REFRACTORY HYPOTENSION WITH ABOUT 20 MG/KG/MIN ANGIOTENSIN II IN A PATIENT RECEIVING VASOPRESSOR
- U-2222 RELIEVES REDNESS OF THE EYE DUE TO MINOR EYE IRRITATIONS
- U-2223 METHOD OF TREATING ANGINA PECTORIS